# Status of the Pacific Hake (whiting) stock in U.S. and Canadian waters in 2024 



Joint Technical Committee of the Pacific Hake/Whiting Agreement Between the Governments of the United States and Canada

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This document reports the collaborative efforts of the official U.S. and Canadian members of the Joint Technical Committee, and others that contributed significantly.

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## ONE-PAGE SUMMARY

- The stock assessment model for 2024 has the same population dynamics structure as the 2023 model. The model is fit to an acoustic survey index of biomass, a relative index of age- 1 fish, annual commercial catch data, and age-composition data from the survey and commercial fisheries.
- Data for 2023 were included for each data set and minor changes to pre-2023 data were made as necessary. In addition, a new model-based approach was used to develop the input weight-at-age matrix, and time-varying temperature-dependent maturity was introduced to better inform fecundity.
- Coast-wide catch in 2023 was $263,981 \mathrm{t}$ [t represents metric tons], $22 \%$ below the average over the most recent 10 years ( $338,606 \mathrm{t}$ ), out of a total allowable catch (TAC), adjusted for carryovers, of $625,000 \mathrm{t}$. The U.S. caught $240,424 \mathrm{t}$ ( $52.1 \%$ of their quota) and Canada caught $23,557 \mathrm{t}$ ( $14.4 \%$ of their quota).
- The median estimate of the 2024 relative spawning biomass (female spawning biomass at the start of 2024 divided by that at unfished equilibrium, $\mathrm{B}_{0}$ ) is $99 \%$ but is highly uncertain (with $95 \%$ credible interval from $45 \%$ to $230 \%$ ). After declining from 2018-2022, the median relative spawning biomass increased in 2023 and 2024, due to the estimated above average, but uncertain, size of the 2020 and 2021 cohorts entering maturity.
- The median estimate of female spawning biomass at the start of 2024 is $1,884,950 \mathrm{t}$ (with $95 \%$ credible interval from 853,207 to $4,828,382 \mathrm{t}$ ). This is an upward shift from this assessment's estimate for the 2023 female spawning biomass of 1,335,485 t (with $95 \%$ credible interval from 652,495 to $3,224,819 \mathrm{t}$ ).
- The estimated probability that female spawning biomass at the start of 2024 is below the $B_{40 \%}\left(40 \%\right.$ of $\left.B_{0}\right)$ reference point is $1.3 \%$, and the probability that the relative fishing intensity exceeded 1 in 2023 is $0.4 \%$. The joint probability of both these occurring is $0.2 \%$.
- Based on the default harvest rule, the estimated median catch limit for 2024 is $747,588 \mathrm{t}$ (with $95 \%$ credible interval from 298,355 to 2,124,832 t).
- Projections were conducted across a wide-range of catch levels due to high uncertainty in estimates of recent and forecasted recruitment. Projections setting the 2024 and 2025 catches equal to the 2023 coast-wide TAC of 625,000 t show the estimated median relative spawning biomass decreasing from $99 \%$ in 2024 to $94 \%$ in 2025 and then to $83 \%$ in 2026, with a $11 \%$ chance of the female spawning biomass falling below $B_{40 \%}$ in 2026. There is an estimated $76 \%$ chance of the female spawning biomass declining from 2024 to 2025, an $84 \%$ chance of it declining from 2025 to 2026, and an $83 \%$ chance of it declining from 2026 to 2027 with a constant catch of 625,000 t.
- Despite estimates of a healthy stock status, the recent lack of survey abundance and fishery catch in Canada suggests a population structure not conducive to achieving harvest quotas in northern fisheries over recent years.


## EXECUTIVE SUMMARY

## Stock

This assessment reports the status of the coastal Pacific Hake (or Pacific whiting, Merluccius productus) stock off the west coast of the United States and Canada at the start of 2024. This stock exhibits seasonal migratory behavior, ranging from offshore and generally southern waters during the winter spawning season to coastal areas between northern California and northern British Columbia during the spring, summer, and fall when the fishery is conducted. The stock tends to move farther to the north during the summer in years with warmer water compared to years with colder waters. Older Pacific Hake tend to migrate farther north than younger Pacific Hake in all years, with catches in Canadian waters typically consisting of fish greater than four years old. Separate, and much smaller, populations of Pacific Hake occurring in the major inlets of the Northeast Pacific Ocean, including the Strait of Georgia, Puget Sound, and the Gulf of California, are not included in this analysis.

## Catches



Figure a. Total Pacific Hake catch used in the assessment by sector, 1966-2023. U.S. tribal catches are included in the sectors where they are represented.

Coast-wide fishery landings of Pacific Hake averaged 243,288 t from 1966 to 2023, with a low of $89,930 \mathrm{t}$ in 1980 and a peak of $440,849 \mathrm{t}$ in 2017 (Figure a). Prior to 1966, total removals were negligible compared to the modern fishery. Over the early period (1966-1990) most removals were from foreign or joint-venture fisheries. Across the time series, annual catch in U.S. waters averaged $186,041 \mathrm{t}(76.5 \%$ of the total catch $)$, while catch from Canadian
waters averaged 57,247 t. Over the last 10 years, 2014-2023 (Table a), the average coastwide catch was $338,606 \mathrm{t}$ with U.S. and Canadian catches averaging $275,957 \mathrm{t}$ and $62,648 \mathrm{t}$, respectively. Since 2017, the coast-wide catch has been declining annually through 2023 to $263,981 \mathrm{t}$ out of a total allowable catch (TAC, adjusted for carryovers) of $625,000 \mathrm{t}$. Attainment in the U.S. was $52.1 \%$ of its quota and in Canada it was $14.4 \%$.

Table a. Total Pacific Hake catch used in the assessment by sector for the most recent ten years. U.S. tribal catches are included in the sectors where they are represented.

| Year | U.S. <br> Mother- <br> ship | U.S. <br> Catcher- <br> processor | U.S. <br> Shore- | U.S. <br> besearch | U.S. <br> Total | Canada <br> Joint- <br> venture | Canada <br> Shore- <br> side | Canada <br> Freezer- <br> trawler | Canada <br> Total |
| :--- | :---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 2014 | 62,102 | 103,203 | 98,640 | 197 | 264,141 | 0 | 13,326 | 21,792 | 35,118 |
| Total |  |  |  |  |  |  |  |  |  |
| 2015 | 27,665 | 68,484 | 58,011 | 0 | 154,160 | 0 | 16,775 | 22,909 | 39,684 |
| 2016 | 65,036 | 108,786 | 87,760 | 745 | 262,327 | 0 | 35,012 | 34,731 | 69,743 |
| 2017 | 66,428 | 136,960 | 150,741 | 0 | 354,129 | 5,608 | 43,427 | 37,686 | 86,721 |
| 2018 | 67,121 | 116,073 | 135,112 | 0 | 318,306 | 2,724 | 50,747 | 41,942 | 95,413 |
| 2019 | 52,646 | 116,146 | 148,210 | 0 | 317,002 | 0 | 40,794 | 54,218 | 95,013 |
| 2020 | 37,978 | 111,147 | 138,688 | 95 | 287,908 | 0 | 312,019 |  |  |
| 2021 | 35,208 | 104,030 | 129,319 | 917 | 269,473 | 0 | 11,269 | 62,404 | 92,489 |
| 2022 | 59,516 | 126,247 | 105,939 | 0 | 291,702 | 0 | 3,868 | 27,807 | 57,076 |
| 2023 | 32,911 | 107,117 | 100,396 | 0 | 240,424 | 0 | 3,657 | 19,901 | $23,397,557$ |

In this document, the terms catch and landings are used interchangeably. Estimates of discard within the target fishery are included but discarding of Pacific Hake in non-target fisheries is not. Discard from all fisheries, including those that do not target Pacific Hake, is estimated to be less than $1 \%$ of landings in recent years. During the last five years, catches were above the long-term average catch $(243,288 \mathrm{t})$ but have been declining over that period (especially in Canada). Landings between 2001 and 2008 were predominantly comprised of fish from the very large 1999 year class, with a cumulative removal (through 2023) from that cohort of $2.13 \mathrm{Mt}[1 \mathrm{Mt}=1$ megatonne $=1$ million metric tonnes]. Through 2023, the cumulative catches of the 2010, 2014, and 2016 year classes were $2.56 \mathrm{Mt}, 1.76$ Mt , and 1.13 Mt , respectively. In the 2023 catch, the 2021 cohort was the largest ( $35 \%$ ), followed by the 2020 cohort ( $25 \%$ ), and then the 2016 cohort ( $13 \%$ ).

## Data and assessment

This Joint Technical Committee (JTC) assessment depends on fishery landings (1966-2023), an acoustic survey biomass index of age-2+ fish (Figure b) and age compositions (1995-2023), a relative index of age-1 fish (Figure c; 1995-2023), fishery age compositions (1975-2023), and mean weight-at-age data (1975-2023). In 2011 the survey biomass index was the lowest in the time series and was followed by the index increasing in 2012, 2013, and again in 2015 before decreasing to near the time series average in 2017. The survey shows a decline from 2019 (the fourth highest of the series) to 2023 (the third lowest of the series). Age-composition data from the aggregated fisheries and the acoustic survey, along with the age- 1 index, provide data that facilitates estimating relative cohort strength, i.e., strong and weak cohorts. The age- 1 index suggests particularly large numbers of age-1 fish in 2009, 2011, 2015, and 2021 (2008, 2010, 2014, and 2020 year classes, respectively), and is not available for most even years (odd year classes). There are no data to inform the size of the 2023 year class.


Figure b. Acoustic survey biomass index of age-2+ fish (Mt). Approximate $95 \%$ confidence intervals are based on sampling variability (intervals without the additional squid/Pacific Hake apportionment uncertainty included in 2009, black line).

The assessment uses a Bayesian estimation approach, sensitivity analyses, and retrospective investigations to evaluate the potential consequences of parameter uncertainty, alternative structural models, and historical performance of the assessment model, respectively. The Bayesian approach combines prior knowledge about natural mortality, stock-recruitment steepness (a parameter for stock productivity), and several other parameters, with likelihoods for the acoustic survey biomass index, acoustic survey age-composition data, the relative age- 1 index, and fishery age-composition data. Integrating the joint posterior distribution over model parameters provides probabilistic inferences about uncertain model parameters and forecasts derived from those parameters; this is done via Markov chain Monte Carlo sampling using the efficient No-U-Turn Sampler (NUTS). Sensitivity analyses are used to identify alternative model assumptions that may also be consistent with the data. All models, including bridging, sensitivity, and retrospective models, use a Bayesian framework for estimation. Retrospective analyses identify possible poor performance of the assessment model with respect to future predictions. Past assessments have conducted closed-loop simulations that provide insights into how alternative combinations of survey frequency, assessment model selectivity assumptions, changes in the distribution or Pacific Hake, and harvest control rules affect expected management outcomes given repeated application of these procedures over the long-term. The results of past (and ongoing) closed-loop simulations help inform decisions made for this assessment.


Figure c. Relative index of age-1 fish (numbers of fish) and approximate $95 \%$ confidence intervals based on sampling variability. The index is relative because the survey does not attempt to sample all available age- 1 fish and the analysis does not include kriging as is done to estimate age- $2+$ biomass.

This 2024 assessment retained the same general population dynamics structure as the base assessment model from 2023 and again is configured using Stock Synthesis. Updates in this assessment include incorporating the new biomass estimate and age-composition data from the acoustic survey conducted in 2023, fishery catch and age-composition data from 2023, weight-at-age data for 2023, the 2023 age- 1 index estimate, and minor changes to pre-2023 data. In addition, a new model-based approach was used to develop the input weight-at-age matrix, and time-varying temperature-dependent maturity was introduced to better inform fecundity.

This assessment continues to use (since 2014) time-varying (rather than constant) selectivity to maintain flexibility within the fishery dynamics given variability in Pacific Hake distribution patterns. The Dirichlet-multinomial estimation approach to weighting composition data was retained, and sensitivity to an alternative data-weighting approach was investigated. Time-varying fecundity, which was introduced in 2019, was retained and improved upon with time-varying estimates of maturity. Assumptions for the forecast period for weight at age and selectivity continue to be based on conditions during the last five years, as done since the 2020 assessment.

## Stock Biomass

Results from the base model indicate that since the 1960s, Pacific Hake female spawning biomass has ranged from well below to above unfished equilibrium (Figures d and e). Model estimates suggest that it was below the unfished equilibrium in the 1960s, at the start of the assessment period, due to lower than average recruitment.


Figure d. Median (solid line) of the posterior distribution for beginning of the year female spawning biomass ( $B_{t}$ in year $t$; Mt) through 2024 with $95 \%$ posterior credibility intervals (shaded area). The left-most circle with a $95 \%$ posterior credibility interval is the estimated unfished equilibrium biomass, $B_{0}$.

The stock is estimated to have increased rapidly and was above unfished equilibrium in the mid-1970s and mid-1980s (after two large recruitment events in the early 1980s). It then declined steadily to a low in 1999. This was followed by a brief increase to a peak in 2003 as the very large 1999 year class matured. The 1999 year class largely supported the fishery for several years due to relatively small recruitment events between 2000 and 2007. With the aging 1999 year class, median female spawning biomass declined throughout the late 2000s, reaching a time-series low of 0.616 Mt in 2009 . Median female spawning biomass is estimated to have peaked again in 2014 due to a very large 2010 year class and an above-average 2008 year class. The subsequent decline from 2014 to 2016 is primarily from the 2010 year class surpassing the age at which the gains in weight from growth are greater than the losses in weight from mortality (growth-mortality transition). The 2014 year class is estimated to be large, though not as large as the 1999 and 2010 year classes, increasing the biomass in 2017. The estimated biomass mostly declined from 2018 to 2022
due to the 2014 and 2016 year classes moving through the growth-mortality transition during a period of high catches. The increase in female spawning biomass in 2023 and 2024 is due to the expected above-average 2020 and potentially large 2021 cohorts entering maturity and the recent declining trend in catch.

The median estimate of the 2024 relative spawning biomass (spawning biomass at the start of 2024 divided by that at unfished equilibrium, $B_{0}$ ) is $99 \%$. However, the uncertainty is large, with a $95 \%$ posterior credibility interval from $45 \%$ to $230 \%$ (Table b), partly due to remaining unknowns about the size of the potentially large 2021 cohort because the acoustic survey has only provided one year of information about it.
The median estimate of the 2024 female spawning biomass is 1.885 (with a $95 \%$ posterior credibility interval from 0.853 to 4.828 ) Mt. The current estimate of the 2023 female spawning biomass is $1.335(0.652-3.225) \mathrm{Mt}$, giving less uncertainty than the estimate from the 2023 assessment of $1.910(0.757-5.610) \mathrm{Mt}$. The current median is reduced from that in the 2023 assessment due to the tail of the distribution being greatly curtailed and a slight lowering of the lower end of the interval. The decrease appears to be due to the addition of the age- $2+$ biomass index pulling down the estimated biomass for recent years, plus the addition of the survey age compositions lowering the estimated 2020 recruitment.


Figure e. Median (solid line) of the posterior distribution for relative spawning biomass ( $B_{t} / B_{0}$ ) through 2024 with $95 \%$ posterior credibility intervals (shaded area). Dashed horizontal lines show $10 \%, 40 \%$, and $100 \%$ of the unfished equilibrium $\left(B_{0}\right)$.

Table b. Recent trends in estimated beginning of the year female spawning biomass (SB; kt) and SB relative to estimated SB at unfished equilibrium (Rel. SB; \%).

| Year | SB <br> $2 . .^{\text {th }}$ <br> percentile | SB <br> Median | SB <br> 97.5 <br> percentile | Rel. SB <br> $2.5^{\text {th }}$ <br> percentile | Rel. SB <br> Median | Rel. SB <br> 9.5.5 <br> percentile |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 2015 | $1,105.1$ | $1,447.8$ | $2,310.3$ | $50.0 \%$ | $76.9 \%$ | $119.8 \%$ |
| 2016 | 940.3 | $1,223.3$ | $1,954.2$ | $42.2 \%$ | $65.0 \%$ | $101.5 \%$ |
| 2017 | $1,248.0$ | $1,646.4$ | $2,720.8$ | $56.4 \%$ | $87.5 \%$ | $139.1 \%$ |
| 2018 | $1,259.3$ | $1,711.2$ | $2,948.4$ | $57.8 \%$ | $90.9 \%$ | $148.5 \%$ |
| 2019 | $1,004.3$ | $1,402.1$ | $2,500.9$ | $46.5 \%$ | $74.5 \%$ | $125.4 \%$ |
| 2020 | 911.0 | $1,349.6$ | $2,566.1$ | $43.3 \%$ | $71.5 \%$ | $125.4 \%$ |
| 2021 | 699.4 | $1,118.3$ | $2,268.8$ | $34.3 \%$ | $58.9 \%$ | $110.0 \%$ |
| 2022 | 627.0 | $1,116.3$ | $2,453.2$ | $31.7 \%$ | $58.6 \%$ | $118.5 \%$ |
| 2023 | 652.5 | $1,335.5$ | $3,224.8$ | $34.2 \%$ | $69.9 \%$ | $154.7 \%$ |
| 2024 | 853.2 | $1,884.9$ | $4,828.4$ | $45.0 \%$ | $98.7 \%$ | $229.8 \%$ |

## Recruitment

The addition of 2023 data changes the estimates of absolute recruitments for the most recent years, while the improved methods for modeling temporal weight-at-age and spatio-temporal maturity have slightly changed some historical estimated recruitments.

The estimate of 2020 recruitment in last year's assessment was based on only two years of data and thus was highly uncertain. It suggested the 2020 cohort could potentially be huge ( $95 \%$ credible interval: $2.9-47.6$ billion fish), but now with information from the age- $2+$ biomass index and survey age-composition data the 2020 cohort looks to be less but still above average ( $95 \%$ interval: $2.1-12.7$ billion fish). The median has consequently fallen from 11.4 to 4.7 billion fish between the two assessments.

The 2021 recruitment is estimated to be potentially large, whereas it was estimated to be below average in last year's assessment (with very limited data); the median has increased by 9.7 billion fish. The general notion remains that recent Pacific Hake recruitment is highly uncertain, and estimates for recent years (based on limited data) can change substantially.

Table c. Estimates of recent recruitment (millions of age-0 fish) and recruitment deviations, where deviations below (above) zero indicate recruitment below (above) that estimated from the stock-recruitment relationship.

| Year | $\begin{aligned} & \text { Recruit- } \\ & \text { ment } \\ & 2.5^{\text {th }} \\ & \text { percentile } \end{aligned}$ | Recruitment Median | $\begin{gathered} \text { Recruit- } \\ \text { ment } \\ 97.5^{\text {th }} \\ \text { percentile } \end{gathered}$ | Rec. Deviations $2.5^{\text {th }}$ percentile | Rec. Deviations Median | Rec. Deviations $97.5^{\text {th }}$ percentile |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2014 | 5,667.0 | 8,255.9 | 14,926.2 | 1.701 | 2.150 | 2.591 |
| 2015 | 8.5 | 34.2 | 122.2 | -4.674 | -3.335 | -2.134 |
| 2016 | 3,715.2 | 5,638.3 | 10,990.7 | 1.334 | 1.810 | 2.299 |
| 2017 | 848.7 | 1,565.0 | 3,553.1 | -0.127 | 0.499 | 1.152 |
| 2018 | 112.1 | 397.2 | 1,287.3 | -2.130 | -0.887 | 0.130 |
| 2019 | 46.9 | 273.3 | 1,015.1 | -2.959 | -1.264 | -0.063 |
| 2020 | 2,063.0 | 4,747.9 | 12,728.4 | 0.849 | 1.615 | 2.455 |
| 2021 | 4,085.1 | 10,187.3 | 29,499.4 | 1.556 | 2.394 | 3.281 |
| 2022 | 288.9 | 1,881.2 | 8,858.6 | -1.148 | 0.699 | 2.142 |
| 2023 | 42.6 | 979.1 | 20,271.6 | -3.038 | 0.014 | 3.018 |

Pacific Hake have low to moderate recruitment with occasional large year classes (Table c and Figure f). Very large year classes in 1980, 1984, and 1999 supported much of the commercial catch from the 1980s to the mid-2000s. From 2000 to 2007, estimated recruitment was at some of the lowest values in the time series but this was followed by an above average 2008 year class and a very strong 2010 year class. Above average year classes occurred in 2014 and 2016, which have been sustaining the fishery in recent years, with small year classes for all other years from 2011-2019 (median recruitment well below the mean of all median recruitments).

The 2020 cohort is estimated to be above average, and the 2021 cohort is estimated to be potentially large from limited fishery data and the 2023 survey. The 2022 cohort was observed by the age-1 index in 2023, suggesting it is average to below average in size.


Figure f. Medians (solid circles) and means (X) of the posterior distribution for recruitment (billions of age-0 fish) with $95 \%$ posterior credibility intervals (vertical lines). The median of the posterior distribution for mean unfished equilibrium recruitment $\left(R_{0}\right)$ is shown as the horizontal dashed line with the $95 \%$ posterior credibility interval shaded between the dotted lines.

There is no information in the data to estimate the sizes of the 2023 and 2024 year classes. Retrospective analyses of year class strength for young fish have shown the estimates of recent recruitment to be unreliable prior to at least a modelled age of 3 (observed as age- 2 fish the previous year) without a survey in the most recent year and a modelled age of two (observed as age-1 fish) with a survey.

## Default harvest policy

The default $F_{40 \%}-40: 10$ harvest policy prescribes the maximum rate of fishing mortality to equal $F_{\text {SPR }=40 \%}$. This rate gives a spawning potential ratio (SPR) of $40 \%$, meaning that the female spawning biomass per recruit with $F_{\text {SPR }=40 \%}$ is $40 \%$ of that without fishing. If female spawning biomass is below $B_{40 \%}\left(40 \%\right.$ of $\left.B_{0}\right)$, the policy reduces the TAC linearly until it equals zero at $B_{10 \%}\left(10 \%\right.$ of $\left.B_{0}\right)$. Relative fishing intensity for fishing rate $F$ is ( $1-$ $\operatorname{SPR}(F)) /\left(1-\mathrm{SPR}_{40 \%}\right)$, where $\mathrm{SPR}_{40 \%}$ is an SPR of $40 \%$; it is reported here interchangeably as a proportion or a percentage. A relative fishing intensity above 1.0 means fishing at a rate above $F_{\mathrm{SPR}=40 \%}$.

Table d. Recent estimates of relative fishing intensity, ( $1-\mathrm{SPR}$ )/( $1-\mathrm{SPR}_{40 \%}$ ), and exploitation fraction (catch divided by age- $2+$ biomass).

| Year | Rel. <br> Fishing Intensity $2.5^{\text {th }}$ percentile | Rel. Fishing Intensity Median | Rel. <br> Fishing Intensity $97.5^{\text {th }}$ percentile | Exploit. Fraction $2.5^{\text {th }}$ percentile | Exploit. <br> Fraction Median | Exploit. Fraction $97.5^{\text {th }}$ percentile |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2014 | 0.393 | 0.623 | 0.858 | 0.045 | 0.071 | 0.094 |
| 2015 | 0.264 | 0.455 | 0.666 | 0.042 | 0.067 | 0.087 |
| 2016 | 0.477 | 0.736 | 0.983 | 0.057 | 0.093 | 0.122 |
| 2017 | 0.504 | 0.793 | 1.135 | 0.077 | 0.128 | 0.168 |
| 2018 | 0.441 | 0.722 | 1.030 | 0.060 | 0.104 | 0.142 |
| 2019 | 0.497 | 0.803 | 1.078 | 0.067 | 0.122 | 0.173 |
| 2020 | 0.363 | 0.625 | 0.873 | 0.072 | 0.136 | 0.203 |
| 2021 | 0.353 | 0.624 | 0.877 | 0.072 | 0.145 | 0.232 |
| 2022 | 0.330 | 0.618 | 0.914 | 0.052 | 0.116 | 0.213 |
| 2023 | 0.267 | 0.551 | 0.872 | 0.026 | 0.066 | 0.138 |

## Exploitation status

The median estimated relative fishing intensity on the stock is below the management level of 1.0 for all years (see Table d for recent years and Figure g).

Over the last five years, it was the highest in 2019 at $80.3 \%$, dropped in 2020 to $62.5 \%$, then remained stable for 2021 and 2022 at $62.4 \%$ and $61.8 \%$ respectively. The 3-year stable trend from 2020-2022 was ended in 2023 when the relative fishing intensity dropped to 55.1\% (Table d and Figure g).

The median exploitation fraction (catch divided by biomass of fish of age-2 and above) peaked in 1999 and 2008 (Figure h). The median exploitation fraction has decreased from a recent high in 2021 of 0.15 to 0.07 in 2023, which is a comparable level to 10 years ago (Table d and Figure h).

Although there is a considerable amount of imprecision around these estimates due to uncertainty in recruitment and spawning biomass, the $95 \%$ posterior credibility interval of relative fishing intensity was below 1.0 from 2012-2016 and again in 2020-2023 (Figure g).

## Management performance

Over the last decade (2014-2023), the mean coast-wide utilization rate (proportion of catch target removed) has been $63.5 \%$ and catches have been below coast-wide targets (Table e). From 2019 to 2023, the mean utilization rates differed between the United States ( $67.4 \%$ ) and Canada ( $48.1 \%$ ), though Canada's was higher than the U.S.'s in 2020. From 2020 the Canadian rate steadily declined to a time-series low of $14.4 \%$ in 2023, and the U.S. rate fell to $52.1 \%$ in 2023 . The usual $73.88 \%$ and $26.12 \%$ allocation of coast-wide TAC, as specified in the Joint U.S.-Canada Agreement for Pacific Hake, was implemented in 2022 and 2023. Total landings last exceeded the coast-wide quota in 2002 when utilization was $112 \%$.


Figure g. Trend in median relative fishing intensity (relative to the $F_{\mathrm{SPR}=40 \%}$ management level) through 2023 with $95 \%$ posterior credibility intervals. The $F_{\mathrm{SPR}=40 \%}$ management level defined in the Joint U.S.-Canada Agreement for Pacific Hake is shown as a horizontal line at 1.0.


Figure h. Trend in median exploitation fraction (catch divided by age-2+ biomass) through 2023 with $95 \%$ posterior credibility intervals.

Table e. Recent trends in Pacific Hake landings and management decisions. Catch targets in 2020 and 2021 were specified unilaterally. All landings and catch targets are given in tonnes.

| Year | U.S. <br> landings | Canada landings | Total landings |  | Canada prop. of total catch | U.S. catch target | Canada catch target | Total catch target | U.S. prop. of catch target removed | Canada prop. of catch target removed | Total prop. of catch target removed |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2014 | 264,141 | 35,118 | 299,259 | 88.3\% | 11.7\% | 316,206 | 111,794 | 428,000 | 83.5\% | 31.4\% | 69.9\% |
| 2015 | 154,160 | 39,684 | 193,844 | 79.5\% | 20.5\% | 325,072 | 114,928 | 440,000 | 47.4\% | 34.5\% | 44.1\% |
| 2016 | 262,327 | 69,743 | 332,070 | 79.0\% | 21.0\% | 367,553 | 129,947 | 497,500 | 71.4\% | 53.7\% | 66.7\% |
| 2017 | 354,129 | 86,721 | 440,849 | 80.3\% | 19.7\% | 441,433 | 156,067 | 597,500 | 80.2\% | 55.6\% | 73.8\% |
| 2018 | 318,306 | 95,413 | 413,719 | 76.9\% | 23.1\% | 441,433 | 156,067 | 597,500 | 72.1\% | 61.1\% | 69.2\% |
| 2019 | 317,002 | 95,013 | 412,015 | 76.9\% | 23.1\% | 441,433 | 156,067 | 597,500 | 71.8\% | 60.9\% | 69.0\% |
| 2020 | 287,908 | 92,489 | 380,397 | 75.7\% | 24.3\% | 424,810 | 104,480 | 529,290 | 67.8\% | 88.5\% | 71.9\% |
| 2021 | 269,473 | 57,076 | 326,549 | 82.5\% | 17.5\% | 369,400 | 104,480 | 473,880 | 72.9\% | 54.6\% | 68.9\% |
| 2022 | 291,702 | 31,671 | 323,372 | 90.2\% | 9.8\% | 402,646 | 142,354 | 545,000 | 72.4\% | 22.2\% | 59.3\% |
| 2023 | 240,424 | 23,557 | 263,981 | 91.1\% | 8.9\% | 461,750 | 163,250 | 625,000 | 52.1\% | 14.4\% | 42.2\% |



Figure i. Estimated historical path of median relative spawning biomass in year $t$ and corresponding median relative fishing intensity in year $t-1$. Labels show the time series start and end years; labels correspond to year $t$ (i.e., year of the relative spawning biomass). Gray bars span the $95 \%$ credibility intervals for 2024 relative spawning biomass (horizontal) and 2023 relative fishing intensity (vertical).

The median relative spawning biomass was above the $B_{40 \%}$ reference level in all years except 2007-2011 (Figures e and i), and the median relative fishing intensity has always been below 1.0 (Figure i). Relative spawning biomass increased from the lows in 2007-2012 with above average recruitment in 2008, 2010, 2014, 2016, and 2020. Correspondingly, median relative fishing intensity has remained below 1, and total catch has been declining since the time series high in 2017. While there is large uncertainty in the 2023 estimates of relative fishing intensity and relative spawning biomass, the model estimates a $0.2 \%$ joint probability of being both above the target relative fishing intensity ( $F_{\mathrm{SPR}=40 \%}$ ) in 2023 and below the relative spawning biomass level of $B_{40 \%}$ at the start of 2024.

## Reference points

The term 'reference points' is used throughout this document to describe common conceptual summary metrics (Table f). The Agreement specifically identifies $F_{\text {SPR }=40 \%}$ as the default harvest rate and $B_{40 \%}$ as a point where the 40:10 TAC adjustment is triggered (see the Glossary in Appendix C). The medians of sustainable yields and biomass reference points are similar to what was reported in the 2023 assessment. The probability that female spawning biomass at the beginning of 2024 is below $B_{40 \%}$ is $\mathrm{P}\left(B_{2024}<B_{40 \%}\right)=1.3 \%$, and of being below $B_{25 \%}$ is $\mathrm{P}\left(B_{2024}<B_{25 \%}\right)=0.1 \%$. The probability that the relative fishing intensity was above the $F_{\mathrm{SPR}=40 \%}$ level of 1.0 at the end of 2023 is $0.4 \%$.

Table f. Summary of median and $95 \%$ credibility intervals of equilibrium conceptual reference points for the base assessment model. Equilibrium reference points were computed using 1975-2023 averages for mean weight-at-age and baseline selectivity-at-age (1966-1990; prior to time-varying deviations). Dashes (-) indicate values that are static at one value and do not have a credible interval associated with them.

| Quantity | 2.5\% | Median | 97.5\% |
| :---: | :---: | :---: | :---: |
| Unfished female spawning biomass ( $B_{0}, \mathrm{kt}$ ) | 1,235 | 1,919 | 3,132 |
| Unfished recruitment ( $R_{0}$, millions) | 1,394 | 2,600 | 5,383 |
| Reference points (equilibrium) based on $F_{\text {SPR }=40 \%}$ |  |  |  |
| Female spawning biomass at $F_{\text {SPR }=40 \%}\left(B_{\text {SPR }}=40 \%, \mathrm{kt}\right)$ | 409 | 681 | 1,127 |
| SPR at $F_{\text {SPR }=40 \%}$ | - | 40\% | - |
| Exploitation fraction corresponding to $F_{\text {SPR }}=40 \%$ | 16.3\% | 19.1\% | 22.0\% |
| Yield associated with $F_{\text {SPR }=40 \% \text { ( }}(\mathrm{kt})$ | 180 | 317 | 594 |
| Reference points (equilibrium) based on $B_{40 \%}\left(40 \%\right.$ of $\left.B_{0}\right)$ |  |  |  |
| Female spawning biomass ( $B_{40 \%}$, kt ) | 494 | 767 | 1,253 |
| SPR at $\mathrm{B}_{40} \%$ | 40.7\% | 43.5\% | 50.8\% |
| Exploitation fraction resulting in $B_{40 \%}$ | 12.9\% | 16.8\% | 20.2\% |
| Yield at $B_{40 \%}$ (kt) | 177 | 309 | 580 |
| Reference points (equilibrium) based on estimated MSY |  |  |  |
| Female spawning biomass ( $B_{\text {MSY }}, \mathrm{kt}$ ) | 297 | 490 | 867 |
| SPR at MSY | 22.8\% | 29.6\% | 45.1\% |
| Exploitation fraction corresponding to SPR at MSY | 15.8\% | 27.0\% | 36.5\% |
| MSY (kt) | 188 | 336 | 639 |

## Unresolved problems and major uncertainties

Measures of uncertainty in the base model underestimate the total uncertainty in the current stock status and projections because they do not account for possible alternative structural models for hake population dynamics and fishery processes (e.g., selectivity) and the scientific basis for prior probability distributions. To address such structural uncertainties, we performed sensitivity analyses to investigate a range of alternative assumptions and present the key ones in the main document.

The Pacific Hake stock displays high recruitment variability relative to other West Coast groundfish stocks, resulting in large and rapid biomass changes. This leads to a dynamic fishery that potentially targets strong cohorts and results in time-varying fishery selectivity. This volatility results in a high level of uncertainty in estimates of current stock status and stock projections because, with limited data to estimate incoming recruitment, the cohorts are fished before the assessment can accurately determine how big they are (i.e., cohort strength is typically not well known until it is observed by the fishery and survey, typically at a minimum age of three). While the addition of the age- 1 index helps inform recent recruitment, the survey is conducted every other year and does not directly address current or future recruitment expectations. In particular, while the model estimates the 2020 and 2021 cohorts as above average in size, their absolute size remains highly uncertain. This uncertainty propagates directly into current and forecasted estimates of female spawning biomass. The 2023 acoustic survey provided additional information on the size of the 2020 year-class (as well as informed the 2021 year class), and the 2023 age- 1 index helped inform the size of the 2022 year class. Collectively, these lessened uncertainty around estimates of female spawning biomass. Further, the interactions among variance parameters that govern variability in fishery selectivity and recruitment parameters through time, as well as those used in relative data weighting, are not well understood and could propagate uncertainty beyond what is presented in this assessment.

## Forecast decision tables

The catch limit for 2024 based on the default $F_{40 \%}-40: 10$ harvest policy has a median of $747,588 \mathrm{t}$ with a wide range of uncertainty, the $95 \%$ credibility interval being 298,355-2,124,832 t.

Decision tables give the projected population status (relative spawning biomass) and fishing intensity relative to the target under different catch alternatives for the base model (Tables g and h ). The tables are organized to show the projected outcome for each potential catch level and year (row) across the quantiles (columns) of the posterior distribution. Tables show results for up to three years of future catch levels based on subsequent estimates of stock status and fishing intensity. Figure j shows the projected relative spawning biomass for several of the catch alternatives. Population dynamics and governing parameters assumed during the forecast period include random recruitment; selectivity, weight-at-age and fecundity averaged over the five most recent years (2019-2023); and constant values for all other parameters.
A relative fishing intensity of 1 should indicate fishing at the $F_{\mathrm{SPR}=40 \%}$ default harvest rate catch target. But, the projected median relative fishing intensity can be slightly different than the target because the $F_{\mathrm{SPR}=40 \%}$ default harvest-rate catch limit is calculated using baseline selectivity-at-age (1966-1990; prior to time-varying deviations), whereas the forecasted catches are removed using selectivity averaged over the last five years. Recent changes in selectivity will thus be reflected in the determination of fishing relative to the default harvest policy. For example, fishing at the $F_{\mathrm{SPR}=40 \%}$ default harvest-rate catch limit (scenario n: default HR) in 2024 results in a median relative fishing intensity of 0.94 (Table h).

Table g. Forecast quantiles of Pacific Hake relative spawning biomass at the beginning of the year. Catch alternatives are defined by letters a-o and are a constant value across all forecasted years unless otherwise defined in the first column. Acronyms are defined in the glossary (Appendix C).

| Catch alternative |  |  | Biomass at start of year | Relative spawning biomass |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Catch year | Catch (t) |  | 5\% | 50\% | 95\% |
|  |  |  | Start of 2024 | 0.51 | 0.99 | 2.01 |
| a: | 2024 | 0 | Start of 2025 | 0.57 | 1.11 | 2.23 |
|  | 2025 | 0 | Start of 2026 | 0.59 | 1.13 | 2.35 |
|  | 2026 | 0 | Start of 2027 | 0.57 | 1.12 | 2.45 |
| b: | 2024 | 180,000 | Start of 2025 | 0.53 | 1.06 | 2.18 |
|  | 2025 | 180,000 | Start of 2026 | 0.50 | 1.04 | 2.26 |
|  | 2026 | 180,000 | Start of 2027 | 0.46 | 1.00 | 2.32 |
| c: | 2024 | 225,000 | Start of 2025 | 0.52 | 1.05 | 2.16 |
|  | 2025 | 225,000 | Start of 2026 | 0.48 | 1.02 | 2.23 |
|  | 2026 | 225,000 | Start of 2027 | 0.43 | 0.97 | 2.29 |
| d: | 2024 | 320,000 | Start of 2025 | 0.50 | 1.02 | 2.14 |
| $10 \%$ reduction | 2025 | 288,000 | Start of 2026 | 0.45 | 0.98 | 2.20 |
| each year | 2026 | 259,200 | Start of 2027 | 0.39 | 0.93 | 2.24 |
| e: | 2024 | 264,000 | Start of 2025 | 0.51 | 1.04 | 2.15 |
| 2023 catch | 2025 | 264,000 | Start of 2026 | 0.47 | 1.00 | 2.21 |
|  | 2026 | 264,000 | Start of 2027 | 0.41 | 0.94 | 2.26 |
| f: | 2024 | 350,000 | Start of 2025 | 0.49 | 1.01 | 2.13 |
|  | 2025 | 350,000 | Start of 2026 | 0.42 | 0.96 | 2.17 |
|  | 2026 | 350,000 | Start of 2027 | 0.35 | 0.88 | 2.20 |
| g: | 2024 | 350,000 | Start of 2025 | 0.49 | 1.01 | 2.13 |
| $10 \%$ reduction | 2025 | 315,000 | Start of 2026 | 0.43 | 0.97 | 2.18 |
| each year | 2026 | 283,500 | Start of 2027 | 0.37 | 0.91 | 2.23 |
| h: | 2024 | 380,000 | Start of 2025 | 0.49 | 1.01 | 2.12 |
|  | 2025 | 380,000 | Start of 2026 | 0.41 | 0.94 | 2.16 |
|  | 2026 | 380,000 | Start of 2027 | 0.33 | 0.86 | 2.17 |
| i: | 2024 | 380,000 | Start of 2025 | 0.49 | 1.01 | 2.12 |
| 10\% reduction | 2025 | 342,000 | Start of 2026 | 0.42 | 0.95 | 2.17 |
| each year | 2026 | 307,800 | Start of 2027 | 0.36 | 0.89 | 2.21 |
| j: | 2024 | 430,000 | Start of 2025 | 0.47 | 0.99 | 2.11 |
|  | 2025 | 430,000 | Start of 2026 | 0.39 | 0.92 | 2.14 |
|  | 2026 | 430,000 | Start of 2027 | 0.30 | 0.83 | 2.13 |
| k: | 2024 | 545,000 | Start of 2025 | 0.45 | 0.96 | 2.08 |
| 2022 TAC | 2025 | 545,000 | Start of 2026 | 0.33 | 0.86 | 2.08 |
|  | 2026 | 545,000 | Start of 2027 | 0.22 | 0.75 | 2.05 |
| 1: | 2024 | 625,000 | Start of 2025 | 0.43 | 0.94 | 2.06 |
| 2023 TAC | 2025 | 625,000 | Start of 2026 | 0.30 | 0.83 | 2.03 |
|  | 2026 | 625,000 | Start of 2027 | 0.18 | 0.70 | 1.99 |
| m: | 2024 | 875,262 | Start of 2025 | 0.37 | 0.88 | 1.99 |
| Fishing intensity | 2025 | 861,614 | Start of 2026 | 0.22 | 0.71 | 1.91 |
| at 100\% | 2026 | 782,426 | Start of 2027 | 0.13 | 0.57 | 1.86 |
| n : | 2024 | 747,588 | Start of 2025 | 0.40 | 0.91 | 2.02 |
| Default HR | 2025 | 772,111 | Start of 2026 | 0.24 | 0.76 | 1.97 |
| $\left(F_{\text {SPR }=40 \%}-40: 10\right)$ | 2026 | 717,464 | Start of 2027 | 0.14 | 0.62 | 1.91 |
| o: | 2024 | 767,382 | Start of 2025 | 0.39 | 0.90 | 2.02 |
| Equal catch | 2025 | 767,382 | Start of 2026 | 0.24 | 0.76 | 1.96 |
| $\left(\mathrm{C}_{2024} \approx \mathrm{C}_{2025}\right)$ | 2026 | 712,782 | Start of 2027 | 0.14 | 0.62 | 1.91 |

Table h. Forecast quantiles of Pacific Hake relative fishing intensity ( $1-\mathrm{SPR}$ )/( $1-\mathrm{SPR}_{40 \%}$ ), expressed as a proportion. Catch alternatives are defined by letters a-o and are a constant value across all forecasted years unless otherwise defined in the first column. Acronyms are defined in the glossary (Appendix C).

| Catch alternative |  |  | Relative fishing intensity |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Catch year | Catch (t) | 5\% | 50\% | 95\% |
| a: | 2024 | 0 | 0.00 | 0.00 | 0.00 |
|  | 2025 | 0 | 0.00 | 0.00 | 0.00 |
|  |  | 0 | 0.00 | 0.00 | 0.00 |
| b: | 2024 | 180,000 | 0.22 | 0.43 | 0.69 |
|  | 2025 | 180,000 | 0.18 | 0.37 | 0.63 |
|  | 2026 | $180,000$ | 0.16 | 0.33 | 0.59 |
| c: | 2024 | 225,000 | 0.27 | 0.50 | 0.78 |
|  | 2025 | 225,000 | 0.22 | 0.44 | 0.72 |
|  | 2026 | 225,000 | 0.20 | 0.40 | 0.69 |
| d: <br> $10 \%$ reduction each year | 2024 | 320,000 | 0.35 | 0.62 | 0.91 |
|  | 2025 | 288,000 | 0.27 | 0.53 | 0.83 |
|  | 2026 | 259,200 | 0.23 | 0.46 | 0.78 |
| e: | 2024 | 264,000 | 0.30 | 0.55 | 0.84 |
| 2023 catch | 2025 | 264,000 | 0.25 | 0.49 | 0.79 |
|  | 2026 | 264,000 | 0.23 | 0.46 | 0.77 |
| f: | 2024 | 350,000 | 0.38 | 0.66 | 0.94 |
|  | 2025 | 350,000 | 0.32 | 0.60 | 0.92 |
|  | 2026 | 350,000 | 0.29 | 0.57 | 0.92 |
| $\begin{gathered} \text { g: } \\ 10 \% \text { reduction } \\ \text { each year } \end{gathered}$ | 2024 | 350,000 | 0.38 | 0.66 | 0.94 |
|  | 2025 | 315,000 | 0.30 | 0.56 | 0.88 |
|  | 2026 | 283,500 | 0.25 | 0.49 | 0.83 |
| h : | 2024 | 380,000 | 0.40 | 0.69 | 0.97 |
|  | 2025 | 380,000 | 0.34 | 0.63 | 0.96 |
|  | 2026 | 380,000 | 0.31 | 0.60 | 0.97 |
| i:$10 \%$ reductioneach year | 2024 | 380,000 | 0.40 | 0.69 | 0.97 |
|  | 2025 | 342,000 | 0.32 | 0.59 | 0.92 |
|  | 2026 | 307,800 | 0.26 | 0.52 | 0.88 |
| j: | 2024 | 430,000 | 0.44 | 0.73 | 1.02 |
|  | 2025 | 430,000 | 0.38 | 0.68 | 1.02 |
|  | 2026 | 430,000 | 0.35 | 0.66 | 1.05 |
| $\begin{gathered} \text { k: } \\ 2022 \text { TAC } \end{gathered}$ | 2024 | 545,000 | 0.51 | 0.82 | 1.11 |
|  | 2025 | 545,000 | 0.45 | 0.78 | 1.13 |
|  | 2026 | 545,000 | 0.42 | 0.78 | 1.20 |
| $\begin{gathered} \text { 1: } \\ 2023 \text { TAC } \end{gathered}$ | 2024 | 625,000 | 0.56 | 0.87 | 1.16 |
|  | 2025 | 625,000 | 0.50 | 0.85 | 1.20 |
|  | 2026 | 625,000 | 0.47 | 0.85 | 1.26 |
| Fishing intensity at $100 \%$ |  | 875,262 | 0.68 | 1.00 | 1.27 |
|  | 2025 | 861,614 | 0.62 | 1.00 | 1.30 |
|  | 2026 | 782,426 | 0.57 | 1.00 | 1.31 |
| n: <br> Default HR <br> ( $F_{\text {SPR }=40 \%}-40: 10$ ) | 2024 | 747,588 | 0.62 | 0.94 | 1.22 |
|  | 2025 | 772,111 | 0.58 | 0.94 | 1.28 |
|  | 2026 | 717,464 | 0.53 | 0.94 | 1.30 |
| o:Equal catch$\left(\mathrm{C}_{2024} \approx \mathrm{C}_{2025}\right)$ | 2024 | 767,382 | 0.63 | 0.95 | 1.23 |
|  | 2025 | 767,382 | 0.58 | 0.94 | 1.28 |
|  | 2026 | 712,782 | 0.53 | 0.94 | 1.30 |

Management metrics that were identified as important to the Joint Management Committee and the Advisory Panel in 2012 are presented for 2025, 2026, and 2027 projections (Tables i, j, and $k$; Figures $k, 1$, and $m$ ). These metrics summarize the probability of various outcomes from the base model given each potential management action. Although not linear, probabilities can be interpolated from these results for intermediate catch values in 2024 (Table i and Figure k). However, interpolation is not appropriate for all catches in 2025 or 2026 because they are conditional on previous year(s) catch levels. This explains why probabilities can sometimes decline (rather than rise) with increased 2025 and 2026 catch levels (Tables j and k; Figures 1 and m).


Figure j. Median and $95 \%$ posterior credibility intervals of estimated relative spawning biomass to the start of 2024 from the base model and projections to the start of 2027 for several management actions, which are defined in Table g.

With zero catch for the next three years, the biomass has a $3 \%$ probability of decreasing from 2024 to 2025 (Table i; Figure j), a 59\% probability of decreasing from 2025 to 2026 (Table j), and a $66 \%$ probability of decreasing from 2026 to 2027 (Table k).

The probability of the female spawning biomass decreasing from 2024 to 2025 is $36 \%$ for a 2024 catch level similar to that for 2023 (scenario e: 2023 catch). For all explored catches, the maximum probability of female spawning biomass at the start of 2025 dropping below $B_{10 \%}$ is $0.0 \%$ and of dropping below $B_{40 \%}$ is $6.7 \%$ (Table $i$ and Figure k). As the above average 2014 and 2016 cohorts continue to age, total biomass of these cohorts even without fishing mortality is expected to decrease as losses from mortality outweigh increases from growth. The estimated above-average (yet still highly uncertain) 2020 and 2021 cohorts
will continue to play a large role in determining female spawning biomass during the forecast years presented here.


Figure k. Graphical representation of the probabilities related to spawning biomass, relative fishing intensity, and the 2025 default harvest policy catch for alternative 2024 catch options (explained in Table g) as listed in Table i. The symbols indicate points that were computed directly from model output and lines interpolate between the points.

Table i. Probabilities related to spawning biomass, relative fishing intensity, and the 2025 default harvest policy catch for alternative 2024 catch options (catch options are explained in Table g).

|  | Catch (t) <br> in 2024 | $\mathrm{B}_{\mathbf{2 0 2 5}}$ <br> $<\mathrm{B}_{\mathbf{2 0 2 4}}$ | $\mathrm{B}_{\mathbf{2 0 2 5}}$ <br> $<\mathbf{B}_{\mathbf{4 0 \%}}$ | $\mathrm{B}_{\mathbf{2 0 2 5}}$ <br> $<\mathbf{B}_{\mathbf{2 5} \%}$ | $\mathrm{B}_{\mathbf{2 0 2 5}}$ <br> $<\mathbf{B}_{\mathbf{1 0 \%}}$ | $\mathbf{2 0 2 4}$ <br> Fishing <br> intensity <br> $>\mathbf{1 0 0 \%}$ | 2025 <br> Default HR <br> catch <br> $\mathbf{2 0 2 4}$ <br> catch |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| a: | 0 | 0.03 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| b: | 180,000 | 0.22 | 0.01 | 0.00 | 0.00 | 0.00 | 0.00 |
| c: | 225,000 | 0.29 | 0.01 | 0.00 | 0.00 | 0.00 | 0.00 |
| d: | 320,000 | 0.44 | 0.02 | 0.00 | 0.00 | 0.02 | 0.03 |
| e: | 264,000 | 0.36 | 0.01 | 0.00 | 0.00 | 0.00 | 0.01 |
| f: | 350,000 | 0.49 | 0.02 | 0.00 | 0.00 | 0.03 | 0.04 |
| g: | 350,000 | 0.49 | 0.02 | 0.00 | 0.00 | 0.03 | 0.04 |
| h: | 380,000 | 0.53 | 0.02 | 0.00 | 0.00 | 0.04 | 0.06 |
| i: | 380,000 | 0.53 | 0.02 | 0.00 | 0.00 | 0.04 | 0.06 |
| j: | 430,000 | 0.60 | 0.02 | 0.00 | 0.00 | 0.07 | 0.10 |
| k: | 545,000 | 0.71 | 0.03 | 0.00 | 0.00 | 0.16 | 0.23 |
| l: | 625,000 | 0.76 | 0.04 | 0.00 | 0.00 | 0.24 | 0.33 |
| m: | 875,262 | 0.87 | 0.07 | 0.01 | 0.00 | 0.50 | 0.61 |
| n: | 747,588 | 0.83 | 0.05 | 0.01 | 0.00 | 0.37 | 0.48 |
| Continued on next page $\ldots$ |  |  |  |  |  |  |  |

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|  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{aligned} & \text { Catch }(t) \\ & \text { in } 2024 \end{aligned}$ | $\begin{gathered} \mathrm{B}_{2025} \\ <\mathrm{B}_{2024} \end{gathered}$ | $\begin{aligned} & \mathrm{B}_{2025} \\ & <B_{40 \%} \end{aligned}$ | $\begin{aligned} & { }_{2}^{B_{2025}} \\ & <B_{25 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2025} \\ & <B_{10 \%} \end{aligned}$ | 2024 <br> Fishing intensity $>100 \%$ | $\begin{aligned} & 2025 \\ & \text { Default HR } \\ & \text { cath } \\ & >2024 \\ & \text { catch } \end{aligned}$ |
| o: | 767,382 | 0.83 | 0.05 | 0.01 | 0.00 | 0.39 | 0.50 |



Figure 1. Graphical representation of the probabilities related to spawning biomass, relative fishing intensity, and the 2026 default harvest policy catch for alternative 2025 catch options (including associated 2024 catch; catch options explained in Table g) as listed in Table j. The symbols indicate points that were computed directly from model output and lines interpolate between the points.

Table j. Probabilities related to spawning biomass, relative fishing intensity, and the 2026 default harvest policy catch for alternative 2025 catch options, given the 2024 catch level shown in Table i (catch options are explained in Table g).

|  | $\begin{aligned} & \text { Catch }(\mathrm{t}) \\ & \text { in } 2025 \end{aligned}$ | $\begin{gathered} \mathrm{B}_{2026} \\ <\mathrm{B}_{2025} \end{gathered}$ | $\begin{aligned} & \mathrm{B}_{2026} \\ & <B_{40 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2026} \\ & <B_{25 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2026} \\ & <B_{10 \%} \end{aligned}$ | 2025 <br> Fishing intensity $>100 \%$ | $\begin{gathered} 2026 \\ \text { Default HR } \\ \text { catch } \\ >2025 \\ \text { catch } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| a: | 0 | 0.59 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| b : | 180,000 | 0.70 | 0.02 | 0.00 | 0.00 | 0.00 | 0.00 |
| c: | 225,000 | 0.72 | 0.02 | 0.00 | 0.00 | 0.00 | 0.00 |
| d: | 288,000 | 0.75 | 0.03 | 0.00 | 0.00 | 0.01 | 0.01 |
| e: | 264,000 | 0.74 | 0.03 | 0.00 | 0.00 | 0.00 | 0.01 |
| f: | 350,000 | 0.77 | 0.04 | 0.00 | 0.00 | 0.02 | 0.04 |
| g : | 315,000 | 0.76 | 0.03 | 0.00 | 0.00 | 0.01 | 0.03 |
| h: | 380,000 | 0.78 | 0.05 | 0.01 | 0.00 | 0.03 | 0.06 |
| i: | 342,000 | 0.76 | 0.04 | 0.01 | 0.00 | 0.02 | 0.04 |
| j: | 430,000 | 0.79 | 0.06 | 0.01 | 0.00 | 0.06 | 0.10 |
| k: | 545,000 | 0.82 | 0.09 | 0.02 | 0.00 | 0.15 | 0.25 |
| $1:$ | 625,000 | 0.84 | 0.11 | 0.03 | 0.00 | 0.23 | 0.35 |
| m : | 861,614 | 0.87 | 0.20 | 0.07 | 0.00 | 0.50 | 0.64 |
| n : | 772,111 | 0.86 | 0.16 | 0.06 | 0.00 | 0.39 | 0.54 |
| o : | 767,382 | 0.86 | 0.16 | 0.06 | 0.00 | 0.39 | 0.54 |



Figure m. Graphical representation of the probabilities related to spawning biomass, relative fishing intensity, and the 2027 default harvest policy catch for alternative 2026 catch options (including associated 2024 and 2025 catches; catch options explained in Table g) as listed in Table k. The symbols indicate points that were computed directly from model output and lines interpolate between the points.

Table k. Probabilities related to spawning biomass, relative fishing intensity, and the 2027 default harvest policy catch for alternative 2026 catch options, given the 2024 and 2025 catch levels shown in Tables i and j (catch options are explained in Table g).

|  | Catch ( t$)$ in 2026 | $\begin{gathered} \mathrm{B}_{2027} \\ <\mathrm{B}_{2026} \end{gathered}$ | $\begin{aligned} & \mathrm{B}_{2027} \\ & <B_{40 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2027} \\ & <B_{25 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2027} \\ & <B_{10 \%} \end{aligned}$ | $\begin{gathered} 2026 \\ \begin{array}{c} \text { Fishing } \\ \text { intensity } \\ >100 \% \end{array} \end{gathered}$ | $\begin{gathered} 2027 \\ \begin{array}{c} \text { Default HR } \\ \text { catch } \\ >2026 \\ \text { catch } \end{array} \\ \hline \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| a: | 0 | 0.66 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| b: | 180,000 | 0.73 | 0.03 | 0.00 | 0.00 | 0.00 | 0.00 |
| c: | 225,000 | 0.74 | 0.04 | 0.00 | 0.00 | 0.00 | 0.00 |
| d: | 259,200 | 0.75 | 0.05 | 0.01 | 0.00 | 0.01 | 0.01 |
| e: | 264,000 | 0.75 | 0.05 | 0.01 | 0.00 | 0.01 | 0.01 |
| f: | 350,000 | 0.78 | 0.07 | 0.02 | 0.00 | 0.03 | 0.05 |
| g: | 283,500 | 0.75 | 0.06 | 0.01 | 0.00 | 0.01 | 0.02 |
| h : | 380,000 | 0.78 | 0.09 | 0.02 | 0.00 | 0.04 | 0.07 |
| i: | 307,800 | 0.76 | 0.07 | 0.01 | 0.00 | 0.02 | 0.03 |
| j: | 430,000 | 0.79 | 0.11 | 0.03 | 0.00 | 0.07 | 0.12 |
| k: | 545,000 | 0.82 | 0.17 | 0.06 | 0.01 | 0.18 | 0.29 |
| $1:$ | 625,000 | 0.83 | 0.21 | 0.09 | 0.01 | 0.27 | 0.41 |
| m: | 782,426 | 0.84 | 0.35 | 0.19 | 0.03 | 0.50 | 0.65 |
| n : | 717,464 | 0.84 | 0.29 | 0.14 | 0.02 | 0.41 | 0.56 |
| o: | 712,782 | 0.84 | 0.29 | 0.15 | 0.02 | 0.41 | 0.56 |

## Research and data needs

There are many research projects that could improve the stock assessment for Pacific Hake and lead to improved biological understanding and decision-making. The most important are as follows:

1. Continue to conduct research to evaluate ways to improve recent, current, and future estimates of recruitment for use in stock assessment. This could include the development of time series of recruitment indices, time series of informative environmental or ecosystem variables, and models that have predictive skill (e.g., Vestfals et al. 2023). Explorations should also consider options for incorporating information on recruitment into the assessment model and the management framework for Pacific Hake. For example, time series could be included in the stock assessment as a standalone data source (similar to the acoustic indices) or improvements could be made to the modeling framework such that these environmental time series could impact the stock-recruitment relationship directly. Results from such work should be connected to or in cooperation with ongoing research related to recruitment variability as discussed in Section 3.3. Related, there is a need to streamline and broaden the availability of products from oceanographic models (e.g., Regional Ocean Modeling System) so they are available across international boundaries and updated on a recurring basis, thereby allowing for their use as informative links in operational stock assessments. A successful example of this has been the annual production of Pacific Hake distribution forecasts that depend on 6-9 month forecasts of subsurface (i.e., 100 m depth) temperature from J-SCOPE. Furthermore, the existing management strategy evaluation framework should be used, or further developed, to examine how information on recruitment can inform robust management decisions.
2. Conduct research on estimates of uncertainty for the relative age- 1 index and the age- $2+$ index and investigate alternative ways to utilize survey age-composition information in the assessment model. Bootstrapping of the acoustic survey time series, or related methods, could help incorporate uncertainty related to the target-strength relationship, subjective scoring of echograms, thresholding methods, and methods used to estimate the species mixes for interpreting the acoustic backscatter into the variance calculations. Research should be communicated with those involved in developing the U.S. West Coast Integrated Survey Initiative. The management strategy evaluation framework should be used, or further developed, to examine how changes in survey methods can be used to inform robust management decisions.
3. Work with regional partners to develop an annual workflow that provides key metrics, indicators, or other summaries of general ecosystem conditions relevant to the coast-wide population of Pacific Hake. In particular, include indicators that are potentially associated with Pacific Hake biology and ecology (e.g., recruitment, distribution, predation, prey, and communities). Such information can broaden the context within which a single species stock assessment is interpreted, be used to support model development, refine uncertain assessment conclusions (e.g., productivity), and provide other non-assessment indicators of the system's state to management.

## 1 INTRODUCTION

The Joint U.S.-Canada Agreement for Pacific Hake (called the Agreement) was signed in 2003, went into force in 2008, and was implemented in 2010. The committees defined by the Agreement were first formed in 2011, and 2012 was the first year for which the process defined by the Agreement was followed. This is the thirteenth annual stock assessment conducted under the Agreement process.

Under the Agreement, Pacific Hake (Merluccius productus, also referred to as Pacific whiting) stock assessments are to be prepared by the Joint Technical Committee (JTC) comprised of both U.S. and Canadian scientists and reviewed by the Scientific Review Group (SRG) that consists of representatives from both nations. Additionally, the Agreement calls for both of these bodies to include scientists nominated by an Advisory Panel (AP) of fishery stakeholders.

The primary data sources for this assessment include an acoustic survey, annual fishery catch, mean weight-at-age data, maturity-at-age data, as well as survey and fishery agecomposition data. The assessment depends primarily upon an index of biomass from the acoustic survey for information on the scale of the current population. Age-composition data provide additional information allowing the model to resolve strong and weak cohorts. The catch is an important source of information regarding changes in abundance and places a lower bound on the available population biomass in each year.
This assessment is fully Bayesian, with the base model incorporating prior information on several key parameters (including informative priors on natural mortality, $M$, and steepness of the stock-recruitment relationship, $h$ ) and integrating over parameter uncertainty to provide results that can be probabilistically interpreted. From a range of alternate models investigated by the JTC, a subset of sensitivity analyses are also reported to provide a broad qualitative comparison of structural uncertainty with respect to the base model (Section 3.8). The structural assumptions of this 2024 base model, implemented using version 3.30.22 of the Stock Synthesis software (Methot, Jr. and Wetzel 2013), differ from the 2023 base model in that the distribution assumed for the relative age- 1 index is now a student-t distribution rather than a log-normal distribution (Berger et al. 2023), and maturity-at-age is now year-specific (Section 2.4.2 and Appendix G). All model runs reported in this document are performed in a Bayesian context. Responses to 2023 SRG requests are in Section 3.3 and a Glossary of terms appears in Appendix C.

### 1.1 Stock structure and life history

Pacific Hake is a semi-pelagic, schooling species distributed along the west coast of North America, generally ranging in latitude from $25^{\circ} \mathrm{N}$ to $55^{\circ} \mathrm{N}$ (Figure 1). It is among 18 species of hake from four genera (being the majority of the family Merluccidae), which are found in both hemispheres of the Atlantic and Pacific Oceans (Alheit and Pitcher 1995; Lloris et al. 2005). The coastal population of Pacific Hake is currently the most abundant groundfish population in the California Current system. Smaller populations of this species occur in the major inlets of the Northeast Pacific Ocean, including the Strait of Georgia, the Puget Sound, and the Gulf of California. Each of these smaller populations
are genetically distinct from the coastal population (Vrooman and Paloma 1977; Iwamoto et al. 2004; King et al. 2012; García-De León et al. 2018; Longo et al. In press). The coastal population is also distinguished from the inshore populations by larger size at age and seasonal migratory behavior and from fish off the west coast of Baja California by smaller size at age and later spawning (Zamora-García et al. 2020).

The coastal population of Pacific Hake typically ranges from the waters off southern California to northern British Columbia and rarely into southern Alaska, with the northern boundary related to fluctuations in annual migration (Hamel et al. 2015) depending, in part, on water temperature (Malick et al. 2020a, 2020b). In spring, adult Pacific Hake migrate onshore and northward to feed along the continental shelf and slope from Northern California to Vancouver Island. In summer, Pacific Hake often form extensive mid-water aggregations in association with the continental shelf break, with the highest densities located over bottom depths of 200-300 m (Dorn and Methot 1991, 1992).

Older Pacific Hake exhibit the greatest northern migration each season, with two- and three-year old fish rarely being observed in Canadian waters north of Southern Vancouver Island. During El Niño events (warm ocean conditions such as in 1998 and 2016), a larger proportion of the population migrates into Canadian waters (Figure 2), due to temperature effects (Malick et al. 2020a) and possibly intensified northward transport during the period of active migration (Dorn 1995; Agostini et al. 2006). In contrast, La Niña conditions (colder water, such as in 2001, 2011, and 2021) result in a southward shift in the distribution of Pacific Hake, with a much smaller proportion of the population found in Canadian waters compared to during El Niño years, a trend evident from the acoustic surveys (Figure 2). In general, warmer than average thermal habitat conditions for mature Pacific Hake lead to relatively higher biomass further north and relatively lower biomass around the U.S.-Canadian border, while cooler than average conditions lead to relatively higher biomass of immature Pacific Hake generally spread evenly across their distribution (Malick et al. 2020a). The distribution of age-1 fish also changes between years (Figure 3).

### 1.2 Ecosystem considerations

Pacific Hake are important to ecosystem dynamics in the Eastern Pacific Ocean due to their relatively large total biomass and potentially large role as both prey and predator (Hicks et al. 2013). Ongoing research investigating abiotic (i.e., environmental conditions) and biotic (e.g., maturity and diet) drivers of the distribution, recruitment, growth, and survival of Pacific Hake could provide insight into how the population is linked with broader ecosystem considerations. For example, Turley and Rykaczewski (2019) found decreased survival of larval Pacific Hake as storm events increased, contrary to many other species in the Southern California Current Ecosystem. An analysis of drivers of recruitment across the maternal preconditioning, egg, and larval phases of Pacific Hake recruitment indicated recruitment is associated with eddy kinetic energy, the location of the North Pacific Current bifurcation, and upwelling during maternal preconditioning, as well as with northward long-shore transport and the number of days between storm events during larval stages (Vestfals et al. 2023). Phillips et al. (2022) suggest temperature dynamically
influences the co-occurrence of Pacific Hake and krill (i.e., euphausiids; Euphausiacea), which can influence annual Pacific Hake growth and recruitment as the availability of key prey species shifts. Temperature was also found to impact the co-occurrence of Pacific Hake and shortbelly rockfish (Sebastes jordani, Sebastidae) during the 2014-2016 marine heatwave (Free et al. 2023). An index of abundance for Humboldt Squid (Dosidicus gigas) suggests that the abundance of Pacific Hake decreases with increasing squid abundance (Stewart et al. 2014; Taylor et al. 2015). Many additional research topics relevant to Pacific Hake distribution, recruitment, and growth patterns in relation to oceanographic conditions have been investigated (Ressler et al. 2007; Hamel et al. 2015; Malick et al. 2020a, 2020b) and provide a foundation for further research on these topics.

Recent oceanographic trends and large-scale ecosystem conditions, as summarized in NOAA's annual California Current Ecosystem Status Reports, provide insight into potential drivers of Pacific Hake population dynamics and fleet operations. Periods of high productivity are often marked by strong winter upwelling which brings nutrients to coastal waters, cooler temperatures, an energy-rich copepod community, and high productivity of krill, a key food source for Pacific Hake (Buckley and Livingston 1997; Harvey et al. 2021). During 2023 (Leising et al. 2024), basin-scale climate patterns suggest average conditions for ecosystem productivity as El Niño conditions continue to strengthen. Despite being highly variable in strength, upwelling was generally less than in previous years. Weaker upwelling conditions have been associated with higher Pacific Hake recruitment (Vestfals et al. 2023). In the southern California Current Ecosystem, the abundance of Pacific Hake larvae in the CalCOFI survey area has been above average for two consecutive years, with 2023 being the highest since 2011. At the same time, hake predation by bluefin and swordfish were also well above average in recent years (2021 and 2022). Abundance of krill, a common prey of Pacific Hake, was below average in 2023. Lipid-rich copepod biomass was average and relatively stable throughout the summer ('northern' copepod index) of 2023 relative to previous years, but was below average in the winter ('southern' copepod index).

Fitting the assessment model to empirical weight-at-age data allows for time-varying growth without needing a mechanistic relationship or environmental data, which facilitates an 'Ecosystem Approach to Fisheries Management' (a priority for DFO and NOAA; see Section 2.4.1). Furthermore, the new year-specific maturity-at-age specifically includes a temperature effect from a spatiotemporal model (Section 2.4.2 and Appendix G). Related ongoing research should provide more insights into the specific mechanisms affecting changes in growth and fecundity, which will enable further condition-specific prediction capabilities (e.g., assumptions of growth, or weight at age, and fecundity during forecast years). It is hypothesized that temporal changes in weight-at-age data may be due to ecosystem effects such as prey availability, predator abundance, and ocean temperature (Chittaro et al. 2022).

### 1.3 Management of Pacific Hake

Since the implementation of the Magnuson-Stevens Fishery Conservation and Management Act in the U.S. and the declaration of a 200-mile fishery-conservation zone in the
U.S. and Canada in the late 1970s, annual quotas (or catch targets) have been used to limit the catch of Pacific Hake in both countries' zones. Scientists from both countries historically collaborated through the Technical Subcommittee of the Canada-U.S. Groundfish Committee (TSC), and there were informal agreements on the adoption of annual fishing policies. During the 1990s, however, disagreements between the U.S. and Canada on the allotment of the catch limits between U.S. and Canadian fisheries led to quota overruns; the 1991-1992 national quotas summed to $128 \%$ of the coast-wide limit, while the 1993-1999 combined quotas were an average of $112 \%$ of the limit. The Agreement establishes U.S. and Canadian shares of the coast-wide total allowable catch (TAC) at $73.88 \%$ and $26.12 \%$, respectively, and this distribution has largely been adhered to since 2005. A bilateral agreement on the coast-wide TAC could not be reached in 2020 and 2021; so, catch targets were set unilaterally during these years for the first time since the inception of the Agreement. Catch allocations as specified in the Agreement have since been applied.

Since 1999, an upper limit on catch has been calculated using an $F_{\mathrm{SPR}=40 \%}$ default harvest rate with a 40:10 adjustment. This decreases the catch linearly from the catch at a relative spawning biomass of $40 \%$ to zero catch at a relative spawning biomass values of $10 \%$ or less (called the default harvest policy in the Agreement); relative spawning biomass is the female spawning biomass divided by that at unfished equilibrium. Further considerations have almost always resulted in catch targets being set lower than the recommended catch limit. Total catch has not exceeded the coast-wide quota since 2002, and harvest rates are likely to have never exceeded the $F_{\mathrm{SPR}=40 \%}$ target.

### 1.3.1 Management of Pacific Hake in the United States

In the U.S. zone, participants in the directed fishery are required to use pelagic trawls with a codend mesh of at least 7.5 cm . Regulations have also restricted the area and season of fishing to reduce the bycatch of Chinook Salmon (Oncorhynchus tshawytscha), depleted rockfish populations, and other species as related to their specific harvest specifications. The current allocation agreement, effective since 1997, divides the U.S. harvest into tribal ( $17.5 \%$ ) and non-tribal ( $82.5 \%$, including a small amount set aside for research) components. Starting in 1996, the Makah Tribe has conducted a fishery with the tribal allocation in its 'usual and accustomed fishing area'. The non-tribal harvest allocation is divided among catcher-processors (34\%), motherships (24\%), and shore-based vessels (42\%). Since 2011, the non-tribal U.S. fishery has been fully rationalized with allocations in the form of Individual Fishing Quotas (IFQs) to the shore-based sector and group shares to cooperatives in the at-sea mothership (MS) and catcher-processor (CP) sectors. The At-Sea Hake Observer Program has been monitoring fishing vessel activity since 1975, originally monitoring foreign and joint-venture vessels. Observer coverage has been 100\% on all domestic vessels since 1991, including the 2020 and 2021 fishing seasons, despite the COVID-19 pandemic.

Shortly after the 1997 allocation agreement was approved by the Pacific Marine Fisheries Commission, fishing companies owning catcher-processor vessels with U.S. West Coast groundfish permits established the Pacific Whiting Conservation Cooperative (PWCC).

The primary role of the PWCC is to distribute the catcher-processor allocation among its members to achieve greater efficiency and product quality, as well as promoting reductions in waste and bycatch rates relative to the former 'derby' fishery in which all vessels competed for a fleet-wide quota. The mothership fleet has also formed a cooperative where bycatch allocations are pooled and shared among the vessels. The individual cooperatives have internal systems of in-season monitoring and spatial closures to avoid and reduce bycatch of salmon and rockfish.

### 1.3.2 Management of Pacific Hake in Canada

Canadian groundfish managers distribute their portion of the coast-wide TAC as quota to individual license holders. In 2023, Canadian hake fishermen were allocated a TAC of $163,250 \mathrm{t}$, which did not include any carryover quota. Canadian priority lies with the domestic fishery. However, when there is determined to be an excess of fish for which there is not enough domestic processing capacity, fisheries managers give consideration to a Joint-Venture fishery in which foreign processor vessels are allowed to accept codends from Canadian catcher vessels while at sea. The last year there was Joint-Venture quota allocation was in 2018.

In 2023, all Canadian Pacific Hake trips were subject to $100 \%$ observer coverage by electronic monitoring for both the shoreside component of the domestic fishery and the freezer-trawler component. There is no in-person observer program for the Canadian Pacific Hake fisheries.

Retention of all catch, with the exception of prohibited species, was mandatory. The retention of groundfish other than Sablefish, Mackerel, Walleye Pollock, and Pacific Halibut on dedicated Pacific Hake trips using electronic monitoring was not allowed to exceed $10 \%$ of the landed catch weight. The bycatch allowance for Walleye Pollock was $30 \%$ of the total landed weight.

### 1.4 Fisheries

The fishery for the coastal population of Pacific Hake occurs along the coasts of Northern California, Oregon, Washington, and British Columbia primarily during May-November. The fishery is conducted with mid-water trawls and has met the Marine Stewardship Council (MSC) Fisheries Standard to be certified as meeting sustainable fishing benchmarks since 2009. Foreign fleets dominated the fishery until 1991, when domestic fleets began taking the majority of the catch. Catches were occasionally greater than 200,000 t prior to 1986, and since then, they have been greater than 200,000 $t$ for all except four years.
In 2021, the Pacific Hake fishery was Canada's largest commercial wild fishery (species with the largest catch), representing 10\% of Canada's total landings of all species (www.dfompo.gc.ca). Over CA\$19 million in wages was estimated to have been paid to employees of the processing industry in British Columbia in 2021, with an exported value of $>\mathrm{CA} \$ 60 \mathrm{mil}-$ lion in product to Ukraine ( $>$ CA\$25 million), China, South Africa, Lithuania, and other countries (Fisheries and Oceans Canada 2023).

In the U.S.A., over U.S. $\$ 75.1$ million in wages is estimated to have been paid to employees in 2021 (www.noaa.gov). This includes wages paid to crew and captains fishing on catcher vessels that deliver shoreside and at-sea to motherships; workers in shore-based processing facilities; crew, captains, and workers on catcher-processor vessels; and workers on mothership vessels. The exported value of Pacific Hake was U.S. $\$ 163$ million in 2021, including to Ukraine, Nigeria, and Netherlands, which make up about $73 \%$ of the total (www.noaa.gov). The total economic impact of the Pacific Hake fishery on the U.S. West Coast in 2021 was U.S. $\$ 335$ million in income and 4,450 jobs.

The Joint Management Committee (JMC) determined an adjusted (for carryovers) coastwide TAC of $625,000 \mathrm{t}$ for 2023 . The U.S. catch target was set at $461,750 \mathrm{t}$ and the Canadian catch target at 163,250 t . A brief review of the 2023 fishery is presented here by country (Tables 1-3 and Figure 4). Additional information is available in annual U.S. and Canada Advisory Panel reports (Appendices E and D).

### 1.4.1 Fisheries for Pacific Hake in the United States

The U.S. specified catch target (i.e., adjusted for carryovers) of 461,750 t was further divided among the research, tribal, catcher-processor, mothership, and shore-based sectors. After the tribal allocation of $17.5 \%(80,806 t)$ and a $750 t$ allocation for research catch and bycatch in non-groundfish fisheries, the 2023 non-tribal U.S. catch limit of $380,194 \mathrm{t}$ was allocated to the catcher-processor (34\%), mothership ( $24 \%$ ), and shore-based ( $42 \%$ ) commercial sectors. Reallocation of $45,000 \mathrm{t}$ of tribal quota to non-tribal sectors on September 27 resulted in final quotas for the catcher-processor, mothership, and shore-based sectors of $144,566 \mathrm{t}, 102,047 \mathrm{t}$, and $178,581 \mathrm{t}$, respectively.

The U.S. fishery for Pacific Hake began on May 1. Prior to 2015, the shore-based sector was allowed to fish starting June 15 north of $42^{\circ} \mathrm{N}$ latitude (the Oregon-California border) and April 1 between $40^{\circ} 30^{\prime} \mathrm{N}$ and $42^{\circ} \mathrm{N}$ latitudes, whereas the at-sea sectors were allowed to fish starting May 15. Between 2015-2022, the shore-based sector was allowed to fish north of $40^{\circ} 30^{\prime} \mathrm{N}$ latitude starting May 15 and south of $40^{\circ} 30^{\prime} \mathrm{N}$ latitude starting April 15, although only $5 \%$ of the shore-based allocation was released for this early period. Since 2023, all sectors have been allowed to fish starting May 1. Regulations do not allow at-sea processing or night fishing (midnight to one hour after official sunrise) south of $42^{\circ} \mathrm{N}$ latitude at any time during the year.
The total catch of Pacific Hake in U.S. waters was the 13th highest value ever recorded (Table 1) and the U.S. utilization rate ( $52.1 \%$ ) was the lowest it has been since 2015 (see Appendix E for more details). The catcher-processor, mothership, and shore-based sectors caught $74.1 \%, 32.3 \%$, and $56.2 \%$ of their final reallocated quotas, respectively. Tribal landings, which are included in the shoreside sector totals were 0 t . The median fishing depth for the at-sea sectors was the same as last year (Figure 5). The shore-based sector had the largest monthly catches during July, August, and September. While, the at-sea sectors had the largest monthly catches during May, September, and October.
In both U.S. at-sea sectors, age-2, age-3, and age-7 fish, associated with the 2021, 2020, and 2016 year classes, were the most common ages. Both the age- 3 and age- 7 fish were
seen last year in appreciable numbers as age-2 and age-6 fish. The reported proportions at age summarize sampling efforts on 391 catcher-processor hauls and 127 mothership hauls (Table 4). For the catcher-processor sector, the four most abundant age classes (by numbers) seen in 2023 were age-2 (51.3\%), age-3 (24.0\%), age-7 (8.9\%), and age-9 (5.0\%); Table 5. For the mothership sector, the four most abundant age classes for 2023 were age- 2 (39.6\%), age-3 (33.2\%), age-7 (7.5\%), and age-6 (5.5\%) (Table 6).

Age-samples from 66 shoreside trips showed similar age compositions in the catch compared to the at-sea sectors, though not nearly as many age- 2 fish. The four most abundant age classes for highest occurrences being for 2023 were age-3 (27.0\%), age-7 (19.3\%), age-2 ( $16.4 \%$ ), and age- 9 ( $11.6 \%$ ) (Table 7). Age-compositions from the at-sea and shoreside sectors during this last year were more similar than they were the previous year.

### 1.4.2 Fisheries for Pacific Hake in Canada

The 2023 Canadian Pacific Hake domestic fishery removed 23,557 t from Canadian waters (Table 2), which was $14.4 \%$ of the Canadian TAC of $163,250 \mathrm{t}$. For the second year in a row, the attainment for Canada was much lower than usual, due to the fishing vessels having a difficult time finding fish in Canadian waters (see Appendix D and last year's assessment, Berger et al. (2023), for more details).

The shoreside vessels, which land fresh round product onshore, landed 3,657 t in 2023, the lowest on record since 1990, and a little less than the 2022 landings of 3,868 t. The Freezer trawlers, which freeze headed and gutted product while at sea, landed 19,901 t. This was the lowest amount the Freezer trawlers have landed since 2013, despite doubling the number of vessels since then.

Fishing started in early April and ended in November. The general view of the Canadian fleet is that abundance in Canadian waters remained below normal levels in 2023, including the normally-abundant areas outside of Southwest Vancouver Island. Reports of difficulties finding fish in 2023 led to some vessels not leaving the dock, therefore amplifying the effect of low catches. The fish caught in Canada appeared to be mostly from four age classes (see below), with very few smaller fish (less than 500 grams) caught.

The most abundant year classes in the Canadian shoreside catch (by numbers; Table 8) were age-7 (31.5\%), age-6 (17.3\%), age-9 (13.2\%), and age-13 (9.4\%). The most abundant year classes in the Canadian freezer-trawler catch (by numbers; Table 9) were age-7 ( $21.6 \%$ ), age-9 (19.6\%), age-6 (16.0\%), and age-13 (13.9\%).

## 2 DATA

Fishery-dependent and fishery-independent data used in this assessment (Figure 7) include the following sources:

- Total catch from all U.S. and Canadian fisheries that targeted Pacific Hake from 1966 to 2023 (Tables 1-3).
- Fishery age compositions aggregated by year and country-specific sector for the last ten years are available (Tables 5-9) to investigate region-specific trends; age compositions aggregated by year, composed of data from the U.S. fishery (1975-2023) and the Canadian fishery (1988-2023), are used to fit the model (Table 10 and Figure 8).
- An age-2+ biomass index and age compositions from the Joint U.S. and Canadian Integrated Acoustic and Trawl Survey (1995, 1998, 2001, 2003, 2005, 2007, 2009, 2011, 2012, 2013, 2015, 2017, 2019, 2021, 2023; Tables 11, 12, and 13; Figures 8 and 9).
- The relative age- 1 index (billions of age- 1 fish) derived from the Joint U.S. and Canadian Integrated Acoustic and Trawl Survey (1995, 1998, 2003, 2005, 2007, 2009, 2011, 2012, 2013, 2015, 2017, 2019, 2021, 2023; Table 12; Figure 9).

The following biological relationships, derived from external analysis of auxiliary data, were input as fixed values in the assessment model:

- Ageing-error matrices based on cross-read and double-blind-read otoliths.
- Annual proportions of female Pacific Hake at each age that are mature, as developed from a new analysis (Section 2.4.2 and Appendix G) of histological analyses of ovary samples (Table 14; Figure 11).
- Mean observed weight-at-age data from fishery and survey catches (1975-2023; Figures 12-15) and, thus, derived fecundity-at-age as well (Figures 11 and 14).

Additional data sources not used in this assessment are discussed in Section 2.3.

### 2.1 Fishery-dependent data

### 2.1.1 Total catch

The catch of Pacific Hake for 1966-2023 is summarized by country-specific sectors (Tables 1-3) and modeled as annual coast-wide catches. Catches in U.S. waters prior to 1978 are available only by year from Bailey et al. (1982) and historical assessment documents. Canadian catches prior to 1989 are also unavailable in disaggregated form. U.S. shorebased landings are from the Pacific Fishery Information Network (PacFIN) database. Foreign and Joint-Venture catches for 1981-1990 and U.S. domestic at-sea catches for 1991-2023 are calculated from the Alaska Fisheries Science Center's North Pacific Groundfish and Halibut Observer (NORPAC) database, which also stores data from the At-Sea Hake Observer Program. Canadian Joint-Venture catches from 1989 are from the Groundfish Biological (GFBio) database. Canadian shoreside landings are from the Groundfish Catch
(GFCatch) database for 1989-1995, the Pacific Harvest Trawl (PacHarvTrawl) database for 1996-March 31, 2007, and the Fisheries Operations System (FOS) database for April 2007-present.

Minor updates to catches used in previous assessments were made based on the best available information extracted from the aforementioned databases. Tribal catches were available in PacFIN for the U.S. tribal fishery at the time the data were extracted and were cross-checked with numbers based on information provided by the Makah Tribe. The Makah Tribe is also working on providing historical catches such that shore-based catches can be summarized separately from tribal catches since the onset of the fishery.

Historically, the fishery for Pacific Hake has been well covered by observers with slight differences in coverage by sector. Currently, U.S. shore-based vessels carry observers and are required to retain all catch and bycatch for sampling by plant observers. Vessels participating in the U.S. at-sea sectors are also required to have observers on board and have since 1990. U.S. foreign and Joint-Venture vessels had varying levels of coverage from 1975-1990 ranging from $21-100 \%$ coverage. Canadian Joint-Venture and Canadian freezertrawler vessels were monitored by at-sea observers from 1996-2019. In 2020 and 2021 there were no observers on Canadian freezer trawlers due to staffing issues and in 2022 the decision was made to stop providing observers on board all Canadian vessels. Early in 2022 a sampling plan designed by Canadian managers, scientists, and the sampling contractor, Archipelago Marine Research Ltd. (AMR), was put into place to ensure the ongoing sampling of Pacific Hake on Canadian vessels (see Section 2.1.2). Canadian trawl catches are monitored autonomously at sea by cameras onboard vessels. Catch is recorded by dockside samplers within the Groundfish Trawl Dockside Monitoring Program using total catch weights provided by processing plants. Discards are negligible relative to the total fishery catch for all sectors.

### 2.1.2 Fishery biological data

Biological information from the U.S. at-sea sectors was extracted from the NORPAC database. This included sex, length, weight, and age information from the foreign and Joint-Venture fisheries from 1975-1990 and from the domestic at-sea fishery since 1990. Observers collected data by selecting fish randomly from each haul. The number of otoliths collected per haul has varied over time but is currently three fish every third haul.

Since 1991, biological samples from the U.S. shore-based sector have been sampled by port samplers located where there are substantial landings of Pacific Hake, primarily Eureka, Newport, Astoria, and Westport. Port samplers took one sample per offload (or trip) consisting of 100 randomly selected fish for individual length and weight. From those 100 fish, typically 20 fish were randomly subsampled for otolith extraction.

When there were observers (1996-2019) aboard Canadian freezer trawlers, they collected 50 otoliths and 300 lengths per sample, sampling once per day during trips that on average last approximately seven days. Since 2022, freezer-trawler employees have collected fish for sampling from two tows per trip and delivered them to the dock as frozen specimens where they are eventually sampled for length, weight, sex, and otoliths. Each delivery
consists of approximately 100 fish in two separate bags. The bags each hold approximately 50 fish which were removed from a single tow. Due to unforeseen circumstances while at sea, some trips did not deliver any bags and some only delivered single bags upon arrival from a trip.

For electronically observed Canadian shoreside trips, port samplers obtained biological data from the landed catch. For each sampled trip, approximately 50 ages and 300 lengths were sampled from the catch. Observed domestic haul-level information is aggregated to the trip level to be consistent with the unobserved trips that were sampled in ports.

In years when there was a Canadian Joint-Venture fishery, length samples were collected every second day of fishing operations and otoliths were collected once per week. Length and age samples were taken randomly from a given codend. The sample weight from which biological information was collected was inferred from length-weight relationships.
The sampling unit for shore-based samples is the trip, while the haul is the primary unit for the at-sea samples (Table 4). There is no least common denominator for aggregating at-sea and shore-based samples because detailed haul-level information is not recorded for shore-based trips and hauls sampled in the at-sea sectors cannot be aggregated to a comparable trip level. As a result, initial sample sizes are simply the summed hauls and trips for fishery biological data.
Biological data were analyzed based on the sampling protocols used to collect them and expanded to estimate the corresponding statistic from the entire landed catch by fishery and year when sampling occurred. A description of the analytical steps for expanding the age compositions can be found in earlier stock assessment documents (Hicks et al. 2013; Taylor et al. 2014).
The aggregate fishery age-composition data (1975-2023) confirm the well-known pattern of large cohorts born in 1973, 1977, 1980, 1984, 1987, 1999, 2008, 2010, 2014, and 2016, and suggests large cohorts for 2020 and 2021 (Table 10 and Figure 8). Recent age-composition data still easily track the 2010 cohort, as well as the large cohorts born since then (Table 10 and Figure 8). Currently, the 2021 cohort is the largest observed cohort in the U.S. at-sea sectors (Tables 5-6), the 2020 cohort is the largest observed cohort in the U.S. shore-based sector (Table 7), and the 2016 cohort is the largest observed cohort in both Canadian fleets (Tables 8-9). Age-1 fish were observed by the fishery this year (Table 10) in the U.S. For the combined data in 2023, the 2021 cohort was the largest (35\%), followed by the 2020 cohort ( $25 \%$ ), and then the 2016 cohort ( $13 \%$ ). For the combined data in 2022, the 2020 cohort was the largest ( $32 \%$ ), followed by the 2016 cohort ( $24 \%$ ), and then the 2014 cohort (15\%).

We caution that proportion-at-age data contain information about the relative numbers at age, and these can be affected by changing recruitment, selectivity, or fishing mortality, making these data difficult to interpret on their own. For example, the above-average 2005 and 2006 year classes declined in proportion in the 2011 fishery samples but persisted in small proportions for years in the fishery catch but were much reduced starting in 2011 due to mortality and the overwhelming size of the more recent large cohorts. The assessment model is fit to these data to estimate the absolute sizes of incoming cohorts,
which become more precise after they have been observed several times (i.e., encountered by the fishery and survey over several years).
Both the weight- (Figure 15; Section 2.4.1) and length-at-age information suggest that growth of Pacific Hake has fluctuated markedly over time (see Figure 7 in Stewart et al. 2011). This is particularly evident in the frequency of larger fish ( $>55 \mathrm{~cm}$ ) before 1990 and a recent linear shift towards larger fish. Although length-composition data are not fit explicitly in the base model presented here, the presence of the 2008 and 2010 year classes have been clearly observed in length data from the U.S. fishery sectors and the 2014 year class has been apparent since 2016.

### 2.2 Fishery-independent data

### 2.2.1 Acoustic survey

The Joint U.S. and Canadian Integrated Acoustic and Trawl Survey (Stewart et al. 2011) has been the primary fishery-independent tool used to assess the distribution, abundance, and biology of coastal age- $2+$ Pacific Hake along the west coasts of the U.S.A. and Canada. The acoustic surveys performed in 1995, 1998, 2001, 2003, 2005, 2007, 2009, 2011, 2012, 2013, 2015, 2017, 2019, 2021, and 2023 were used in this assessment (Table 12). The acoustic survey samples transects that represent all waters off the coasts of the U.S.A. and Canada thought to contain all portions of the age- $2+$ Pacific Hake stock. Observations of age- 0 and age- 1 Pacific Hake are excluded from the age- $2+$ index due to largely different schooling behavior relative to older Pacific Hake, concerns about their catchability by the trawl gear, and differences in expected location during the summer months when the survey takes place. Observations of age- 1 Pacific Hake are recorded during the survey, and additional analyses, described below, are conducted to develop a relative age- 1 index.

The 2023 survey covered U.S. and Canadian waters from Point Conception to north of Haida Gwaii using 116 transects (Figure 2). In the U.S.A., transects were mostly separated by 10 nmi ; six transects were dropped to account for available ship days at sea. In Canada, transects were separated by 10 nmi along Vancouver Island and then 20 nmi further north. The NOAA Ship Bell M. Shimada and the Canadian Coast Guard Ship Sir John Franklin worked collaboratively to complete the full extent of the survey in 2023.

Distributions of the backscatter of Pacific Hake plotted for each acoustic survey since 1995 illustrate the variable spatial patterns across time of age- $2+$ fish (Figure 2). This variability is due in part to changes in the composition of the age- $2+$ population because older Pacific Hake tend to migrate farther north and partially due to environmental and/or climatic factors. The 1998 acoustic survey is notable because it shows an extreme northward distribution that is thought to be related to the strong 1997-1998 El Niño. In contrast, the distribution of Pacific Hake during the 2001 acoustic survey was compressed into the lower latitudes off the coast of Oregon and Northern California following the strong La Niña event in 2000. In 2003, 2005, and 2007 the distribution of Pacific Hake did not show an unusual coast-wide pattern despite 2003 and 2007 being characterized as El Niño years. In 2009, 2011, 2012, and 2013 the majority of the distribution of Pacific Hake was again found in U.S. waters, which is more likely due to age composition than the environment
because 2013 had some warmer than average sea-surface temperatures. In 2015, sea-surface temperatures were warmer again, resulting in a northern shift in the overall distribution. The distribution of Pacific Hake in 2017 was more latitudinally uniform than observed in 2015, which is likely a result of having large proportions of both the 2010 and 2014 cohorts (Figure 2). Weak 2019 El Niño conditions decreased in their prevalence starting in March of that year, leading to neutral conditions by July. Consequently, during the 2019 survey Pacific Hake were found on all survey transects from just north of Morro Bay, California to the northern end of Vancouver Island, with the greatest offshore extent found off of Cape Mendocino. During the 2021 survey, the majority of Pacific Hake were found in U.S. waters, congruent with the continuation of La Niña conditions in the California Current from 2020 to 2021. Despite the switch to El Niño conditions in April of 2023, very few fish were seen in Canadian waters during the 2023 survey. Ongoing research is looking into relationships between environmental conditions and Pacific Hake distribution and recruitment that will help to inform the mechanisms behind observations (Malick et al. 2020b; Phillips et al. 2023).

During the acoustic surveys, mid-water trawls are made opportunistically to determine the species composition of the observed acoustic sign and to obtain the length data necessary to scale the acoustic backscatter into biomass (see Table 12 for the number of trawls in each survey year). Biological samples collected from these trawls are post-stratified, based on similarity in size composition, and the composite length frequency is used to characterize the size distribution of Pacific Hake along each transect and to predict the expected backscattering cross section for Pacific Hake based on the fish-size target-strength (TS) relationship. Any potential biases that might be caused by factors such as alternative TS relationships are partially accounted for in catchability. But variability in the estimated survey biomass due to uncertainty in TS is not explicitly accounted for in the assessment.
Data from the acoustic survey are analyzed using kriging, which accounts for spatial correlation, to provide an estimate of total biomass as well as an estimate of the yearspecific sampling variability due to patchiness of schools of Pacific Hake and irregular transects (Petitgas 1993; Rivoirard et al. 2000; Mello and Rose 2005; Simmonds and MacLennan 2006). Advantages to the kriging approach are discussed in the 2013 stock assessment (Hicks et al. 2013).
For the 2016 assessment (Grandin et al. 2016), the data from all surveys since 1998 were scrutinized and reanalyzed using consistent assumptions, an updated version of the EchoPro software, and a common input-file structure because some previously generated files had spurious off-transect zeros because of how the data were exported. The same analytical procedure was carried out during the reanalysis of 1995 survey data (Berger et al. 2017) and during the preparation of survey data collected since 2017. The assumptions are as follows:

- fixed minimum $\left(k_{\min }=3\right)$ and maximum $\left(k_{\min }=10\right)$ number of points used to calculate the value in a cell;
- search radius is three times the length scale that is estimated from the variogram; and
- biomass decays with distance from the end of the transect when extrapolating biomass beyond the western end of a transect, which was refined and supported by the SRG starting with the 2016 assessment (Grandin et al. 2016).

Starting in 2021, the EK 60 echosounders were replaced with EK 80 echosounders, and thus, survey estimates from years using the new echosounders are scaled by factor of 1.06 to convert the EK 80 to EK 60 acoustic data. The survey team will eventually be converting all pre-2021 EK 60 data to an equivalent EK 80 format. Thus, a full time series of consistently analyzed survey biomass (Table 12 and Figure 9) and age compositions (Table 11 and Figure 8) since 1995 are used to fit the stock assessment model. These data contain many sources of variability (see Stewart et al. 2011) but results from research done in 2010 and 2014 on their representativeness show that trawl sampling and poststratification is only a small source of variability. Specifically, repeated trawls at different depths and spatial locations on the same aggregation of Pacific Hake were similar and analyses regarding the method used to stratify the data led to similar overall conclusions.

Estimated age-2+ biomass in the survey increased steadily over the four surveys conducted in 2011-2013 and 2015 (Table 12 and Figure 9). It decreased in 2017 to 1.42 Mt , then increased to 1.72 Mt in 2019, and has since decreased to 0.91 Mt in 2023. The 2023 survey age composition was made up of $50.58 \%, 24.66 \%, 8.09 \%, 5.38 \%$, and $2.92 \%$ from the 2021, 2020, 2016, 2014, and 2017 year classes, respectively. Note that the estimate of biomass does not include age- 1 fish and the age compositions used to estimate selectivity of the survey also exclude age-1 fish (Table 11). Estimates of country-specific age-2+ biomass are also provided (Table 13).

A separate relative age- 1 index (numbers of fish) is used to fit the assessment model and inform recruitment. For the 2013-2021 assessments (Hicks et al. 2013; Johnson et al. 2021), the relative age- 1 index was only explored as a sensitivity and not included in the base model. The relative index of age- 1 fish fit to in this assessment was estimated similarly to previous years. like the age-2+ index, data collected using EK 80 echosounders are scaled by a factor of 1.06 to account for differences between the EK 60 and EK 80 echosounders. Note that, in error, this scaling factor was not applied to the 2021 estimate in the 2022 (Edwards et al. 2022) and 2023 (Berger et al. 2023) assessments. The index indicates relative changes between years, not absolute values, and confirms the large year classes in 2008, 2010, 2014, and 2016, and suggests large cohorts for 2020 and 2022 (Table 12 and Figure 3).

Incorporating the relative age- 1 index results in estimates of recruitment strength that are informed on average one year earlier than models without the index (e.g., Figures 54 and G. 1 in Johnson et al. 2021). The suite of sensitivity models related to the relative age-1 index explored over the past decade indicate that its use typically informs recruitment such that the direction of cohort strength (i.e., weak, strong, or neutral) remains unchanged in subsequent assessments even after being informed by more data. The utility of an informed recruitment signal is far greater than an uninformed recruitment assumption. Finally, the Joint U.S. and Canadian Integrated Acoustic and Trawl Survey team supports its use in the stock assessment and is committed to continually evaluating and refining
approaches to improve the estimates and related uncertainty. A model without the age- 1 index was explored as a sensitivity.

### 2.3 Other data not used in this assessment

Some data sources were not included in the base model but have been explored, were used for sensitivity analyses, or were included in previous stock assessments. Data sources not discussed here have either been discussed at past Pacific Hake assessment review meetings or are discussed in more detail in the 2013 stock assessment document (Hicks et al. 2013). These primarily include those listed below.

- Fishery and survey length compositions.
- Fishery and survey age-at-length compositions.
- Calculation of a reliable fishery catch-per-unit-effort (CPUE) metric is particularly problematic for Pacific Hake, and it has never been used as an index for the assessment of this stock (see Hicks et al. 2013 for more details).
- Biomass index and age compositions from the following years of the Joint U.S. and Canadian Integrated Acoustic and Trawl Survey: 1977, 1980, 1983, 1986, 1989, and 1992.
- Bottom trawl surveys in the U.S.A. and Canada (various years and spatial coverage from 1977-2023).
- Northwest Fisheries Science Center/Southwest Fisheries Science Center/PWCC coast-wide juvenile Pacific Hake and rockfish surveys (2001-2023). However, the JTC is analysing the age-0 Pacific Hake data from these surveys in collaboration with researchers from the Southwest Fisheries Science Center and Australia, to investigate potential for developing an early indication of potential cohort strength.
- California Cooperative Oceanic Fisheries Investigations (CalCOFI) larval Pacific Hake production index, 1951-2006. The data source was previously explored and rejected as a potential index of Pacific Hake female spawning biomass. However, the JTC are exploring new avenues to utilize CalCOFI data based on recently developed methods (related to previous bullet).
- Bycatch of Pacific Hake in the trawl fishery for Pink Shrimp off the coast of Oregon (2004, 2005, 2007, and 2008).
- Historical biological samples collected in Canada prior to 1990 but currently not available in electronic form.
- Historical biological samples collected in the U.S. prior to 1975 but currently not available in electronic form or too incomplete to allow for their analysis with methods consistent with more current sampling programs.
- Northwest Fisheries Science Center winter 2016 and 2017 acoustic research surveys of spawning Pacific Hake.


### 2.4 Externally analyzed data

### 2.4.1 Weight-at-age

A matrix of empirically derived population weight-at-age data by year (Figures 12-15) is used in the current assessment model to translate numbers at age to biomass at age. Annual weight-at-age data was modelled from biological samples pooled from all fisheries and summer surveys for the years 1975 to 2023 (Figures 12-15). Samples from winter and research surveys were not included, nor were samples from near-shore areas. Past investigations into calculating weight-at-age data for the fishery and survey independently showed little impact on model results, and thus, a single matrix is used for all fleets and beginning and middle-of-the-year population weight-at-age.

New for this assessment, a generalized linear mixed model (GLMM) with a lognormal distribution was used to model weight-at-age data external to the assessment. Weight was thought to relate to a smoothed effect of age between ages zero and fifteen; random effects for cohort and year; and fixed effects for sex. The model is similar to models investigated for Walleye pollock off of Alaska, where models with correlations between age, cohort, and year were found to best fit the data (Cheng et al. 2023). Here, all unsexed fish were removed from the analysis given the small sample sizes. Weights from fish ages 15 and above for each year were pooled, and thus, ages 15-20 are assumed to have the same weight-at-age. Estimated parameters from this model were used to predict weight for ages zero to fifteen from 1975 to 2023 for each sex. The means of annual, age-specific estimates across both sexes were used for input into the assessment model. The number of samples (Figure 13) is generally proportional to the amount of catch.

The biomass at the start of a given year is based on the weight-at-age from the previous year; for example, the biomass at the start of 2022 is modelled using the empirical weight-at-age data from 2021 (Figure 12). Prior to 1975, weight-at-age input to Stock Synthesis is assumed to be equal to the mean of all available years for each respective age (1975-2023) (Figure 12). Forecast weight-at-age data are based on age-specific means from the most recent five years (2019-2023), consistent with forecast selectivity (Figure 12).

The use of empirical weight-at-age data is a convenient method to capture the variability in both the weight-at-length relationship within and among years as well as the variability in length-at-age data, without requiring parametric models to represent growth relationships. Previous attempts to explicitly model year- and cohort-specific growth were not successful for Pacific Hake and have not been revisited since Stewart et al. (2011). The empirical weight-at-age method requires the assumption that observed values are not biased by strong selectivity at length or weight and that the spatial and temporal patterns of the data sources provide a representative view of the underlying population. Simulations show that, in general, using empirical weight-at-age data when many observations are available results in more accurate estimates of spawning biomass than modeling growth (Kuriyama et al. 2016).

### 2.4.2 Maturity and fecundity

Maturity-at-age has always been modelled external to the assessment but up through the the 2023 assessment (Berger et al. 2023) the maturity ogive was time-invariant. Starting with the 2024 assessment, annual age-based maturity ogives (Figure 11) were developed using the same data, i.e., histological estimates of functional maturity, but fit with a spatiotemporal GLMM. The current data set is comprised of 2,836 ovaries with information on histological estimates of functional maturity. The samples were collected from the acoustic survey, winter and summer acoustic research trips, U.S. catcher-processor vessels by observers in the U.S. At-Sea Hake Observer Program, and the U.S. West Coast Groundfish Bottom Trawl Survey (Table 14) since 2009. Samples from south of Point Conception ( $34.44^{\circ} \mathrm{N}$ ) have been excluded from maturity analyses since 2018 (Edwards et al. 2018) because they were thought to mature at earlier ages and smaller sizes. Samples from Canada were excluded from this analysis because the spatiotemporal resolution was insufficient and led to model-convergence issues. Additional samples are available to update the maturity relationship (including samples collected from Canadian waters since 2018) but have yet to be analyzed.

The spatiotemporal GLMM used to estimate the probability of being functionally mature included an estimated spatial field, spatially varying coefficients for the quadratic function of age, and year effects as a quadratic function of subsurface ( 130.67 m ) temperature indices in the domain of the West Coast Groundfish Bottom Trawl Survey. The stochastic partial differential equation approximation to Gaussian Random Fields (Lindgren et al. 2011) was used to approximate the spatial surface with a series of estimated random effects. The temperature covariate has the potential to add mechanistic relationships to the modeling and reduce uncertainty in years where no or few samples are collected. Annual maturity-at-age was then predicted for each year since 2009 using estimated coefficients for non-ASHOP samples and day 278 (i.e., October 5th). Projections were also made forward in time to 2023 using available temperature indices. This forward projection was needed because maturity information was not sampled in 2022 and data from 2023 have yet to be analyzed. More detailed information on the modeling and projection methods is available in Appendix G.
Time-varying fecundity at age (Figure 14) was previously modeled using year-specific weight-at-age values multiplied by maturity-at-age (Berger et al. 2019). For this assessment, the maturity-at-age is also year specific. The same methods were used to estimate early (before 2009) and projection period maturity as was done for the weight-at-age data. Additionally, samples from age-15+ fish were pooled for both the maturity and weight-at-age estimation due to limited sample sizes. Consequently, the age $15+$ estimates were applied to ages 15-20 in the population dynamics model (Figure 11).

Some fish at almost every age were found to be functionally immature based on histological criteria. Older, functionally immature fish are a combination of 'skip spawners' that will not be spawning in the upcoming year and senescent fish that appear to no longer have viable ovaries. Results from ongoing research investigating the impacts of functionally immature individuals on estimates of female spawning biomass could help refine the fraction of fish mature at each age.

Tissue samples have been collected from many of the same fish from which ovaries were sampled. In the future, these tissue samples may help determine whether the fish south of $34.44^{\circ} \mathrm{N}$ are from the same population as the rest of the coastal population via genetic analyses.

### 2.4.3 Ageing error

The large inventory of Pacific Hake age determinations includes many duplicate reads of the same otolith, either by more than one laboratory or by more than one age reader within a laboratory. Recent west coast assessments have utilized the cross- and doubleread approach to generate an ageing-error matrix describing the imprecision and bias in the observation process as a function of fish age. New data and analyses were used in the 2009 assessment to address an additional process influencing the ageing of Pacific Hake, namely cohort-specific ageing error related to the relative strength of a year-class. This process reflects a tendency for uncertain age determinations to be assigned to predominant year classes. The result is that the presence of strong year classes is inflated in the age data while neighboring year classes are under-represented relative to what would be observed if ageing error was consistent with age across cohorts.

To account for these observation errors in the model, year-specific ageing-error matrices (defined via vectors of standard deviations of observed age at true age) are applied, where the standard deviations of strong year classes are reduced by a constant proportion. For the 2009 and 2010 assessments, this proportion was determined empirically by comparing double-read error rates for strong year classes with rates for other year classes. In 2010, a blind double-read study was conducted using otoliths collected across the years 2003-2009. One read was conducted by a reader who was aware of the year of collection, and therefore of the age of the strong year classes in each sample, while the other read was performed by a reader without knowledge of the year of collection, and therefore with little or no information to indicate which ages would be more prevalent. The results were analyzed via an optimization routine to estimate both ageing error and cohort effect. The resultant ageing error was similar to the ageing error derived from the 2008 analysis. Since 2011, cohort-specific ageing error has been used to reduce the ageing-error standard deviation by a factor of 0.55 for the following largest cohorts: 1980, 1984, 1999, 2010, and 2014. In the 2014 base model (Taylor et al. 2014), the 2008 cohort was also included in this set, but subsequent estimates show this year class to not be as strong as previously thought, and thus, cohort-specific ageing error has not been included for the 2008 cohort since 2015. Also, cohort-specific ageing error does not include the reduction in ageing error for age- 1 fish under the assumption that they never represent a large enough proportion of the samples to cause measurement error related to the cohort-effect.

Additional exchanges of otoliths between ageing labs within the U.S.A. and Canada are in process but were not completed in time for this assessment. The additional across-lab double reads will be informative for updating the ageing-error matrix. Unfortunately, increased protocols for moving samples across the border have led to delays.

### 2.5 Estimated parameters and prior probability distributions

Several prior distributions (Table 15) are used to fit the model. The priors that are assumed to be informative are discussed below.

### 2.5.1 Natural Mortality

Since the 2011 assessment, a combination of the informative prior for natural mortality used in previous Canadian assessments and results from analyses using Hoenig's (Hoenig 1983) method support the use of a lognormal distribution with a median of 0.20 and a standard deviation (in $\log$ space) of 0.10 . Sensitivity to this prior has been evaluated extensively in many previous assessments of Pacific Hake (see Hicks et al. (2013) for a discussion of the historical treatment of natural mortality and its prior) and is repeated here (see Section 3.8), including increasing the prior standard deviation and using an alternative prior distribution altogether based on a life history meta-analysis Hamel and Cope (2022). The Hamel-Cope prior used a lognormal prior distribution with a median of 0.22 (based on a maximum age of 25) and a standard deviation (in log space) of 0.31. Alternative prior distributions for natural mortality typically have a significant impact on the model results. But in the absence of new information on natural mortality there has been little option to update the prior.

### 2.5.2 Steepness

The prior for the steepness parameter of the stock-recruitment relationship is based on the median ( 0.79 ) and the 20th (0.67) and 80th (0.87) percentiles from the meta-analysis of the family Gadidae (Myers et al. 1999) and has been used in U.S. assessments since 2007. This prior has a beta distribution with parameters 9.76 and 2.80 , which translate to a mean of 0.777 and a log-standard deviation of 0.113 . Sensitivities to the variance on the prior on steepness were evaluated in the 2012 and 2013 assessments (Stewart et al. 2012; Hicks et al. 2013). Sensitivities to the mean of the prior are explored in this assessment (see Section 3.8).

### 2.5.3 Variability on fishery selectivity deviations

Time-varying selectivity was introduced in the 2014 assessment (Taylor et al. 2014) and is modeled using annual deviations since 1991 applied to the selectivity-at-age parameters for the fishery. A normal distribution with a fixed standard deviation ( $\Phi=1.4$; see Edwards et al. (2018) for justification) is used as a penalty function to keep deviations from straying far from zero. Selectivity for age- 0 fish is fixed at 0.0 and parameters for ages that are estimated represent the change in selectivity from the next youngest age. Beyond the age of 6 , age-specific parameters are fixed at zero giving constant selectivity beyond the last estimated value. The condition that maximum selectivity equals 1.0 results in one fewer degrees of freedom than the number of estimated parameters. Further testing of alternative methods for parameterizing time-varying selectivity (e.g., Xu et al. 2019) should be investigated in conjunction with the estimation of additional time-varying parameters.

### 2.5.4 Age composition likelihood

Since 2018, the assessment has used the linear formulation of the Dirichlet-multinomial (DM) likelihood (Thorson et al. 2017) to fit the age-composition data. Estimated parameters $\theta_{\text {fish }}$ and $\theta_{\text {surv }}$ serve to automatically adjust the weight given to the fishery- and the surveycomposition data, respectively. As of 2021, Stock Synthesis includes the constant of integration in the likelihood calculation for the D-M model such that likelihoods are comparable across weighting methods.
Integration of weighting the composition data within the assessment increases the efficiency of the assessment process, removes the subjective choice of how many iterations are required, and ensures that the results of model sensitivities, retrospective analyses, and likelihood profiles are automatically tuned, rather than having the age compositions be given the same weight as the base model. Note that the following description holds for both the survey data and the fishery data, with $\theta$ equal to $\theta_{\text {surv }}$ or $\theta_{\text {fish }}$.
The likelihood function for the linear parameterization of the D-M likelihood (see Equation 10 of Thorson et al. (2017)) is

$$
\begin{equation*}
\mathrm{L}(\boldsymbol{\pi}, \theta \mid \tilde{\boldsymbol{\pi}}, n)=\frac{\Gamma(n+1)}{\prod_{a=1}^{A_{\max }} \Gamma\left(n \tilde{\pi}_{a}+1\right)} \frac{\Gamma(\theta n)}{\Gamma(n+\theta n)} \prod_{a=1}^{A_{\max }} \frac{\Gamma\left(n \tilde{\pi}_{a}+\theta n \pi_{a}\right)}{\Gamma\left(\theta n \pi_{a}\right)} \tag{1}
\end{equation*}
$$

where $\tilde{\pi}_{a}$ is the observed proportion at age $a, \pi_{a}$ is the corresponding expected proportion at age $a$ estimated by the model, $\tilde{\pi}$ and $\pi$ designate the vectors of these proportions, $A_{\max }$ is the maximum age in the model, and $n$ is the input sample size. The parameter $\theta$ is defined as a linear scaling parameter such that $\theta n$ is the variance-inflation parameter of the D-M distribution. The linear parameterization has been shown to be superior over the saturation parameterization in simulation testing (Fisch et al. 2022), and thus, corroborates our decision to continue to use it even though the saturation parameterization is available in Stock Synthesis.
The effective sample size ( $n_{\text {eff }}$ ) associated with this likelihood is given by

$$
\begin{equation*}
n_{\mathrm{eff}}=\frac{1}{1+\theta}+\frac{n \theta}{1+\theta} \tag{2}
\end{equation*}
$$

The input sample sizes used in this assessment, which are based on the number of trips and/or hauls, are large enough that the first term is insignificant compared to the second term. Consequently, $\theta /(1+\theta)$ can be compared to the sample size multipliers used in the McAllister-Ianelli (M-I) data-weighting method (McAllister and Ianelli 1997) that was used for assessments prior to 2018 (Table 16) and as a sensitivity here (see Section 3.8). In short, the M-I method involves iteratively adjusting multipliers of the input sample sizes passed to the multinomial likelihoods until they are roughly equal to the harmonic mean of the effective sample sizes. The effective sample size is dependent on how well the
model expectation matches the observed values. Typically, this process involves no more than four to five iterations.

A uniform prior between -5 and 20 for $\log \theta_{\text {fish }}$ and $\log \theta_{\text {surv }}$ tends to lead to inefficient sampling of $\log \theta_{\text {surv }}$ because many samples occur in a part of the parameter space where the effective sample size multiplier, $\theta_{\text {surv }} /\left(1+\theta_{\text {surv }}\right)$, is between 0.99 and 1.0 (Berger et al. 2019). In that area, the input sample sizes given the uniform prior have full weight and the likelihood surface is almost completely flat with respect to $\log \theta_{\text {surv }}$. The current prior on $\log \theta_{\text {surv }}$ can be associated with an approximately uniform prior of the weight $\theta_{\text {surv }} /\left(1+\theta_{\text {surv }}\right)$, where the parameters of the normal distribution were back-calculated from a uniform distribution with the bounds of 0 and 1 (Grandin et al. 2020). The normal prior for both $\log \theta_{\text {fish }}$ and $\log \theta_{\text {surv }}$ has a mean of 0 and a standard deviation of 1.813.

Composition data can also be weighted using the Francis method (T2.6 in Table 2 of Francis 2011), which is based on variability in the observed ages by year. This method, like the M-I method, is iterative, where the sample sizes are adjusted such that the fit of the expected mean age should be within the estimated uncertainty at a rate that is consistent with the variability expected given the effective sample sizes. The Francis method is known to be sensitive to outliers and prone to convergence issues when selectivity varies with time. As a result, the Francis method was not included as a sensitivity.

## 3 ASSESSMENT

### 3.1 Modeling history

In spite of the relatively short history of fishing, Pacific Hake have surely been subject to a larger number of stock assessments than any marine species off the west coast of the U.S.A. and Canada. These assessments have included a large variety of age-structured models. Initially, a cohort analysis tuned to fishery CPUE was used (Francis et al. 1982). Later, the cohort analysis was tuned to National Marine Fisheries Service (NMFS) triennial acoustic survey estimates of absolute biomass at age (Hollowed et al. 1988). Since 1989, Stock Synthesis models (or base versions of it) fit to fishery catch-at-age data and acoustic survey estimates of population biomass and age composition have been the primary assessment method.

While the general form of the age-structured assessment has remained similar since 1991, modeling procedures have been modified in a variety of ways. There have been alternative data choices, post-data collection processing routines, data-weighting schemes, structural assumptions for the stock assessment model, MCMC sampling algorithms, and control rules (Table 16). Analysts are constantly trying to improve the caliber and relevance of the assessment by responding to new scientific developments related to statistics and biological dynamics, policy requirements, and different or new insights brought up during the peer review process to ensure a robust stock assessment.

Data processing, filtering, and weighting choices have been modified several times since the first assessment. For example, modifications to the target-strength relationship used to scale acoustic data changed in 1997 (Dorn and Saunders 1997), and kriging was im-
plemented to account for the spatial correlation in the acoustic data in 2010 (Stewart and Hamel 2010). While survey data have been the key index for biomass since 1988, surveys that have been used have varied considerably. The Alaska Fisheries Science Center/Northwest Fisheries Science Center West Coast Triennial Shelf Survey was used from 1988 before being discarded from the 2009 assessment (Hamel and Stewart 2009). Acoustic surveys from the years prior to 1995 were used for assessments in the early 1990s, but Stewart et al. (2011) reviewed these early surveys and deemed that sampling was insufficient to be comparable with more recent data. Several recruitment indices have been considered but ultimately none were identified as adding appreciable contribution to model results (Helser et al. 2002; Helser et al. 2005; Stewart and Hamel 2010), except for the fishery-independent acoustic-based relative age- 1 index that has been included in the base model since the 2022 assessment. The process for generating fecundity-at-age from the combination of weight-at-age and maturity data changed in 2019 from using time-invariant to year-specific weight-at-age values. In 2024, time-varying maturity ogives were also added to the calculation of fecundity-at-age (see Section 2.4.2 for details). Even where data have been consistently used, the weighting of these data in the statistical likelihood has changed through the use of various emphasis factors (e.g., Dorn 1994; Dorn et al. 1999), a multinomial sample size on age compositions (e.g., Dorn et al. 1999; Helser et al. 2002; Helser et al. 2005; Stewart et al. 2011), internal estimations of effective sample size using the Dirichlet-multinomial distribution (Edwards et al. 2018), and assumptions regarding year-specific survey variance. Since 2021, a more computationally efficient Bayesian MCMC sampler [No-U-Turn Sampler; NUTS; Hoffman and Gelman (2014)] was used to estimate posterior distributions (Monnahan and Kristensen 2018; Monnahan et al. 2019), a change from previous assessments that used the random walk Metropolis Hastings (rwMH) sampler (details described in Johnson et al. 2021). The list of changes discussed above is for illustrative purposes only and represents a small fraction of the different choices analysts have made and that reviewers have required.

The structure of the assessment models has perhaps had the largest number of changes. In terms of spatial models, analysts have considered spatially explicit forms (Dorn 1994, 1997), spatially implicit forms (Helser et al. 2006), and single-area models (Stewart et al. 2012). Predicted recruitment has been modeled by sampling historical recruitment (e.g., Dorn 1994; Helser et al. 2005), using a stock-recruitment relationship parameterized using maximum sustainable yield (MSY) and the fishing mortality rate estimated to produce the MSY [ $F_{\text {MSY }}$; Martell (2010)], and using several alternative steepness priors (Stewart et al. 2012; Hicks et al. 2013). Selectivity has also been modeled in several ways, invariant (Stewart et al. 2012; Hicks et al. 2013), time-varying with (Helser et al. 2002) and without (Dorn 1994; Dorn and Saunders 1997; Stewart et al. 2012; Hicks et al. 2013) a random walk, alternative levels of allowable deviation through time (Hicks et al. 2013; Berger et al. 2017), age-based (Dorn 1994; Dorn and Saunders 1997; Stewart et al. 2012; Hicks et al. 2013), and length-based (Helser and Martell 2007).

Several harvest control rules have been explored for providing catch limits from stock assessment output. Pacific Hake stock assessments have presented decision makers with constant $F$, variable $F$, and the following hybrid control rules: $F_{\mathrm{SPR}=35 \%}, F_{\mathrm{SPR}=40 \%}, F_{40 \%}-40: 10$, $F_{\mathrm{SPR}=45 \%}, F_{45 \%}-40: 10$, and $F_{\mathrm{SPR}=50 \%}$ (e.g., Dorn 1996; Hicks et al. 2013). Changes to poli-
cies such as the United States' National Standards Guidelines in 2002 and the $F_{40 \%}-40: 10$ harvest control rule in the Agreement (Appendix C) have required specific changes to control rules.

In addition to the examples given above and changes documented in stock assessments, there have been many more investigations conducted at review panel meetings. Starting in 2013, the addition of the MSE (Hicks et al. 2013; Jacobsen et al. 2021) facilitated investigating changes to the modeling procedure in terms of pre-specified objectives that aim for a sustainable coast-wide fishery.

### 3.2 Description of base model

The 2024 base model has the same general population dynamics structure as the 2023 assessment's base model. The statistical-catch-at-age model assumes that the Pacific Hake population is a single coast-wide stock subject to one aggregated fleet with combined male and female population dynamics. Stock Synthesis (Methot, Jr. and Wetzel 2013) version 3.30 .22 was the modeling platform used. The largest changes between the 2023 and 2024 stock assessments are the addition of another year of fishery and survey data, an age- 1 index data point, a model-based method for empirical weight-at-age data, and time-varying maturity information integrated into the base model.

The 2024 base model includes a time series (1995 to 2023) of acoustic age-2+ biomass estimates and acoustic estimates of relative numbers of age- 1 fish (see Section 2.2.1 for more details on the age-1 index). Maturity is assumed to be time-invariant prior to 2009, and time-varying, with the integration of annual maturity ogives informed by sea temperature at depth, since 2009 (see Section 2.4.2). Fecundity is time-varying as defined by annual weight-at-age multiplied by annual maturity ogives (1975-2023; additionally see Section 2.4.1). The D-M likelihood approach (Thorson et al. 2017) is used to estimate the weights associated with age-composition data, rather than iteratively tuning the sample size multiplier as in 2017 and earlier assessments (see Section 2.5.4). Time-varying fishery selectivity is retained in the 2024 base model with the magnitude of the allowable deviations unchanged from the 2023 base model (see Section 2.5.3). The general parameterization of selectivity was retained, although additional parameters were required to estimate an additional year of deviations. The selectivity of the acoustic survey is assumed to be time invariant. Selectivity curves were modeled as non-parametric functions estimating age-specific values for each age beginning at age two for the index of age- $2+$ biomass and age one for the fishery until a maximum age of 6 , after which all ages are assumed to have the same selectivity. Selectivity for the relative age- 1 index was set to one for age one and zero for all other ages.
Prior probability distributions are used for a select few parameters and fixed values are used for several parameters. For the base model, the instantaneous rate of time-invariant natural mortality $(\mathrm{M})$ is estimated with a lognormal prior having a median of 0.20 and a standard deviation (in log-space) of 0.1 (see Section 2.5.1). The stock-recruitment relationship is a Beverton-Holt parameterization, with the log of the mean unexploited recruitment $\left(\log R_{0}\right)$ freely estimated. This assessment uses the same beta-distributed prior for stock-recruitment steepness (h), based on Myers et al. (1999), that has been
applied since 2011 (Stewart et al. 2011). Year-specific recruitment deviations were estimated from 1966-2022. The standard deviation, $\sigma_{r}$, of recruitment variability serves as a recruitment deviation constraint and is fixed at 1.4 in this assessment. This value is based on consistency with the observed variability in the time series of recruitment deviation estimates and is the same as assumed in assessments from 2013 to 2023 (Table 16). Catchabilities associated with the biomass index $\left(q_{\mathrm{b}}\right)$ and with the relative age- 1 index $\left(q_{1}\right)$ were calculated analytically as per Ludwig and Walters (1981) for each sample of posterior parameters, resulting in a distribution of catchability for each.

Statistical likelihood functions used for data fitting are typical of many stock assessments. The biomass index was fit via a lognormal likelihood function, using the observed (and extra 2009) sampling variability, estimated via kriging, as year-specific weightings. The relative age-1 index was specified as having a Student's t-distribution for its error structure with the number of degrees of freedom equal to one less than the number of available data points. An additional constant and additive standard deviation on the log-scale component is included for both the biomass index and the relative age- 1 index, which were freely estimated to accommodate unaccounted-for sources of process and observation error. A D-M likelihood was applied to age-composition data, with input sample sizes equal to the sum of the number of trips and hauls sampled across all fishing fleets or the number of trawl sets in the research surveys (see Section 2.5.4).

Model results and statistical inference were based on 8,000 MCMC samples [using the adnuts R package; Monnahan and Kristensen (2018)] compiled across 8 chains, each with a 250 sample burn in period, to describe posterior distributions for model parameters and derived quantities. The number of samples used for bridging models, sensitivity models, and retrospective models was also 8,000. Medians ( $50 \%$ quantiles) are reported together with the bounds of $95 \%$ credibility intervals calculated as the $2.5 \%$ quantile and the $97.5 \%$ quantile of posterior distributions from the MCMC samples, to give equal-tailed intervals. A full explanation of the NUTS algorithm and the adnuts package, including an analysis with the Pacific Hake stock can be found in Monnahan et al. (2019).

### 3.3 Response to 2023 Scientific Review Group (SRG) review

The Scientific Review Group (SRG) meeting was held from February 7-10th, 2023, at the Graduate Hotel, Seattle, WA, U.S.A.

The following are the 'SRG Recommendations and Conclusions for the Stock Assessment' from the 2023 SRG report and the associated responses from the JTC:

1. Pacific Hake dynamics are highly variable even without fishing mortality. The SRG applauds the efforts of the JTC and the MSE Working Group to add capabilities for estimating dynamic reference points within the assessment and MSE, and encourage those groups to jointly develop alternative reference points, including dynamic reference points, for future SRG consideration.

Response - An evaluation of alternative reference points and how they may best be used in Pacific Hake management continues to be an important area for ongoing research. Future reference point discussions stemming from simulation work, preferably through
the MSE, would be beneficial. The JTC, nor the MSE Working Group, have had the capacity to initiate simulations that explore alternative reference points, including dynamic reference points, to date. Several alternative simulation-based study designs have been fleshed out but the resources to complete such research is currently the limiting factor.
2. The SRG also recommends continuing sensitivities for steepness, natural mortality, $\sigma_{R}$, excluding the age-1 index, alternative standard deviations for time-varying selectivity, and using the McAllister-Ianelli method to weight fishery age- composition data.
Response - The JTC has conducted all of the requested sensitivities and provides summaries in written (Section 3.8), tabular (beginning with Table 32), and graphical (beginning with Figure 48) formats in this document. Many other model explorations were conducted during the development and exploration of the base model to understand model performance and sensitivity to data and structural decisions.
3. The SRG recommends that the JTC include dynamic unfished spawning biomass in the 2024 assessment as a comparator with the equilibrium unfished spawning biomass used to provide management advice. The SRG also encourages the continued outreach regarding the use of dynamic reference points to stakeholders and managers, including identifying pros and cons of using dynamic unfished spawning biomass.
Response - Two figures have been added to the assessment document. The first shows the unfished spawning stock biomass time series relative to the standard 'fished' spawning stock biomass time series (see Figure 41). The second shows a comparison of relative spawning biomass when spawning biomass in year $t$ is related to unfished equilibrium biomass, $B_{0}$ (static $B_{0}$; as in Figure 31) and when spawning biomass in year $t$ is related to the unfished biomass time series in year $t$ (dynamic $B_{0}$; see Figure 42). These can be used to visually compare equilibrium versus non-equilibrium assumptions and estimated changes in population dynamics in the absence of fishing.

The JTC once again delivered a presentation on dynamic reference points at the October 2023 JMC and AP meeting as a way to provide education and outreach opportunities related to dynamic reference point methods and approaches.
4. The SRG recommends that the JTC explore alternative ways of estimating natural mortality to update the current approach in the model, which is based on methods from more than a decade ago, since newer methods are available. Information presented during the SRG meeting implies that natural mortality on age-2 Pacific Hake is higher than currently assumed in the assessment model and should be explored more fully.

Response - The JTC investigated estimating a vector of natural mortality parameters instead of a single age-invariant parameter. Several configurations, i.e., breakpoints at different ages, were explored but no combination of breakpoints led to natural mortality decreasing with age as hypothesized given the diet data of predators of Pacific Hake. Instead parameters were estimated near the median of the prior. When the prior was
removed, natural mortality at younger ages was estimated to be lower than natural mortality at older ages largely due to the lack of information in the data. If age-specific natural mortality is desired in the future, tagging studies could be used to help inform the parameters. Or, diet data could be used to externally estimate natural mortality at age. Other modeling frameworks, such as the Woods Hole Assessment Model have more flexibility in modeling natural mortality and might also be an option for future explorations.
5. The SRG encourages an analysis of catch and CPUE distribution for Canada and US that examines latitudinal shifts in fishing over time.
Response - The JTC has not officially investigated CPUE data since prior to the 2013 assessment because of the many difficulties in accurately characterizing catch rates given the difficulties inherent in calculating effort within the Pacific Hake fishery (Hicks et al. 2013). Furthermore, vessel locations for the U.S. shore-based sector must be extracted from log-book information, which is known to be problematic, and current data confidentiality agreements do not allow the sharing of vessel-specific information amongst the JTC members making it difficult to do a comprehensive analysis. Maps of CPUE for the U.S. at-sea sectors and Canadian vessels were presented in the data presentation at the SRG meeting as a first take at fulfilling this request.
6. The SRG recommends continued work to collect ovary samples and data to develop a picture of the Pacific Hake reproductive cycle both seasonally, interannually and at the life-time scale based on histological and physiological measurements. Given the inter-annual variability in age-at-maturity, the SRG recommends that the JTC explore ways to incorporate time-varying maturity in the stock assessment model.
Response - Melissa Head has been working hard to continuously collect maturity information for Pacific Hake since 2009. Up until this assessment, maturity was included in the fecundity relationship as a single ogive. Maturity-at-age ogives are now estimated for each year since 2009 and used to calculate fecundity (i.e., maturity * weight at age). Moreover, the relationship between age and maturity includes a coefficient for temperature (see Appendix G). This relationship could be enhanced in the future by including samples from Canada, samples that have previously been analyzed, collected samples that have yet to be analyzed, and samples collected during future endeavors. During the JTC meeting, it was also brought to our attention that there is the potential for samples to be collected from the U.S. shoreside sector as well. The JTC, Melissa Head, and the Sarah Nayani are investigating the feasibility of this in attempts to increase the seasonal coverage of sampling, though winter months will still remain sparsely sampled without a dedicated winter survey.
7. The SRG noted that the age- 1 index did not include a value for 2001 because it was zero. Although this decision had negligible influence on the results because the estimate for 2000 recruitment was close to zero, the SRG noted that Stock Synthesis uses a lognormal likelihood which does not handle zero values. Given that future zero values are expected to have a bigger influence on the results in the short-term, the SRG recommends that the JTC explore likelihood forms that
can fit to very low index values from the age-1 survey (e.g., robust likelihood). The SRG acknowledges that implementing new likelihoods will require changes to the Stock Synthesis platform.
Response - A Student's t-distribution was implemented for the relative age- 1 index of abundance. The Student's $t$-distribution cannot accommodate values of zero but it does allow for fatter tails than the lognormal distribution. Secondly, the Student's $t$-distribution is known to provide more accurate estimates of the variability of the sample mean for small sample sizes than the lognormal distribution when the standard deviation is unknown or imprecise. In the future, additional distributions should be investigated that can accommodate values of zero but this change was seen as a positive step forward until the survey team can estimate the variability of the survey.
8. Given the importance of the age-1 index in estimating the size of the age-classes entering the fishery, the SRG recommends that the JTC implement updated age-1 index CVs, when they are provided by the Survey Team, in the stock assessment model.

Response - The joint survey team continues to be short staffed. During 2023, priority was given to completing the 2023 survey, including providing the 2023 survey biomass estimate, associated age compositions, and the 2023 relative age- 1 index in a timely manner for the stock assessment. Age-1 index CVs were not available for inclusion in the 2024 assessment; so, the fixed values of 0.5 were retained but a Student's $t$ distribution was used to fit the survey instead of a lognormal distribution (see the above response).
9. A new at-sea sampling program in the Canadian freezer-trawler fleet was implemented in 2022 involving vessel crews sampling 50 fish per tow from two tows per trip. The SRG recommends that Canada consider sampling fewer fish from more tows to spread the sampling out and provide a more representative sample of fishery catches.
Response - This was discussed briefly in a meeting involving industry representatives and one member of the Canadian JTC and determined to be too much to ask of crew at the time. They would have to deal with more than two bags on a trip which would lead to space limitations in the freezer that they are currently using for the purpose and organizational difficulties. The space limitations could be overcome by storing bags of fish in the galley but after a short discussion on the idea, it was determined not to be feasible for food safety and storage-space reasons. The idea will be introduced again for the 2024 fishing season.
10. The SRG encourages the JTC to consider methods to determine the maximum input sample size for the age compositions (e.g., Stewart and Hamel 2014).

Response - Determining input sample sizes is important for how annual fishery and survey age compositions are initially weighted, which then provides the basis from which wholesale re-weighting of data sources (fishery or survey) is done with the Dirichlet-multinomial data weighting model parameters. The JTC has considered alternative methods, including that of Stewart and Hamel (2014), and has determined
that the first step is to explore the handling of survey age-composition data (given that initial effective sample sizes and data source weighting is relative to other data sources in the model). Currently, survey age compositions represent age structure associated with the acoustic survey as viewed through an estimated selectivity curve for the acoustic-trawl sampling net. Yet, selectivity for ages two and older with acoustics is theoretically at or near one. The JTC plans to investigate whether there is a more informative way to utilize survey age-composition information in the stock assessment model in the coming year.
11. The SRG notes that there have been multiple strong cohorts in the stock recently compared to earlier periods where there was only one strong cohort supporting the stock, including during the period of sample collection for the ageing error matrix that supports the assessment model. Therefore, the SRG encourages the resumption of the ageing error study by the Committee of Age Reading Experts (CARE) using samples collected during the past decade.

Response - An ageing error study in conjunction with CARE has been planned for several years but a full exchange remains on hold due to continuing difficulties with permits to send biological specimens across the U.S.-Canada border. The JTC plans to move forward with updating the ageing error analysis in the coming year with the data that are available regardless of the status of the in progress CARE study.
12. The SRG supports the investigation of alternative selectivity functions, which may include a two-dimensional autoregressive (AR) approach, which may use information from the previous year and from adjacent year classes to predict selectivity.

Response - The JTC agrees that investigations into alternative selectivity functions is a priority research area for Pacific Hake. The JTC would like to make incremental progress on this in the coming year as time allows, including also looking into alternative methods (e.g., random effects models) to best capture age, year, and cohort effects.

### 3.4 Modeling results

### 3.4.1 Changes from 2023

A set of 'bridging' models was constructed to evaluate the component-specific effects of the steps to change from the 2023 base model to the 2024 base model. The steps are as follows:

- Update to the latest version of Stock Synthesis, version 3.30.22, to follow current best practices;
- Incrementally update catch, weight-at-age, age-1 index, and fishery age-composition data from years previous to 2023 (in that order);
- Incrementally add 2023 catch, weight-at-age, biomass survey, survey agecomposition, age- 1 index, and fishery age-composition data (in that order);
- Change the error distribution associated with the age-1 index from lognormal to Student's t-distribution;
- Use a model-based matrix of input weight-at-age data; and
- Incorporate model-based estimates of time-varying maturity ogives into the calculation of time-varying fecundity.

Stock Synthesis version 3.30.22 includes a number of changes since the version used by Berger et al. (2023). However, none of the changes were specifically relevant to this assessment, and thus, the software update had no effect on assessment results (Figure 16).

The update of pre-2023 data occurs because databases are continually updated; this yielded minor adjustments to the data. For example, samples that were recently aged but not available for the 2023 assessment were included. Updates to pre-2023 data were small enough that they had little impact on the model results.

The addition of the 2023 catch and weight-at-age data extends the model to the start of 2024. Recruitment estimates and historical stock trajectory were relatively unchanged, and the new data suggest a slight decrease in female spawning biomass from 2023 to 2024 (Figure 16).

Including the 2023 fishery-independent biomass estimate led to a downward shift in the stock trajectory going back to 2017 and a similar shift in the fit to the survey (Figure 16). The addition of the 2023 survey biomass age compositions led to an estimated increase in stock biomass from 2023 to 2024 as a result of small shifts in expected recruitment, particularly for the 2020 and 2021 year classes. While the addition of the 2023 relative age- 1 index had a negligible effect on the stock trajectory, it did slightly adjust estimates of recent recruitment strength, in particular raising the 2022 recruitment somewhat.

The final step of adding 2023 data involved incorporating fishery age-composition information, which shifted the ending year of the deviations in the selectivity parameters from 2022 to 2023. These data had relatively little impact on the historical biomass estimates but did shift recent estimates of spawning biomass upwards (Figure 16). Recruitment increased in 2021 and 2022, while the 2020 cohort was reduced. The increase in 2021 and 2022 recruitment contributes to the increase in female spawning biomass by the start of 2024, as these fish are considered mostly mature at the start of 2024. Despite both fishery age compositions and the relative age-1 index pointing towards above average cohorts in 2020, 2021 and, to a lesser extent, 2022, estimates of 2024 female spawning biomass remain highly uncertain (Figure 16).

Structurally changing the error distribution for the relative age-1 index from a lognormal to a Student's t-distribution (with 13 degrees of freedom) had negligible effect on assessment results (Figure 17). Nonetheless, the t-distribution is better suited for these data given the low sample size (14) and broader distribution tails compared to a lognormal when sample sizes are less than 30 .

Input weight-at-age data were constructed using a model-based approach (see Section 2.4.1 for details) to better inform changes in weight-at-age for years and ages when there is little or no data. Model predictions were based on a smoothed fixed effect for age and random effects for year and cohort. During periods of more consistent sampling protocols (since the mid-1990s), there was little effect on overall stock size or status (Figure 17). However,
there were refinements to stock size and status during the 1970s and 1980s, a period when sampling protocols were not as well documented (e.g., from foreign vessels).

Lastly, model-based estimates of time-varying maturity ogives were incorporated into the calculation of time-varying fecundity (Section 2.4.2 and Appendix G). This is in addition to using time-varying weight-at-age inputs, which improves the assessment model inputs that translate total biomass to spawning biomass. The addition of time-varying maturity resulted in annual differences in the age at which Pacific Hake mature, such that there were minor shifts in the translation of total biomass to spawning biomass by one year in some cases within the time series (Figure 17), but the general trend in spawning biomass and population status remained largely the same.

### 3.4.2 Assessment model results

## Model Fit

Stationarity of the posterior distribution for model parameters was assessed via a suite of standard single-chain and multi-chain diagnostic tests via graphical summaries and interactive web applications (ShinySTAN; Appendix A). All estimated parameters showed good mixing during sampling, no evidence for lack of convergence, and acceptable autocorrelation (results for some key parameters are shown in Figures A.1-A.4). Correlationcorrected effective sample sizes were sufficient to summarize the posterior distributions and neither the Geweke nor the Heidelberger and Welch statistics for these parameters exceeded critical values more frequently than expected via random chance (Figure A.5). The Gelman-Rubin multi-chain diagnostic test, which compares within-chain variance to among-chain variance, further indicated that convergence was adequately achieved (examined via ShinySTAN). Correlations among key parameters were generally low, with the exception of $M$ and $\log R_{0}$ (Figure A.6). Estimates of recruitment in 2014 and 2016 were correlated with the derived quantity of catch from the default harvest rule in 2024, as to be expected given the dependencies among these quantities (Figure A.6). An examination of deviations in recruitment (log-scale differences between estimated and expected recruitment values) from recent years (Figure A.7) indicates the highest correlation (0.92) was between the 2014 and 2016 recruitment deviations. This is the same as in the last assessment despite the fact that each cohort has been observed for an additional year.
Regarding the Dirichlet-multinomial parameter $\theta_{\text {fish }}$, the estimate (median and $95 \%$ credible interval) for $\log \theta_{\text {fish }}$ was -0.663 ( $-0.853--0.470$ ), giving an effective sample size multiplier $\theta_{\text {fish }} /\left(1+\theta_{\text {fish }}\right)$ of $0.340(0.299-0.385)$. The related $\log$ of the survey age-composition parameter $\theta_{\text {surv }}$, i.e., $\log \theta_{\text {surv }}$, was 2.770 (1.541-4.986), and the resulting effective sample size multiplier $\theta_{\text {surv }} /\left(1+\theta_{\text {surv }}\right)$ of 0.941 ( $\left.0.824-0.993\right)$.
The base model fit to the acoustic survey biomass index (Figure 18) remains similar to the 2023 base model, up to 2017. The low 2023 survey biomass pulls down the last few years of estimated biomass, such that the fit to the 2019 data point is very good (for the 2023 assessment it was overestimated), the 2021 fit is underestimated (for the 2023 assessment it was very good). The median of the posterior distribution for the analytically-derived catchability associated with the acoustic survey biomass index ( $\mathrm{q}_{\mathrm{b}}$ ) was 0.838 (Figure 20).

The 2023 biomass index is the third lowest in the series (Table 12), and is well below the model estimate, similar to the 2001 index that has always been below model estimates (Berger et al. 2023). While no direct cause for the 2001 index anomaly is known, the survey did begin earlier that year than all other surveys between 1995 and 2009 (Table 12), which may explain some portion of the anomaly, along with age structure. For 2023, the survey timing is not anomalous. The estimated biomass increase from 2023 to 2024 is driven by the addition of 2023 survey age-composition data (Figure 16). The addition of the 2023 relative age- 1 index suggested an above-average 2022 cohort (and also increased the 2021 relative age- 1 index compared to the previous assessment due to a previous omission; Section 2.2.1).

The relatively stable estimated biomass from 2013-2019 is unchanged from the previous assessment. The underestimation of the 2009 and 2023 biomass estimates are larger than the underestimation of any other year. The uncertainty of the 2009 value (both modeled and actual) is high because of the presence of large numbers of Humboldt Squid during the survey. Humboldt Squid have similar target strength to hake which could introduce bias in the biomass estimate for that year, which also likely influenced the population dynamics of Pacific Hake through predation in that year. Future data will reduce the large uncertainty in the 2023 biomass estimate, which may reduce the underestimation.

Differences between the median posterior density estimates from the fit to the survey index are likely due to slight differences in what the fishery composition data and survey composition data, when considered independently, would otherwise suggest as population trends. Additionally, the population has undergone recent high, but declining, catch levels and produced a couple of above-average cohorts that are now mature.

The base model fit to the relative index of age-1 fish highlights an overall general confirmation of relative cohort strength (Figure 19). In particular, the 2008, 2014, and 2018 cohorts were estimated to be less than the index, while the 1994 and 2016 cohorts were estimated to be larger than indicated by the index. The 2011 value (the large 2010 cohort) was closely fit. Age-1 fish in 2021 (2020 cohort) were estimated slightly below the index value (in last year's assessment they were estimated slightly above) and, being so young, include a large amount of uncertainty. The median of the posterior distribution for the analytically-derived catchability associated with the age-1 index ( $q_{1}$ ) was 0.490 (Figure 21).
Fits to the age-composition data continue to show close correspondence to the dominant and small cohorts observed in the data when the data give a consistent signal (Figures 22 and 23). Because of the time-varying fishery selectivity, the fit to commercial age-composition data is particularly good, although models with time-invariant selectivity used in previous years also fit the age compositions well. In the 2023 fishery, the 2021 cohort was the largest ( $35 \%$ ), followed by the 2020 cohort ( $25 \%$ ), and then the 2016 cohort (13\%). Age compositions from the 2023 acoustic survey suggest a similar age structure for older fish.

The 2020 cohort has now been observed by, and is well fit by, the acoustic survey (Figure 23), with the survey's inclusion decreasing its estimated size (Figure 16). Combined, the 2015-2023 fishery age-composition data and the 2017-2023 acoustic survey age-composition data suggest that 2014 was a strong recruitment year, and the model was
able to adequately fit to these observations (Figure 23). The 2016 cohort, which has been observed three times by the survey, still appears to be smaller than the 2014 cohort.

The 2023 survey was the first to sample the 2021 cohort, suggesting that it was a large contingent of the population ( $50.6 \%$ of the 2023 survey catch). The 2020 cohort, which has now been observed by the acoustic survey, is expected to be above average in size. Residual patterns to the fishery and survey age data do not show patterns that would indicate systematic bias in model predictions (Figure 24).

The median estimates for numbers, biomass, exploitation rate, and catch (in numbers and in biomass) for each age class in each year are given in Tables 17-21. For the major cohorts, the resulting estimated age-specific catch, natural mortality, and surviving biomasses are given in Table 22. For example, at age-5 the catch weight of the 2014 cohort was slightly more than that of the 2010 cohort, and the resulting surviving biomass of the 2014 cohort was approximately half of the surviving biomass of the 2010 cohort.

Posterior distributions for both steepness and natural mortality are influenced by priors (Figures 25-26). The posterior for steepness is only slightly updated by the data, as expected given the low level of information available to inform steepness as found in previous hake assessments. The posterior of natural mortality, on the other hand, is shifted to the right of the prior distribution and the prior may be constraining the posterior distribution from shifting further. Broadening the prior distribution by increasing the prior standard deviation for the natural mortality parameter is examined in sensitivity runs (see Section 3.8). Other parameters showed updating from diffuse priors to posterior distributions, including $\log \theta_{\text {fish }}$ and $\log \theta_{\text {surv }}$ (as outlined in Section 2.5.4).

The 2024 base model specified the same level of variation (standard deviation of $\Phi=$ 1.4) associated with time-varying fishery selectivity as the 2023 base model, effectively allowing the model flexibility (i.e., a lower penalty on the overall likelihood) to fit to data that suggests high variability among years for each age. This level of variation led to results that remained consistent with the 2023 acoustic survey age-composition data (but not the biomass index) and gave reasonable fits to the fishery age-composition data, given that there is considerable uncertainty associated with spatial changes in fish availability (due to movement) and recent variability in oceanographic conditions. Estimated selectivity deviations for age- 3 and age- 4 fish are larger from 2010 to 2012 than in subsequent years until 2020 when the deviation for age- 4 was large again (Figures 27 and 28). The median selectivity peaks at age-4 in 2010, 2012 and 2020 and at age-3 in 2011 suggesting targeting (or generally higher availability) of the younger cohorts in those years. This pattern is consistent with the 2008 cohort appearing strong in the fishery age compositions initially, but decreasing in prominence from 2013 onward (Figure 22). Fishery selectivity on age-2 fish was at its highest in 2016. Fishery selectivity for 2023 was characterised by a local peak at age 3 rather than a logistic pattern (Figure 28), likely as a result of increased availability of the above-average 2020 cohort. Even though the survey selectivity is time invariant, the posterior shows a broad band of uncertainty between ages 2 and 5 (Figure 29). The decline in survey selectivity between ages 3 and 4 may be an artifact of the interaction between large cohorts and the biennial timing of recent surveys, with the 2010, 2014, 2016, and 2020 cohorts occurring in the survey at ages 3 and/or 5 but not age 4 . Fishery selectivity is
likewise very uncertain (Figures 28 and 29), but in spite of this uncertainty, changes in year-to-year patterns in the estimates are still evident, particularly for age-2, age-3, and age- 4 fish, though these patterns might also reflect time-varying mortality processes.

## Stock biomass

The base stock assessment model indicates that, since the 1960s, Pacific Hake female spawning biomass has ranged from well below to above unfished equilibrium (Figures 30 and 31 and Tables 23 and 24). The model estimates that it was below the unfished equilibrium in the 1960s, at the start of the assessment period, due to lower than average recruitment. The stock is estimated to have increased rapidly and was above unfished equilibrium in the mid-1970s and mid-1980s (after two large recruitments in the early 1980s). It then declined steadily to a low in 1999. This was followed by a brief increase to a peak in 2003 as the very large 1999 year class matured. The 1999 year class largely supported the fishery for several years due to relatively small recruitments between 2000 and 2007. With the aging 1999 year class, median female spawning biomass declined throughout the late 2000s, reaching a time-series low of 0.616 million $t$ in 2009. The assessment model estimates that median female spawning biomass then peaked again in 2014 due to a very large 2010 year class and an above-average 2008 year class. The subsequent decline from 2014 to 2016 is primarily from the 2010 year class surpassing the age at which gains in weight from growth are greater than the loss in weight from mortality (growth-mortality transition). The 2014 year class is estimated to be large, though not as large as the 1999 and 2010 year classes, resulting in an increased biomass in 2017. The estimated biomass mostly declined from 2018 to 2022 due to the 2014 and 2016 year classes moving through the growth-mortality transition during a period of high catches. The increase in female spawning biomass in 2023 and 2024 is due to the expected above-average 2020 and potentially large 2021 cohorts entering maturity, and the recent declining trend in catch.

The median estimate of the 2024 relative spawning biomass (spawning biomass at the start of 2024 divided by that at unfished equilibrium, $B_{0}$ ) is $99 \%$. However, the uncertainty is large, with a $95 \%$ posterior credibility interval from $45 \%$ to $230 \%$ (Tables 23 and 24), partly due to remaining unknowns about the size of the potentially large 2021 cohort because the acoustic survey has only provided one year of information about it.

The median estimate of the 2024 female spawning biomass is 1.885 million $t$ (with a $95 \%$ posterior credibility interval from 0.853 to 4.828 million $\mathfrak{t}$ ). The current estimate of the 2023 female spawning biomass is 1.335 ( $0.652-3.225$ ) million $t$, giving less uncertainty than the estimate from the 2023 assessment of 1.910 (0.757-5.610) million $t$. The current median is reduced from last year, partly due to the tail of the distribution being greatly curtailed (upper end of the interval is much lower than it was in the 2023 assessment), and a slight lowering of the lower end of the interval. The decrease appears to be due to the addition of the age- $2+$ biomass index pulling down the estimated biomass for recent years, plus the addition of the survey age compositions lowering the estimated 2020 recruitment (Figure 16).

## Recruitment

The new data for this assessment do not significantly change the general pattern of recruitment estimated in recent assessments. However, estimates of absolute recruitment for the most recent years can change with new data, and the improved methods for modeling temporal weight-at-age and spatio-temporal maturity can slightly change some historical estimated recruitments.

The estimate of 2020 recruitment in last year's assessment was based on only two years of data and thus was highly uncertain; it suggested the cohort could potentially be huge ( $95 \%$ credible interval: 2.9-47.6 billion fish). But with the extra data in this year's assessment the 2020 cohort looks to be above average but not huge ( $95 \%$ interval: $2.1-12.7$ billion fish). The median has consequently fallen from 11.4 to 4.7 billion fish between the two assessments.

Similarly, median estimates of 2019 recruitment have changed by $-55 \%$ (which is only 0.3 billion fish because 2019 was already estimated to be a small year class).

The 2021 recruitment is now estimated to be potentially large, whereas it was estimated to be below average in last year's assessment (for which the only information was the proportions of age- 1 fish caught in the 2022 commercial fishery). The $95 \%$ credible interval in the 2023 assessment was $0.03-6.91$ billion fish), expanding in the current assessment to 4.1-29.5 billion fish). Consequently, the median has increased by $2,162 \%$ ( 9.7 billion fish). The general notion remains that recent recruitment is highly uncertain, and estimates for recent years (based on limited data) can change substantially.

Pacific Hake have low average recruitment with occasional large year classes (Figures 32 and 33, Tables 23 and 24). Very large year classes in 1980, 1984, and 1999 supported much of the commercial catch from the 1980s to the mid-2000s. From 2000 to 2007, estimated recruitment was at some of the lowest values in the time series, but this was followed by an above average 2008 year class and a very strong 2010 year class. Above average year classes occurred in 2014 and 2016, which have been sustaining the fishery in recent years (Figure 22).

The current assessment estimates a strong 2014 year class (Figure 34) comprising 50\% of the 2016 catch, $38 \%$ of the 2017 catch, $28 \%$ of the 2018 catch, $33 \%$ of the 2019 catch, $31 \%$ of the 2020 catch, $25 \%$ of the 2021 catch, $15 \%$ of the 2022 catch, and $8 \%$ of the 2023 catch.

The 2016 cohort also appears to be strong, comprising $26 \%$ of the 2018 catch, $21 \%$ of the 2019 catch, $36 \%$ of the 2020 catch, $34 \%$ of the 2021 catch, $24 \%$ of the 2022 catch, and $13 \%$ of the 2023 catch.

The large size of the 2014 and 2016 cohorts is informed by observations from several years of fishery data and the acoustic survey. For all other years from 2011 to 2019, the model currently estimates small year classes (median recruitment below the mean of all median recruitments). As noted above, the 2020 cohort is estimated to be somewhat smaller than in last year's assessment (though last year's estimate was highly uncertain), due to the introduction of new information from the 2023 age- $2+$ biomass index and survey age-composition data (Figure 16). The 2021 cohort strength is only informed by fishery data and the 2023 biomass survey, and is estimated to be potentially large with a median and $95 \%$ credible interval of 10.187 (4.085-29.499) billion fish.

The 2022 cohort was observed by the age- 1 index in 2023, suggesting it is average to below average (Figures 10 and 32), and it will not be observed as part of the age- $2+$ survey index until 2025. There is no information in the data to estimate the sizes of the 2023 and 2024 year classes. Retrospective analyses of year class strength for young fish have shown the estimates of recent recruitment to be unreliable prior to at least model age-3 (observed at age-2 the previous year) without a survey in the most recent year and age-2 (observed at age-1) with a survey.

From Figure 32 it looks as though the 2014 recruitment could be as large as the 2010 recruitment. However, the assessment model estimates a $0 \%$ chance that this could be the case. The overlapping of the credible intervals in Figure 32 is because large MCMC estimates of 2010 recruitment are associated with large estimates of 2014 recruitment (presumably with large estimates of $R_{0}$ ). By scaling all recruitments by the 2010 recruitment, Figure 35 provides an intuitive way to compare recruitment across years (see Edwards et al. 2022 for motivation and methods). It shows that only the 1980 recruitment is probably larger than 2010 (median relative values > 1), and the 1984 recruitment has a small chance of being as large as 2010. Whereas Figure 32 suggests that 1967, 1973, 1977, 1999, 2014, and 2020 could also possibly be larger than in 2010, giving an over-optimistic impression of how often we can expect cohorts the size of the 2010 cohort to occur. The 2021 cohort is still very uncertain but has a small chance of exceeding the 2010 cohort (Figure 35). Participants in the Pacific Hake process have an intuition that the 2010 is a very large recruitment event - Figure 35 shows how it is the largest in the last 30 years, and that such large cohorts are rarer than is inferred from Figure 32.
The estimated recruitments with uncertainty for each year and the overall stock-recruitment relationship are provided in Figure 36. Extremely large variability about the expectation and about the joint uncertainty of individual recruitment and female spawning biomass pairs are evident. High and low recruitments have been produced throughout the range of observed female spawning biomass (Figure 36). The standard deviation of the time series of median recruitment deviation estimates for the years 1970-2022, which are informed by the age compositions and the age- 1 index, is 1.73 .

## Exploitation status

The median estimated relative fishing intensity on the stock is below 1.0 for all years (Figure 37 and Tables 23 and 24). It was closest to 1.0 in 1999 and 2008, but catch in 2008 did not exceed the catch limit that was specified, based on the best available science and harvest control rules in place at the time; however, catch did exceed the catch limit in 1999 (Table 3). Exploitation fraction (catch divided by biomass of fish of age-2 and above) has shown relatively similar patterns (Figure 38 and Tables 23 and 24). Although displaying similar patterns, the exploitation fraction does not necessarily correspond to fishing intensity because fishing intensity more directly accounts for the age-structure of both the population and the catch. Median relative fishing intensity is estimated to have declined from $87.6 \%$ in 2010 to $45.5 \%$ in 2015, and then leveled off around $73-80 \%$ from 2016 to 2019 before declining to $55.1 \%$ in 2023. The median exploitation fraction has increased from a recent low of 0.06 in 2012 to 0.15 in 2021 then declined to 0.07 in 2023. Although there is a considerable amount of imprecision around these recent estimates due
to uncertainty in recruitment and spawning biomass, the $95 \%$ posterior credibility interval of relative fishing intensity was below 100\% from 2012-2016 and again from 2020-2023 (Figure 37).

## Management performance

Over the last decade (2014-2023), the mean coast-wide utilization rate (i.e., proportion of catch target removed) has been $63.5 \%$ and catches have been below coast-wide targets (Table 3). From 2019 to 2023, the mean utilization rates differed between the United States ( $67.4 \%$ ) and Canada ( $48.1 \%$ ), though Canada's rate was higher than the U.S.'s in 2020. In 2015, the utilization rate for the coast-wide fishery was the lowest of the previous decade $(44.1 \%)$ due, in part, to difficulties locating aggregations of fish and possibly economic reasons. Before 2015, the under-utilization in the United States was mostly a result of unrealized catch in the tribal apportionment, while reports from stakeholders in Canada suggested that hake were less aggregated in Canada and availability had declined. In 2016, the utilization rate increased but remained below pre-2015 levels, despite the total 2016 catch being one of the highest of the preceding years. This is in large part due to increasing catch targets as biomass continues to increase. While the total utilization rate between 2017-2021 was relatively steady, it decreased to $59.3 \%$ in 2022 and to $42.2 \%$ in 2023 . This is due to the utilization rate in Canada steadily declining since 2020 to a time-series low of $16.5 \%$ in 2023 , and also a fall in the U.S. utilization rate to $59.7 \%$ in 2023. Country-specific quotas (or catch targets) in 2020 and 2021 were specified unilaterally, due to the lack of an agreement on coast-wide 2020 and 2021 TACs. The usual $73.88 \%$ and $26.12 \%$ allocation of coast-wide TAC, as specified in the Joint U.S.-Canada Agreement for Pacific Hake, was once again implemented in 2022 and 2023. Total landings last exceeded the coast-wide quota in 2002 when utilization was $112 \%$.
As noted above, the median relative fishing intensity was below $100 \%$ (i.e. median fishing intensity below $F_{\mathrm{SPR}=40 \%}$ ) in all years. The median relative spawning biomass was above $40 \%$ (the $B_{40 \%}$ reference point) in all years except 2007-2011 (Table 23 and Figure 31). These are also shown on a phase plot of the joint history of relative spawning biomass and relative fishing intensity (Figure 39). Relative spawning biomass increased from the lows in 2007-2012 with above average recruitment in 2008, 2010, 2014, 2016, and 2020. Correspondingly, median relative fishing intensity has remained below $100 \%$, and total catch has been declining since the time series high in 2017. While there is large uncertainty in the 2023 estimates of relative fishing intensity and relative spawning biomass, the model estimates a $0.2 \%$ joint probability of being both above the $F_{\mathrm{SPR}=40 \%}$ fishing intensity in 2023 and below the $B_{40 \%}$ spawning biomass level at the start of 2024.

### 3.5 Model uncertainty

The base assessment model integrates over the substantial uncertainty associated with several important model parameters including: biomass index and age-1 index catchabilities ( $q_{\mathrm{b}}$ and $q_{1}$, respectively), the magnitude of the stock (via the $R_{0}$ parameter for equilibrium recruitment), productivity of the stock (via the steepness parameter, $h$, of the stock-recruitment relationship), the rate of natural mortality ( $M$ ), annual selectiv-
ity for key ages, recruitment deviations, and survey and fishery data weights (via the Dirichlet-multinomial parameters $\theta_{\text {fish }}$ and $\left.\theta_{\text {surv }}\right)$.
The medians of the key parameters from the posterior distribution are generally similar to those in last year's base model (Table 25). The largest change was a reduction of the 2020 recruitment by more than half, as discussed above, leading to a fall in the estimated median relative spawning biomass at the start of 2023. Medians of the 2014 and 2016 recruitment also both decreased, by about $10 \%$ from those estimated in the 2023 assessment.
The Pacific Hake stock displays a very high degree of recruitment variability, perhaps the largest of any west coast groundfish stock, resulting in large and rapid biomass changes. This volatility, coupled with a dynamic fishery that potentially targets strong cohorts (resulting in time-varying selectivity) will in most circumstances continue to result in highly uncertain estimates of current stock status and even less-certain projections of the stock trajectory. This is particularly true for female spawning biomass estimates in 2024 and throughout the current forecast period, because there is considerable uncertainty associated with the absolute size of the, now mostly mature, 2020 and 2021 year classes that propagates into forecasts. Although the 2023 acoustic survey helped to refine these estimates and reduce uncertainty, further observations of these year classes will improve estimates. The inclusion of the age- 1 index in this assessment will, in some cases, also help to reduce this uncertainty (as it currently does in this case; see Figure 52 discussed later). However, further work is needed to improve upon the characterization of uncertainty in the age- 1 index itself, which is based on a time invariant assumption about index observation error and catchability.

Uncertainty measures in the base model underestimate the total uncertainty in the current stock status and projections, because they do not account for alternative structural models for hake population dynamics and fishery processes (e.g., recruitment, selectivity, or spatial fleet or population structure), the effects of alternative data-weighting choices, survey catchability, and the scientific basis for prior probability distributions. To address structural uncertainties, the JTC investigated a range of alternative models, and we present the key sensitivity analyses along with other informative sensitivity analyses using full MCMC results (Section 3.8).

The JTC continues to be committed to advancing MSE analyses, by coordinating research with the Pacific Hake MSE Working Group and other scientists in the region engaged in similar research. Incorporating feedback from the Working Group and stakeholders will ensure that operating models will be able to provide insight into the important questions defined by interested parties. Specifically, the development of MSE tools will evaluate major sources of uncertainty relating to data, model structure and the harvest policy for this fishery, and will compare potential methods to address them. In the coming years, this will include a host of research evaluations (see Section 3.3 and Section 3.12), including evaluating the utility of incorporating environmentally-driven age-0 recruitment indices into the stock assessment.

### 3.6 Reference points

The term 'reference points' is used throughout this document to describe common conceptual summary metrics. The Agreement specifically identifies $F_{\text {SPR }=40 \%}$ as the default harvest rate and $B_{40 \%}$ as a point where the 40:10 TAC adjustment is triggered (see the Glossary in Appendix C).

We report estimates of the base reference points (e.g., $F_{\mathrm{SPR}=40 \%}, B_{40 \%}, B_{\mathrm{MSY}}$, and MSY) with posterior credibility intervals in Table 26. The median of the female spawning biomass at $F_{\mathrm{SPR}=40 \%}$ (namely the median of $B_{\mathrm{SPR}=40 \%}$ ) and the median yield at $F_{\mathrm{SPR}=40 \%}$ have remained about the same as estimates in the 2023 assessment (Table 25).

As part of the DFO Sustainable Fisheries Framework, Fisheries and Oceans Canada (2009) defined a limit reference point as being a biomass below which serious harm is believed to be occurring to the stock, and an upper stock reference point above which the stock is considered to be healthy. These would equate to the Agreement reference points of $B_{10 \%}$ and $B_{40 \%}$ (the female spawning biomass being $10 \%$ and $40 \%$, respectively, of the unfished equilibrium female spawning biomass). The probabilities of the female spawning biomass at the start of 2024 being above each of these points are $\mathrm{P}\left(B_{2024}>B_{10 \%}\right)=100 \%$ and $\mathrm{P}\left(B_{2024}\right.$ $>B_{40 \%}$ ) $=98.7 \%$ such that the stock is estimated to be in the 'healthy zone' (above the upper stock reference point of $B_{40 \%}$ ). This probability is slightly higher than in last year's assessment, where the equivalent calculation was $\mathrm{P}\left(B_{2023}>B_{40 \%}\right)=98.1 \%$. Note that a probability of ' $100 \%$ ' ( or ' $0 \%$ ') is based on the MCMC results, and is not meant to imply that something definitely occurs (or definitely does not occur).

With respect to DFO's provisional limit reference point of $0.4 B_{\mathrm{MSY}}$ and provisional upper stock reference point of $0.8 B_{\mathrm{MSY}}$, the probabilities are $\mathrm{P}\left(B_{2024}>0.4 B_{\mathrm{MSY}}\right)=100 \%$ and $\mathrm{P}\left(B_{2024}>0.8 B_{\mathrm{MSY}}\right)=100 \%$ such that the stock is estimated to be in the provisional 'healthy zone'. For completeness, we note that $\mathrm{P}\left(B_{2024}>B_{\mathrm{MSY}}\right)=99.9 \%$.
Reference levels of stock status that are used by the U.S. Pacific Fisheries Management Council (PFMC) for Pacific Hake include $B_{40 \%}$ and a Minimum Stock Size Threshold (MSST) of $B_{25 \%}$. For 2024, the estimated posterior median relative spawning biomass is $99 \%$, such that the female spawning biomass is well above $B_{40 \%}$ and $B_{25 \%}$. The probability that female spawning biomass at the beginning of 2024 is above $B_{40 \%}$ is $\mathrm{P}\left(B_{2024}>B_{40 \%}\right)=$ $98.7 \%$ (as noted above), and of being above $B_{25 \%}$ is $\mathrm{P}\left(B_{2024}>B_{25 \%}\right)=99.9 \%$.

### 3.7 Model projections

The catch limit for 2024 based on the default $F_{40 \%}-40: 10$ harvest policy has a median of $747,588 \mathrm{t}$ and a wide range of uncertainty (Figure 40), with the $95 \%$ credibility interval being 298,355-2,124,832 t.

Decision tables give projected population status (relative spawning biomass and relative fishing intensity) under different catch alternatives for the base model (Tables 27 and 28). The tables are organized such that the projected outcome for each potential catch level and year (each row) can be evaluated across the quantiles (columns) of the posterior
distribution. Table 27 shows projected relative spawning biomass outcomes, and Table 28 shows projected fishing intensity outcomes relative to $F_{\mathrm{SPR}=40 \%}$.
Population dynamics and governing parameters assumed during the forecast period include random recruitment; selectivity, weight-at-age and maturity (and thus fecundity) averaged over the five most recent years (2019-2023); and all estimated parameters constant (at their estimates for each particular MCMC sample).

Relative fishing intensity exceeding 1 (or $100 \%$ when shown as a percentage) indicates fishing in excess of the $F_{\mathrm{SPR}=40 \%}$ default harvest rate limit. A slight exceedance can happen for the median relative fishing intensity in 2024, 2025 and 2026 because the $F_{\text {SPR }=40 \%}$ default harvest-rate catch limit is calculated using baseline selectivity-at-age (1966-1990; prior to time-varying deviations), whereas the forecasted catches under the default harvestrate are removed using selectivity averaged over the last five years. Recent changes in selectivity could be reflected in the projection of slight over- or under-fishing relative to the desired $F_{\text {SPR }=40 \%}$ rate.

Key management metrics are presented for 2025, 2026 and 2027 projections (Tables 29-31 and Figures 44-46). These metrics summarize the probability of various outcomes from the base model given each potential management action. Although not linear, probabilities can be interpolated from this table for intermediate catch values in 2024 (Table 29 and Figure 44). However, interpolation may not be applicable for all catches in 2025 and 2026 because they are conditional on catch levels from the previous year or years. This explains why a few probabilities decline (rather than rise) with increased 2025 and 2026 catch levels in Tables 30 and 31 and Figures 45 and 46.

Figure 43 shows the projected relative spawning biomass trajectory through 2027 for several of these management actions. With zero catch for the next three years, the biomass has a 3\% probability of decreasing from 2024 to 2025 (Table 29 and Figure 44), a 59\% probability of decreasing from 2025 to 2026 (Table 30 and Figure 45), and a $66 \%$ probability of decreasing from 2026 to 2027 (Table 31 and Figure 46).

The probability of the female spawning biomass decreasing from 2024 to 2025 is greater or equal to $22 \%$ for all catch levels examined other than zero (Table 29 and Figure 44). The probability is $36 \%$ for the 2024 catch level similar to that for 2023 (catch alternative e). For all explored catches, the maximum probability of the female spawning biomass dropping below $B_{10 \%}$ at the start of 2025 is $0.0 \%$, at the start of 2026 is $0.1 \%$, and at the start of 2027 is $2.6 \%$ (Tables 29-31 and Figures 44-46). The similar maximum probability of dropping below $B_{40 \%}$ at the start of 2025 is $6.7 \%$, at the start of 2026 is $19.9 \%$, and at the start of 2027 is $34.8 \%$.

It should be noted that forecasted biomass is not only influenced by catch levels. As the above average 2014 and 2016 cohorts continue to age, total biomass of these cohorts even without fishing mortality will continue to decrease (Tables 18 and 22) as losses from mortality outweigh increases from growth. The above-average 2020 cohort entered this growth-mortality transition period around 2023 (Tables 18 and 22). During 2024, the age-3 2021 cohort will likely begin the growth-mortality transition where a net increase in total biomass is less likely (note that fecundity will increase which will influence the exact
change in female spawning biomass, Figure 11). The estimated above-average (yet still highly uncertain) 2020 and 2021 cohorts will continue to play a large role in determining female spawning biomass during the forecast years presented here. The below-average 2015, 2018, and 2019 cohorts will contribute much less to forecasted spawning biomass than the larger cohorts.

The age composition (in numbers) of the catch in 2024 is projected to be (using MCMC medians) $38 \%$ age- 3 fish from the 2021 cohort, $23 \%$ age- 4 fish from the 2020 cohort, $9 \%$ age- 8 fish from the 2016 cohort, $7 \%$ age- 2 fish from the 2022 cohort, and $6 \%$ age- 10 fish from the large 2014 cohort (Figure 47). However, those estimates are highly uncertain with the $95 \%$ credibility interval for the age- 3 fraction spanning $21 \%-57 \%$.

Due to the higher average weight of older fish compared to younger fish, the median expected proportion of the 2024 catch by weight is $32 \%$ for the age 32021 cohort (compared to $38 \%$ by numbers) and $23 \%$ for the age- 42020 cohort (compared to $23 \%$ by numbers; Figure 47).

With respect to the DFO reference points, with the largest 2024 catch of $875,262 \mathrm{t}$ given in Table 29, at the start of 2025 the stock is expected to be above the critical zone with a probability of $\mathrm{P}\left(B_{2025}>B_{10 \%}\right)=100 \%$ and in the healthy zone with a probability of $\mathrm{P}\left(B_{2025}>B_{40 \%}\right)=93 \%$. With respect to the DFO provisional reference points (based on $\left.B_{\mathrm{MSY}}\right)$, the stock is expected to be above the provisional critical zone with a probability of $\mathrm{P}\left(B_{2025}>0.4 B_{\mathrm{MSY}}\right)=100 \%$, in the healthy zone with a probability of $\mathrm{P}\left(B_{2025}>0.8 B_{\mathrm{MSY}}\right)$ $=100 \%$, and above $B_{\mathrm{MSY}}$ with a probability of $\mathrm{P}\left(B_{2025}>B_{\mathrm{MSY}}\right)=99 \%$ for this catch.
With respect to PFMC stock size reference points, a level of 2024 catch consistent with the Agreement default harvest control rule (747,588 t) has a 5\% estimated probability of resulting in the biomass going below $B_{40 \%}$ at the start of 2025 (and $1 \%$ probability of going below $B_{25 \%}$; Table 29). If catches in 2024 and 2025 are the same as in 2023 (264,000 t, catch scenario e) then the probability of the biomass going below $B_{40 \%}$ is $1 \%$ for the start of 2025 and $3 \%$ for the start of 2026.

### 3.8 Sensitivity analyses

Sensitivity analyses were conducted to investigate influence of data inputs and structural uncertainty of the base model by investigating how changes to the model affected the estimated values and derived quantities. All sensitivity analyses compared MCMC posteriors with the same number of posterior samples as the base model. Several key underlying structural model assumptions were identified that have persisted across many previous hake assessments, and thus warrant revisiting annually as a set of reference sensitivity examinations to new base models. Many additional sensitivity runs were conducted while developing and testing the 2024 base model. Here we focus on the main sensitivities, relative to the base model, which are as follows:

1. Consideration of higher standard deviations on the prior distribution for natural mortality;
2. Consideration of an alternative prior distribution (mean and standard deviation) for natural mortality based on the Hamel (2015) and Hamel and Cope (2022) life history meta-analytic method;
3. Consideration of an alternative prior distribution and a fixed value for steepness, to change the resiliency of the stock;
4. Consideration of higher and lower variation about the stock-recruitment relationship $\left(\sigma_{r}\right)$;
5. Removal of the age- 1 index as a data source;
6. Downweighting the fishery age-composition data; and
7. Consideration of alternative standard deviations for time-varying selectivity.

None of the sensitivities resulted in a substantial departure from the main population dynamics of the base model (Tables 32-34 and Figures 48-58). All sensitivity models showed large estimated increases in female spawning biomass in the early- to mid-2010s that continues to be driven by the 2010, 2014, and 2016 cohorts, followed by a period of general decline (2018-2023) before increasing again due to the above average 2020 and 2021 cohorts. All sensitivity models indicate that 2024 relative spawning biomass is above $B_{40 \%}$. The overall scale of the population was impacted by various alternative assumptions, and the highly uncertain size of the recent cohorts were more variable across sensitivity analyses than earlier cohorts that have been observed for more years.

The standard deviation of the prior distribution on natural mortality was increased from the base model value of 0.1 to $0.2,0.3$, and 0.31 . Note that the median of the prior was also changed for the latter sensitivity. Each of these sensitivities led to an increase in estimates of natural mortality relative to the base model. The medians of the MCMC posteriors for natural mortality increased from 0.235 to $0.290,0.312$, and 0.315 , respectively. The $95 \%$ credibility intervals also increased, with the largest differences in the upper rather than the lower credible interval. Credible intervals were $0.194-0.280$ for the base model, $0.227-0.343$ for the sensitivity run with the prior standard deviation set to $0.2,0.242-0.360$ for the sensitivity run with the prior standard deviation set to 0.3 , and $0.248-0.361$ for the sensitivity run with the Hamel and Cope (2022) prior (Table 32). In addition to increased estimates of natural mortality, results from these sensitivity models also showed an increase in the overall scale of the population, the estimated stock status relative to $B_{0}$ prior to 1990, the uncertainty in female spawning biomass on both absolute and relative scales, roughly halved estimated relative fishing intensity in 2023, and more than doubled equilibrium yield at $B_{\mathrm{SPR}=40 \%}$ (Table 32 and Figures 48 and 49).
The mean of the prior distribution on steepness was decreased from 0.777 (base) to 0.5 and, separately, steepness was fixed at 1.0. The decrease in the mean of the prior resulted in a decrease in the MCMC estimate of steepness from a median of 0.812 with a $95 \%$ credible interval of $0.582-0.958$ to a median of 0.541 with a $95 \%$ credible interval of $0.345-0.756$ (Table 32). However, neither steepness sensitivity analysis had a large impact on the overall model results (Figures 48 and 49), because Pacific Hake female spawning biomass
has remained above levels where changes in steepness would appreciably influence the stock-recruitment relationship (Figure 36).

Input values of $\sigma_{r}$ were changed from 1.40 (base) to alternative high (1.60) and low (1.00) states. Both sensitivities were similar to the base model in that the calculated standard deviation of recruitment deviations (from the main period) was higher than the input $\sigma_{r}$, i.e., 1.58 and 1.89 when $\sigma_{r}$ was 1.00 and 1.60 , respectively. The calculated standard deviation of recruitment deviations from the base model was intermediate at 1.70. These calculated standard deviations should match the input if the vectors of deviations were from the 'population' of values rather than just a sample. However, this systematic bias to be larger than the input value indicates that the standard deviation of recruitment deviations is accounting for more variability than just variability in recruitment. The high $\sigma_{r}$ model led to a larger difference between the female spawning biomass at unfished equilibrium and the female spawning biomass at the initial year of the model than the low $\sigma_{r}$ model (Figure 48). Similar to previous assessments, estimates of unfished equilibrium recruitment and relative spawning biomass are sensitive to $\sigma_{r}$, whereas absolute estimates of female spawning biomass are relatively insensitive. The method Methot and Taylor (2011) proposed to tune $\sigma_{r}$ was developed in the context of maximum likelihood estimation and not Bayesian inference, where the latter potentially allows for estimating $\sigma_{r}$ using random effects, and thus, this proposed method is not used here to tune the fixed input value.

The sensitivity of the base model to the removal of the relative age- 1 index provides a method to evaluate how the information about juvenile fish is propagated through the model. Estimates of female spawning biomass throughout most of the time series are similar between models with and without the relative age- 1 index but diverge near the end of the time series (Table 32, Figures 50 and 51). The 2024 estimates of relative spawning biomass are $98.7 \%$ for the base model ( $95 \%$ credible interval of $45.0-229.8 \%$ ) and $78.4 \%$ for the model where the relative age- 1 index is removed ( $95 \%$ credible interval of $34.2-184.1 \%$ ). This difference is due to the relative age- 1 index providing additional information on recruitment for cohorts associated with recent age-1 indices (i.e., 2020 and 2022 cohorts detected in the 2021 and 2023 age- 1 indices). In particular, the base model with the relative age-1 index indicates slightly larger 2020 and 2021 year classes than the model removing the age-1 index (Figure 52). Similarly, recruitment in 2022 is estimated to be slightly above average when the model is fit to the relative age- 1 index compared to slightly below average without the index. Removing the relative age- 1 index led to minor changes in fit to the age- $2+$ survey biomass index, with 2021 showing a slight improvement and 2023 a deterioration compared to the base model (Figure 53).

The base model includes a Dirichlet-multinomial likelihood component that includes two estimated parameters to automatically weight each of the fishery and survey age compositions. The base model was compared to a model that downweighted the fishery age compositions relative to the survey age compositions. This downweighting was based on the McAllister-Ianelli method, which requires manual iterative adjustments to the input sample sizes using a derived multiplier. The McAllister-Ianelli method, which was used in assessments prior to 2018 (Table 16), attempts to make the arithmetic mean of the input sample size approximately equal to the harmonic mean of the effective sample size. Here,
this was accomplished with weighting factors of 0.14 and 0.46 (ratio of 0.30 ) for fishery and survey age compositions, respectively. These weighting factors are not specific to this year's base model, rather they are values calculated from previous maximum likelihood estimates. The median estimate from the Dirichlet-multinomial method used in the base model was 0.340 and 0.941 (ratio of 0.36 ). Downweighting fishery composition data led to minor changes in relative spawning biomass, recruitment estimates, and increased uncertainty in estimates of early recruitments compared to the base model (Figures 51 and 52). The largest changes in the time series occurred prior to the availability of survey data.

The degree of flexibility of annual variation in the fishery selectivity was tested using three alternative values of standard deviations $(\Phi)$ (Figures 54-58). The consideration of alternative $\Phi$ values is discussed earlier in Section 2.5.3. Changing $\Phi$, which controls the flexibility in time-varying selectivity, from the base model value of $\Phi=1.40$ to 0.21 , 0.70 , and 2.10 did not appreciably influence the estimates, or precision, associated with recruitment in 2014 or 2016 but it did impact more recent recruitments (Figure 56). In particular, recruitment estimates for 2020 and 2021 are linked to the choice of $\Phi$, where the smallest investigated value of $\Phi(0.21)$ led to the highest estimates of the 2020 and 2021 recruitment deviations of the investigated models (Figure 57). The high estimates of recruitment led to a large increase in female spawning biomass in recent years compared to the base model (Figure 54). When $\Phi=0.21$, the fit to the most recent age- $2+$ survey biomass index was the worst of the three investigated models (Figure 58).

### 3.9 Retrospective analyses

Retrospective analyses were performed by iteratively removing the terminal years' data (going back 10 years) and estimating the posterior distribution of parameters under the assumptions of the base model. This year's base model shows similar retrospective results to last year's (Figure 59 and Edwards et al. 2022) for the older cohorts. Uncertainties are shown for select cohorts in Figures 59 and 60. In previous years, thes figures showed only the median lines. The uncertainty is represented as credible intervals from $2.5 \%$ to $97.5 \%$ as shaded areas surrounding the median lines. For cohorts that have positive recruitment deviations, the uncertainty is a narrower band around the median due to a higher sampling rate over the years than the cohorts with negative recruitments.

The 2020 cohort has been estimated lower this year than in last year's assessment, which is also evident in Figure 60 when excluding the 2023 data - the uncertainty at age- 3 gets reduced when including the 2023 data, shown by the age- 4 intervals being tighter and also lower. The latest data no longer suggest that 2020 is a huge cohort. The 2021 cohort at age- 3 has a similar median to the 2020 cohort at age-3, but with less uncertainty (narrower credibility interval in Figure 60) because it has been seen in the age-2+ biomass survey (whereas the 2020 cohort was not seen in that survey until it was another year older). Although the 2021 displays unusual behaviour in that the median is below 0 at age- 2 and then above 0 at age-3, its uncertainty at age-2 was very large (Figure 60).

Some cohort recruitments are over or under-estimated at age-2. Over-estimation can be seen most clearly with the 2014, 2015, 2017, 2018, and 2020 cohorts (Figures 59 and 61). The

2014 cohort reached high deviation after two years, then even higher after three years only to drop back down to a lower value and then stabilize at around age- 4 with the addition of more data. A similar pattern can be seen with the smaller 2017 and 2018 cohorts. Even with the addition of new data, the size of the very small 2015 cohort has not fully stabilized. Under-estimation is slight, but apparent, for the 2016 cohort as recruitment estimates have risen by a small amount since the estimate at age-3. The under-estimation of the 2021 cohort stands out as it was estimated as being slightly smaller than average at age- 2 in last year's assessment and then estimated to be a very large cohort at age-3 this year, though this is based on medians and the age- 2 estimate was highly uncertain, as mentioned above.

Cohort strength is further informed once at least one year of age- $2+$ survey biomass index age-composition data are available for a cohort, which for even-numbered recruitment years typically does not occur until the cohort reaches age-3, due to the acoustic survey occurring in odd years; though the age- 1 index does provide some information.

The stability of the recruitment estimates seen in this plot is also evident in the absolute estimates of uncertainty for each cohort. Uncertainty of the 2016-2021 cohorts has been substantially reduced compared to removing five years of data (Figure 62, bottom figure). The uncertainty of the 2020 cohort was substantially increased with the removal of only 1 year of data. This increase was exacerbated by the removal of the 2023 survey index as well as the fishery catch, as all data sources are removed for each year of the retrospectives. Medians of various quantities of interest are given in Table 35.

Overall, there is little retrospective change to the relative spawning biomass trajectory up to the mid-2010s, and most retrospective change occurs in the final 5 years of the retrospective model (upper panel of Figure 62). In this assessment, there is very little retrospective bias, with only slight year-specific positive and negative bias in female spawning biomass, some minor adjustments to recruitment deviates, and a slight trend in $B_{0}$ as the retrospective year increases. All of these retrospective differences are well within the range of estimation uncertainty across all retrospective years. There is no indication from retrospective evaluations that the base model is displaying a systematic bias.

### 3.10 Comparison with past assessments

A comparison of the base models, approved for management, used in each year since 1991 indicates that the variability between model results, especially early on in the estimated time series, is larger than the estimated uncertainty reported from the current base model (Figure 63). There have been substantial differences in the structural assumptions of the models and, thus, results submitted each year. Prior to 2004, catchability was fixed at 1.0. This assumption was investigated between 2004 and 2007, leading to variability in model results because of the use of several different, but fixed, values of catchability. Since 2008, catchability has been freely estimated by the model ( $q_{b}=0.84$ ). The fixing of survey catchability had the effect of driving the estimate of initial biomass upward, which in turn scaled the entire biomass trajectory up, leading to higher estimates of relative spawning biomass than in more recent assessments. The median estimates of female spawning biomass for 2016 and 2017 have remained similar to the previous assessment, being somewhat lower than in the 2016 and 2017 assessments. In addition to more information
about the 2014 and 2016 cohorts, the 2018 assessment model also included a change in the data weighting method, an update to maturity and fecundity, and a change to selectivity parameterization (Table 16). The uncertainty interval associated with the 2024 assessment brackets the majority of the historical estimates.

The level of uncertainty associated with each assessment's estimate of that year's current female spawning biomass (i.e., that used to convey current stock status and inform management advice) changes from assessment to assessment given updates in data and Pacific Hake population structure and dynamics. Uncertainty around the absolute amount of 2024 female spawning biomass is similar to the final-year estimates from previous assessments, with both absolute interquartile range and the relative amount of dispersion (or variability relative to the stock size; similar to a coefficient of variation) consistent with previous assessments (Figure 64).

### 3.11 Performance of past projections

Without rigorous simulation experiments it can be difficult to operationally assess the accuracy of projections in stock assessments because the truth is never known with $100 \%$ certainty. For Pacific Hake, hindsight comparisons have been conducted since 2021 (Johnson et al. 2021) to evaluate performance of projections provided in decision tables (such as Tables 29 and 30) of past assessments relative to updated assessments. Overall, results indicate that assessment model projections give a relatively good idea of general projected trends and status.

As an example, the 2019 assessment (Berger et al. 2019) gave the estimated probability of the female spawning biomass declining in the subsequent year, i.e., $\mathrm{P}\left(B_{2020}<B_{2019}\right)$, for several possible catches in 2019, such as $0 t, 180,000 t, 350,000 t, 410,000 t$ etc. Now that we 'know' the catch in 2019 was $412,015 \mathrm{t}$, we can select the $410,000 \mathrm{t}$ row (close enough to $412,015 \mathrm{t})$ in the table from the 2019 assessment to give that assessment's $\mathrm{P}\left(B_{2020}<B_{2019}\right)=$ $61 \%$; Figure 65. We can also calculate this probability from the current assessment model, which implicitly includes the $412,015 \mathrm{t}$ catch from 2019, giving $\mathrm{P}\left(B_{2020}<B_{2019}\right)=83 \%$; Figure 65. We extracted similar probabilities from past assessment documents going back to 2012 and calculate analogous probabilities, $\mathrm{P}\left(B_{\mathrm{t}+1}<B_{\mathrm{t}}\right)$, from the current base model [Figure 65; see Edwards et al. (2022) for full methods].

Each assessment correctly predicted whether the stock would most likely increase or decrease the following year, except for 2017 and 2023; Figure 65. Estimates from previous assessments are almost always closer to $50 \%$ than those from the current base model (Figure 65), because the current assessment model has more information and thus provides a more definitive probability (closer to $0 \%$ or to $100 \%$ ) than year $t$ 's assessment model. It is desirable that the probabilities from the assessment documents are not too definitive (too close to $0 \%$ or to $100 \%$ ), because they are admitting a wide range of uncertainty given unknown recent recruitments.

The 2017 and 2023 assessments 'incorrectly' projected that the stock would likely decline the following year (given the catch that subsequently occurred), because the current assessment model estimates a likely increase (Figure 65). For the 2017 (Berger et al. 2017)
assessment the biomass trend was projected to be relatively flat the following year, so even slight changes in biomass could influence the binomial outcome of an 'increase' or 'decrease' in biomass, despite the overall change in biomass not being very substantial. The 2023 assessment (Berger et al. 2023) had minimal information on the 2021 cohort and predicted the biomass would probably decline in 2024 with any non-zero 2023 catch. However, the current assessment estimates that the 2021 cohort was potentially large, which further highlights how impactful a realized large deviation from average recruitment (rather than assuming average recruitment) can be on forecasted outcomes. Similarly, the 2012 assessment had no information on the very large 2010 recruitment, and so also over-estimated the probability of decline the following year (Figure 65). A range of catch alternatives are shown for the current assessment because realized 2024 catches are not yet known (Figure 65), and give a mostly greater that $50 \%$ chance that the stock will decline from 2024 to 2025.

A similar approach was used to calculate the probability of the biomass falling below $B_{40 \%}$ in the subsequent year, i.e., $\mathrm{P}\left(B_{\mathrm{t}+1}<B_{40 \%}\right)$; Figure 66. The 2012 assessment was the only one that gave a $>50 \%$ chance of the biomass falling below $B_{40 \%}$ in the subsequent year, but later data determined that the 2010 year class was substantial and so in hindsight the probability of going below $B_{40 \%}$ was $0 \%$ (based on the current assessment). From the 2018 assessment onwards, the estimated $\mathrm{P}\left(B_{\mathrm{t}+1}<B_{40 \%}\right)$ probabilities rose, until falling due to the incoming above-average 2020 cohort and lower catches (Figure 66). The same probabilities calculated from the current base model similarly rose, but all remained lower than the previous assessments' calculations, similar to the analogous figure in the 2023 assessment (Berger et al. 2023).

### 3.12 Research and data needs

There are many research projects that could improve the stock assessment for Pacific Hake and lead to improved biological understanding and decision-making. The most important are as follows:

1. Continue to conduct research to evaluate ways to improve recent, current, and future estimates of recruitment for use in stock assessment. This could include the development of time series of recruitment indices, time series of informative environmental or ecosystem variables, and models that have predictive skill (e.g., Vestfals et al. 2023). Explorations should also consider options for incorporating information on recruitment into the assessment model and the management framework for Pacific Hake. For example, time series could be included in the stock assessment as a standalone data source (similar to the acoustic indices) or improvements could be made to the modeling framework such that these environmental time series could impact the stock-recruitment relationship directly. Results from such work should be connected to or in cooperation with ongoing research related to recruitment variability as discussed in Section 3.3. Related, there is a need to streamline and broaden the availability of products from oceanographic models (e.g., Regional Ocean Modeling System) so they are available across international boundaries and updated on a recurring basis, thereby allowing for their use as informative links in operational stock assessments. A successful example of this has been the annual production of Pacific Hake distribution forecasts that depend on 6-9 month forecasts of subsurface (i.e., 100 m depth) temperature from J-SCOPE. Furthermore, the existing management strategy evaluation framework should be used, or further developed, to examine how information on recruitment can inform robust management decisions.
2. Conduct research on estimates of uncertainty for the relative age- 1 index and the age- $2+$ index and investigate alternative ways to utilize survey age-composition information in the assessment model. Bootstrapping of the acoustic survey time series, or related methods, could help incorporate uncertainty related to the target-strength relationship, subjective scoring of echograms, thresholding methods, and methods used to estimate the species mixes for interpreting the acoustic backscatter into the variance calculations. Research should be communicated with those involved in developing the U.S. West Coast Integrated Survey Initiative. The management strategy evaluation framework should be used, or further developed, to examine how changes in survey methods can be used to inform robust management decisions.
3. Work with regional partners to develop an annual workflow that provides key metrics, indicators, or other summaries of general ecosystem conditions relevant to the coast-wide population of Pacific Hake. In particular, include indicators that are potentially associated with Pacific Hake biology and ecology (e.g., recruitment, distribution, predation, prey, and communities). Such information can broaden the context within which a single species stock assessment is interpreted, be used to support model development, refine uncertain assessment conclusions (e.g., productivity), and provide other non-assessment indicators of the system's state to management.
4. Use, build, and expand upon the existing management strategy evaluation framework to evaluate major sources of uncertainty relating to data, model structure, and the harvest policy for this fishery (as needed) and compare potential methods to address them. In particular, utilize and adapt the management strategy evaluation framework to address new and ongoing stock assessment research and data needs through the Pacific Hake Management Strategy Evaluation Working Group, including relevant requests by the Scientific Review Group (see Section 3.3). For example, research investigating links between Pacific Hake biomass, spatial distribution, growth, recruitment, and natural mortality, and how these biological processes vary with ocean conditions and ecosystem variables such as temperature, transport, and prey availability could inform models used in the MSE. Ongoing investigations have the potential to improve the scenarios considered in future work on the MSE framework and the basic understanding of drivers of Pacific Hake population dynamics and availability to fisheries and surveys.
5. Complete the ongoing inter-laboratory otolith exchange and use the results to update estimates of ageing error used in the stock assessment. This would include updated information about ageing imprecision, the effects of large cohorts, and comparisons between ageing methods such as break and burn, surface reads, and Fourier-Transform Near Infrared Spectroscopy. The last inter-laboratory comparison was done in 2010 ('CARE' exchanges). Related, streamlining procedures that ease the exchange of biological materials (e.g., otoliths) across international borders would increase the efficiency at which research products can be produced.
6. Improve stock assessment forecasts through research that identifies linkages between Pacific Hake biology and ecosystem, oceanographic, or climate variables across the population domain. In particular, explore possible relationships with recruitment, growth, fecundity (including weight-at-age and maturity), and population density to improve biomass forecasting capabilities for Pacific Hake.
7. Explore the operational use of environmental DNA data for characterizing aspects of Pacific Hake population dynamics, such as changes in species distribution or density and the incorporation of these data into the assessment. Recent research demonstrated that environmental DNA provides similar information as the acoustic survey at scales relevant to management, i.e., coast-wide and not just sample-tosample comparisons (Shelton et al. 2022), but longer time series are needed before the data can be used to inform trends in abundance. Environmental DNA is now available for 2019, 2021, and 2023 (three years total). Continuing to extend the time series would allow for its incorporation in future stock assessments as a relative index of abundance.
8. Explore alternative approaches and related assumptions for parameterizing timevarying fishery selectivity in the assessment. Simulations that evaluate methods for including multiple variance structures, including interactions, tradeoffs, and related assumptions, across multiple processes (e.g., selectivity, recruitment, data weighting) in integrated stock assessment models would be particularly beneficial.
9. Explore the potential to use acoustic data collected from commercial fishing vessels to study Pacific Hake distributions, schooling patterns, and other questions of interest. This could be similar to the 'acoustic vessels of opportunity' program on fishing vessels targeting Pollock in Alaska (Stienessen et al. 2019).

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## 6 TABLES

Table 1. Annual catches of Pacific Hake (t) in U.S. waters by fleet, 1966-2023. Tribal catches are included in the fleet totals. Research catch includes landed catch associated with research-related activities. Catch associated with surveys and discarded bycatch in fisheries not targeting hake is not currently included in the table or model.

| Year | Foreign | $\begin{aligned} & \text { Joint- } \\ & \text { venture } \end{aligned}$ | Mothership | Catcherprocessor | Shorebased | Research | Total |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1966 | 137,000 | 0 | 0 | 0 | 0 | 0 | 137,000 |
| 1967 | 168,700 | 0 | 0 | 0 | 8,960 | 0 | 177,660 |
| 1968 | 60,660 | 0 | 0 | 0 | 160 | 0 | 60,820 |
| 1969 | 86,190 | 0 | 0 | 0 | 90 | 0 | 86,280 |
| 1970 | 159,510 | 0 | 0 | 0 | 70 | 0 | 159,580 |
| 1971 | 126,490 | 0 | 0 | 0 | 1,430 | 0 | 127,920 |
| 1972 | 74,090 | 0 | 0 | 0 | 40 | 0 | 74,130 |
| 1973 | 147,440 | 0 | 0 | 0 | 70 | 0 | 147,510 |
| 1974 | 194,110 | 0 | 0 | 0 | 0 | 0 | 194,110 |
| 1975 | 205,650 | 0 | 0 | 0 | 0 | 0 | 205,650 |
| 1976 | 231,330 | 0 | 0 | 0 | 220 | 0 | 231,550 |
| 1977 | 127,010 | 0 | 0 | 0 | 490 | 0 | 127,500 |
| 1978 | 96,827 | 860 | 0 | 0 | 690 | 0 | 98,377 |
| 1979 | 114,910 | 8,830 | 0 | 0 | 940 | 0 | 124,680 |
| 1980 | 44,023 | 27,537 | 0 | 0 | 790 | 0 | 72,350 |
| 1981 | 70,365 | 43,557 | 0 | 0 | 838 | 0 | 114,760 |
| 1982 | 7,089 | 67,465 | 0 | 0 | 1,023 | 0 | 75,577 |
| 1983 | 0 | 72,100 | 0 | 0 | 1,051 | 0 | 73,151 |
| 1984 | 14,772 | 78,889 | 0 | 0 | 2,721 | 0 | 96,382 |
| 1985 | 49,853 | 31,692 | 0 | 0 | 3,894 | 0 | 85,439 |
| 1986 | 69,861 | 81,640 | 0 | 0 | 3,432 | 0 | 154,932 |
| 1987 | 49,656 | 105,997 | 0 | 0 | 4,795 | 0 | 160,448 |
| 1988 | 18,041 | 135,781 | 0 | 0 | 6,867 | 0 | 160,690 |
| 1989 | 0 | 195,636 | 0 | 0 | 7,414 | 0 | 203,049 |
| 1990 | 0 | 170,972 | 0 | 4,537 | 9,632 | 0 | 185,142 |
| 1991 | 0 | 0 | 86,408 | 119,411 | 23,970 | 0 | 229,789 |
| 1992 | 0 | 0 | 36,721 | 117,981 | 56,127 | 0 | 210,829 |
| 1993 | 0 | 0 | 14,558 | 83,466 | 42,108 | 0 | 140,132 |
| 1994 | 0 | 0 | 93,610 | 86,251 | 73,616 | 0 | 253,477 |
| 1995 | 0 | 0 | 40,805 | 61,357 | 74,962 | 0 | 177,124 |
| 1996 | 0 | 0 | 62,098 | 65,933 | 85,128 | 0 | 213,159 |
| 1997 | 0 | 0 | 75,128 | 70,832 | 87,416 | 0 | 233,376 |
| 1998 | 0 | 0 | 74,686 | 70,377 | 87,856 | 0 | 232,920 |
| 1999 | 0 | 0 | 73,440 | 67,655 | 83,470 | 0 | 224,565 |
| 2000 | 0 | 0 | 53,110 | 67,805 | 85,854 | 0 | 206,770 |
| 2001 | 0 | 0 | 41,901 | 58,628 | 73,412 | 0 | 173,940 |
| 2002 | 0 | 0 | 48,404 | 36,342 | 45,708 | 0 | 130,453 |
| 2003 | 0 | 0 | 45,396 | 41,214 | 55,335 | 0 | 141,945 |
| 2004 | 0 | 0 | 47,561 | 73,176 | 96,503 | 0 | 217,240 |
| 2005 | 0 | 0 | 72,178 | 78,890 | 109,052 | 0 | 260,120 |
| 2006 | 0 | 0 | 60,926 | 78,864 | 127,165 | 0 | 266,955 |
| 2007 | 0 | 0 | 52,977 | 73,263 | 91,441 | 0 | 217,682 |
| 2008 | 0 | 0 | 72,440 | 108,195 | 67,760 | 0 | 248,395 |
| 2009 | 0 | 0 | 37,550 | 34,552 | 49,222 | 0 | 121,324 |
| 2010 | 0 | 0 | 52,022 | 54,284 | 64,653 | 0 | 170,960 |
| 2011 | 0 | 0 | 56,394 | 71,678 | 102,146 | 1,042 | 231,261 |
| 2012 | 0 | 0 | 38,512 | 55,264 | 65,919 | 448 | 160,144 |
| 2013 | 0 | 0 | 52,470 | 77,950 | 102,141 | 1,018 | 233,578 |
| 2014 | 0 | 0 | 62,102 | 103,203 | 98,640 | 197 | 264,141 |
| 2015 | 0 | 0 | 27,665 | 68,484 | 58,011 | 0 | 154,160 |
| 2016 | 0 | 0 | 65,036 | 108,786 | 87,760 | 745 | 262,327 |
| 2017 | 0 | 0 | 66,428 | 136,960 | 150,741 | 0 | 354,129 |
| 2018 | 0 | 0 | 67,121 | 116,073 | 135,112 | 0 | 318,306 |
| 2019 | 0 | 0 | 52,646 | 116,146 | 148,210 | 0 | 317,002 |

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| :---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| Year | Foreign | Joint- <br> venture | Mother- <br> ship | Catcher- <br> processor | Shore- <br> based | Research | Total |
| 2020 | 0 | 0 | 37,978 | 111,147 | 138,688 | 95 | 287,908 |
| 2021 | 0 | 0 | 35,208 | 104,030 | 129,319 | 917 | 269,473 |
| 2022 | 0 | 0 | 59,516 | 126,247 | 105,939 | 0 | 291,702 |
| 2023 | 0 | 0 | 32,911 | 107,117 | 100,396 | 0 | 240,424 |

Table 2. Annual catches of Pacific Hake ( t ) in Canadian waters by fleet, 1966-2023.

| Year | Foreign | Jointventure | Shoreside | Freezertrawler | Total |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1966 | 700 | 0 | 0 | 0 | 700 |
| 1967 | 36,710 | 0 | 0 | 0 | 36,710 |
| 1968 | 61,360 | 0 | 0 | 0 | 61,360 |
| 1969 | 93,850 |  | 0 | 0 | 93,850 |
| 1970 | 75,010 | 0 | 0 | 0 | 75,010 |
| 1971 | 26,700 | 0 | 0 | 0 | 26,700 |
| 1972 | 43,410 | 0 | 0 | 0 | 43,410 |
| 1973 | 15,130 | 0 | 0 | 0 | 15,130 |
| 1974 | 17,150 | 0 | 0 | 0 | 17,150 |
| 1975 | 15,700 | 0 | 0 | 0 | 15,700 |
| 1976 | 5,970 | 0 | 0 | 0 | 5,970 |
| 1977 | 5,190 | 0 | 0 | 0 | 5,190 |
| 1978 | 3,450 | 1,810 | 0 | 0 | 5,260 |
| 1979 | 7,900 | 4,230 | 300 | 0 | 12,430 |
| 1980 | 5,270 | 12,210 | 100 | 0 | 17,580 |
| 1981 | 3,920 | 17,160 | 3,280 | 0 | 24,360 |
| 1982 | 12,480 | 19,680 |  | 0 | 32,160 |
| 1983 | 13,120 | 27,660 |  | 0 | 40,780 |
| 1984 | 13,200 | 28,910 | 0 | 0 | 42,110 |
| 1985 | 10,530 | 13,240 | 1,190 | 0 | 24,960 |
| 1986 | 23,740 | 30,140 | 1,770 | 0 | 55,650 |
| 1987 | 21,450 | 48,080 | 4,170 | 0 | 73,700 |
| 1988 | 38,080 | 49,240 | 830 | 0 | 88,150 |
| 1989 | 29,750 | 62,718 | 2,562 | 0 | 95,029 |
| 1990 | 3,810 | 68,314 | 4,021 | 0 | 76,144 |
| 1991 | 5,610 | 68,133 | 16,174 | 0 | 89,917 |
| 1992 | 0 | 68,779 | 20,043 | 0 | 88,822 |
| 1993 | 0 | 46,422 | 12,352 | 0 | 58,773 |
| 1994 | 0 | 85,154 | 23,776 | 0 | 108,930 |
| 1995 | 0 | 26,191 | 46,181 | 0 | 72,372 |
| 1996 | 0 | 66,779 | 26,360 | 0 | 93,139 |
| 1997 | 0 | 42,544 | 49,227 | 0 | 91,771 |
| 1998 | 0 | 39,728 | 48,074 | 0 | 87,802 |
| 1999 | 0 | 17,201 | 70,121 | 0 | 87,322 |
| 2000 | 0 | 15,625 | 6,382 | 0 | 22,007 |
| 2001 | 0 | 21,650 | 31,935 | 0 | 53,585 |
| 2002 | 0 | 0 | 50,244 | 0 | 50,244 |
| 2003 | 0 | 0 | 63,217 | 0 | 63,217 |
| 2004 | 0 | 58,892 | 66,175 | 0 | 125,067 |
| 2005 | 0 | 15,695 | 77,335 | 9,985 | 103,014 |
| 2006 | 0 | 14,319 | 65,289 | 15,136 | 94,744 |

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| Year | Foreign | Joint- <br> venture | Shore- <br> side | Freezer- <br> trawler | Total |
| :---: | ---: | ---: | ---: | ---: | ---: |
| 2007 | 0 | 6,780 | 52,649 | 14,121 | 73,550 |
| 2008 | 0 | 3,592 | 57,795 | 13,214 | 74,602 |
| 2009 | 0 | 0 | 44,130 | 13,223 | 57,353 |
| 2010 | 0 | 8,081 | 35,362 | 13,573 | 57,016 |
| 2011 | 0 | 9,717 | 31,760 | 14,596 | 56,073 |
| 2012 | 0 | 0 | 32,147 | 14,912 | 47,059 |
| 2013 | 0 | 0 | 33,665 | 18,584 | 52,249 |
| 2014 | 0 | 0 | 13,326 | 21,792 | 35,118 |
| 2015 | 0 | 0 | 16,775 | 22,909 | 39,684 |
| 2016 | 0 | 0 | 35,012 | 34,731 | 69,743 |
| 2017 | 0 | 5,608 | 43,427 | 37,686 | 86,721 |
| 2018 | 0 | 2,724 | 50,747 | 41,942 | 95,413 |
| 2019 | 0 | 0 | 40,794 | 54,218 | 95,013 |
| 2020 | 0 | 0 | 30,085 | 62,404 | 92,489 |
| 2021 | 0 | 0 | 11,269 | 45,807 | 57,076 |
| 2022 | 0 | 0 | 3,868 | 27,803 | 31,671 |
| 2023 | 0 | 0 | 3,657 | 19,901 | 23,557 |

Table 3. Pacific Hake landings and management decisions, 1966-2023. A dash (-) indicates the management decision was either not specified or was unknown to the authors at the time of this assessment.

| Year | U.S. landings | Canada landings | Total landings | U.S. prop. of total catch | Canada prop. <br> of total catch | U.S. catch target | Canada catch target | Total catch target | U.S. prop. of catch target removed | Canada prop. of catch target removed | Total prop. of catch target removed |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1966 | 137,000 | 700 | 137,700 | 99.5\% | 0.5\% | - | - | - | - | - | - |
| 1967 | 177,660 | 36,710 | 214,370 | 82.9\% | 17.1\% | - | - | - | - | - | - |
| 1968 | 60,820 | 61,360 | 122,180 | 49.8\% | 50.2\% | - | - | - | - | - | - |
| 1969 | 86,280 | 93,850 | 180,130 | 47.9\% | 52.1\% | - | - | - | - | - | - |
| 1970 | 159,580 | 75,010 | 234,590 | 68.0\% | 32.0\% | - | - | - | - | - | - |
| 1971 | 127,920 | 26,700 | 154,620 | 82.7\% | 17.3\% | - | - | - | - | - | - |
| 1972 | 74,130 | 43,410 | 117,540 | 63.1\% | 36.9\% | - | - | - | - | - | - |
| 1973 | 147,510 | 15,130 | 162,640 | 90.7\% | 9.3\% | - | - | - | - | - | - |
| 1974 | 194,110 | 17,150 | 211,260 | 91.9\% | 8.1\% | - | - | - | - | - | - |
| 1975 | 205,650 | 15,700 | 221,350 | 92.9\% | 7.1\% | - | - | - | - | - | - |
| 1976 | 231,550 | 5,970 | 237,520 | 97.5\% | 2.5\% | - | - | - | - | - | - |
| 1977 | 127,500 | 5,190 | 132,690 | 96.1\% | 3.9\% | - | - | - | - | - | - |
| 1978 | 98,377 | 5,260 | 103,637 | 94.9\% | 5.1\% | 130,000 | - | - | 75.7\% | - | - |
| 1979 | 124,680 | 12,430 | 137,110 | 90.9\% | 9.1\% | 198,900 | 35,000 | - | 62.7\% | 35.5\% | - |
| 1980 | 72,350 | 17,580 | 89,930 | 80.5\% | 19.5\% | 175,000 | 35,000 | - | 41.3\% | 50.2\% | - |
| 1981 | 114,760 | 24,360 | 139,120 | 82.5\% | 17.5\% | 175,000 | 35,000 | - | 65.6\% | 69.6\% | - |
| 1982 | 75,577 | 32,160 | 107,737 | 70.1\% | 29.9\% | 175,000 | 35,000 | - | 43.2\% | 91.9\% | - |
| 1983 | 73,151 | 40,780 | 113,931 | 64.2\% | 35.8\% | 175,000 | 45,000 | - | 41.8\% | 90.6\% | - |
| 1984 | 96,382 | 42,110 | 138,492 | 69.6\% | 30.4\% | 175,000 | 45,000 | 270,000 | 55.1\% | 93.6\% | 51.3\% |
| 1985 | 85,439 | 24,960 | 110,399 | 77.4\% | 22.6\% | 175,000 | 50,000 | 212,000 | 48.8\% | 49.9\% | 52.1\% |
| 1986 | 154,932 | 55,650 | 210,582 | 73.6\% | 26.4\% | 295,800 | 75,000 | 405,000 | 52.4\% | 74.2\% | 52.0\% |
| 1987 | 160,448 | 73,700 | 234,148 | 68.5\% | 31.5\% | 195,000 | 75,000 | 264,000 | 82.3\% | 98.3\% | 88.7\% |
| 1988 | 160,690 | 88,150 | 248,840 | 64.6\% | 35.4\% | 232,000 | 98,000 | 327,000 | 69.3\% | 89.9\% | 76.1\% |
| 1989 | 203,049 | 95,029 | 298,079 | 68.1\% | 31.9\% | 225,000 | 98,000 | 323,000 | 90.2\% | 97.0\% | 92.3\% |
| 1990 | 185,142 | 76,144 | 261,286 | 70.9\% | 29.1\% | 196,000 | 73,500 | 245,000 | 94.5\% | 103.6\% | 106.6\% |
| 1991 | 229,789 | 89,917 | 319,705 | 71.9\% | 28.1\% | 228,000 | 98,000 | 253,000 | 100.8\% | 91.8\% | 126.4\% |
| 1992 | 210,829 | 88,822 | 299,650 | 70.4\% | 29.6\% | 208,800 | 90,000 | 232,000 | 101.0\% | 98.7\% | 129.2\% |
| 1993 | 140,132 | 58,773 | 198,905 | 70.5\% | 29.5\% | 142,000 | 61,000 | 178,000 | 98.7\% | 96.3\% | 111.7\% |
| 1994 | 253,477 | 108,930 | 362,407 | 69.9\% | 30.1\% | 260,000 | 110,000 | 325,000 | 97.5\% | 99.0\% | 111.5\% |
| 1995 | 177,124 | 72,372 | 249,495 | 71.0\% | 29.0\% | 178,400 | 76,500 | 223,000 | 99.3\% | 94.6\% | 111.9\% |
| 1996 | 213,159 | 93,139 | 306,299 | 69.6\% | 30.4\% | 212,000 | 91,000 | 265,000 | 100.5\% | 102.4\% | 115.6\% |


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| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | U.S. landings | Canada landings | Total landings | U.S. prop. of total catch | Canada prop. <br> of total catch | U.S. catch target | Canada catch target | Total catch target | U.S. prop. of catch target removed | Canada prop. of catch target removed | Total prop. of catch target removed |
| 1997 | 233,376 | 91,771 | 325,147 | 71.8\% | 28.2\% | 232,000 | 99,400 | 290,000 | 100.6\% | 92.3\% | 112.1\% |
| 1998 | 232,920 | 87,802 | 320,722 | 72.6\% | 27.4\% | 232,000 | 80,000 | 290,000 | 100.4\% | 109.8\% | 110.6\% |
| 1999 | 224,565 | 87,322 | 311,887 | 72.0\% | 28.0\% | 232,000 | 90,300 | 290,000 | 96.8\% | 96.7\% | 107.5\% |
| 2000 | 206,770 | 22,007 | 228,777 | 90.4\% | 9.6\% | 232,000 | 90,300 | 290,000 | 89.1\% | 24.4\% | 78.9\% |
| 2001 | 173,940 | 53,585 | 227,525 | 76.4\% | 23.6\% | 190,400 | 81,600 | 238,000 | 91.4\% | 65.7\% | 95.6\% |
| 2002 | 130,453 | 50,244 | 180,697 | 72.2\% | 27.8\% | 129,600 | - | 162,000 | 100.7\% | - | 111.5\% |
| 2003 | 141,945 | 63,217 | 205,162 | 69.2\% | 30.8\% | 148,200 | - | 228,000 | 95.8\% | - | 90.0\% |
| 2004 | 217,240 | 125,067 | 342,307 | 63.5\% | 36.5\% | 250,000 | - | 514,441 | 86.9\% | - | 66.5\% |
| 2005 | 260,120 | 103,014 | 363,135 | 71.6\% | 28.4\% | 269,069 | 95,128 | 364,197 | 96.7\% | 108.3\% | 99.7\% |
| 2006 | 266,955 | 94,744 | 361,699 | 73.8\% | 26.2\% | 269,545 | 95,297 | 364,842 | 99.0\% | 99.4\% | 99.1\% |
| 2007 | 217,682 | 73,550 | 291,231 | 74.7\% | 25.3\% | 242,591 | 85,767 | 328,358 | 89.7\% | 85.8\% | 88.7\% |
| 2008 | 248,395 | 74,602 | 322,997 | 76.9\% | 23.1\% | 269,545 | 95,297 | 364,842 | 92.2\% | 78.3\% | 88.5\% |
| 2009 | 121,324 | 57,353 | 178,677 | 67.9\% | 32.1\% | 135,939 | 48,061 | 184,000 | 89.2\% | 119.3\% | 97.1\% |
| 2010 | 170,960 | 57,016 | 227,975 | 75.0\% | 25.0\% | 193,935 | 68,565 | 262,500 | 88.2\% | 83.2\% | 86.8\% |
| 2011 | 231,261 | 56,073 | 287,334 | 80.5\% | 19.5\% | 290,903 | 102,848 | 393,751 | 79.5\% | 54.5\% | 73.0\% |
| 2012 | 160,144 | 47,059 | 207,203 | 77.3\% | 22.7\% | 186,036 | 65,773 | 251,809 | 86.1\% | 71.5\% | 82.3\% |
| 2013 | 233,578 | 52,249 | 285,828 | 81.7\% | 18.3\% | 269,745 | 95,367 | 365,112 | 86.6\% | 54.8\% | 78.3\% |
| 2014 | 264,141 | 35,118 | 299,259 | 88.3\% | 11.7\% | 316,206 | 111,794 | 428,000 | 83.5\% | 31.4\% | 69.9\% |
| 2015 | 154,160 | 39,684 | 193,844 | 79.5\% | 20.5\% | 325,072 | 114,928 | 440,000 | 47.4\% | 34.5\% | 44.1\% |
| 2016 | 262,327 | 69,743 | 332,070 | 79.0\% | 21.0\% | 367,553 | 129,947 | 497,500 | 71.4\% | 53.7\% | 66.7\% |
| 2017 | 354,129 | 86,721 | 440,849 | 80.3\% | 19.7\% | 441,433 | 156,067 | 597,500 | 80.2\% | 55.6\% | 73.8\% |
| 2018 | 318,306 | 95,413 | 413,719 | 76.9\% | 23.1\% | 441,433 | 156,067 | 597,500 | 72.1\% | 61.1\% | 69.2\% |
| 2019 | 317,002 | 95,013 | 412,015 | 76.9\% | 23.1\% | 441,433 | 156,067 | 597,500 | 71.8\% | 60.9\% | 69.0\% |
| 2020 | 287,908 | 92,489 | 380,397 | 75.7\% | 24.3\% | 424,810 | 104,480 | 529,290 | 67.8\% | 88.5\% | 71.9\% |
| 2021 | 269,473 | 57,076 | 326,549 | 82.5\% | 17.5\% | 369,400 | 104,480 | 473,880 | 72.9\% | 54.6\% | 68.9\% |
| 2022 | 291,702 | 31,671 | 323,372 | 90.2\% | 9.8\% | 402,646 | 142,354 | 545,000 | 72.4\% | 22.2\% | 59.3\% |
| 2023 | 240,424 | 23,557 | 263,981 | 91.1\% | 8.9\% | 461,750 | 163,250 | 625,000 | 52.1\% | 14.4\% | 42.2\% |

Table 4. Annual summary of U.S. and Canadian fishery sampling included in this stock assessment by fleet, 1975-2023. The majority of values are reported as number of hauls but U.S. Shore-based and Canadian Shoreside fleets are reported as the number of trips. A dash $(-)$ indicates there was no sampled catch. The number of fish with otoliths sampled per haul has varied over time but is typically small.

| Year | U.S. Foreign (hauls) | U.S. Jointventure (hauls) | U.S. <br> Mothership (hauls) | U.S. <br> Combined Mothership Catcherprocessor (hauls) | U.S. Catcherprocessor (hauls) | U.S. Shorebased (trips) | Canada Foreign (hauls) | Canada Jointventure (hauls) | Canada Shoreside (trips) | Canada <br> Freezer trawlers (hauls) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1975 | 13 | - | - | - | - | - | - | - | - | - |
| 1976 | 142 | - | - | - | - | - | - | - | - | - |
| 1977 | 320 | - | - | - | - | - | - | - | - | - |
| 1978 | 336 | 5 | - | - | - | - | - | - | - | - |
| 1979 | 99 | 17 | - | - | - | - | - | - | - | - |
| 1980 | 191 | 30 | - | - | - | - | - | - | - | - |
| 1981 | 113 | 41 | - | - | - | - | - | - | - | - |
| 1982 | 52 | 118 | - | - | - | - | - | - | - | - |
| 1983 | - | 117 | - | - | - | - | - | - | - | - |
| 1984 | 49 | 74 | - | - | - | - | - | - | - | - |
| 1985 | 37 | 19 | - | - | - | - | - | - | - | - |
| 1986 | 88 | 32 | - | - | - | - | - | - | - | - |
| 1987 | 22 | 34 | - | - | - | - | - | - | - | - |
| 1988 | 39 | 42 | - | - | - | - | - | 3 | - | - |
| 1989 | - | 77 | - | - | - | - | - | 3 | - | - |
| 1990 | - | 143 | - | - | - | 15 | - | 5 | - | - |
| 1991 | - | - | - | 116 | - | 26 | - | 18 | - | - |
| 1992 | - | - | - | 164 | - | 46 | - | 33 | - | - |
| 1993 | - | - | - | 108 | - | 36 | - | 25 | 3 | - |
| 1994 | - | - | - | 143 | - | 50 | - | 41 | 1 | - |
| 1995 | - | - | - | 61 | - | 51 | - | 35 | 3 | - |
| 1996 | - | - | - | 123 | - | 35 | - | 28 | 1 | - |
| 1997 | - | - | - | 127 | - | 65 | - | 27 | 1 | - |
| 1998 | - | - | - | 149 | - | 64 | - | 21 | 9 | - |
| 1999 | - | - | - | 389 | - | 80 | - | 14 | 26 | - |
| 2000 | - | - | - | 413 | - | 91 | - | 25 | 1 | - |
| 2001 | - | - | - | 429 | - | 82 | - | 28 | 1 | - |
| 2002 | - | - | - | 342 | - | 71 | - | - | 36 | - |

Continued on next page ...

| Year | U.S. Foreign (hauls) | U.S. Jointventure (hauls) | U.S. <br> Mothership (hauls) | U.S. Combined Mothership Catcherprocessor (hauls) | U.S. Catcherprocessor (hauls) | U.S. Shorebased (trips) | Canada Foreign (hauls) | Canada Jointventure (hauls) | Canada Shoreside (trips) | Canada <br> Freezer trawlers (hauls) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2003 | - | - | - | 358 | - | 78 | - | - | 21 | - |
| 2004 | - | - | - | 381 | - | 72 | - | 20 | 28 | - |
| 2005 | - | - | - | 499 | - | 58 | - | 11 | 31 | 14 |
| 2006 | - | - | - | 549 | - | 83 | - | 21 | 21 | 46 |
| 2007 | - | - | - | 524 | - | 68 | - | 1 | 7 | 29 |
| 2008 | - | - | 324 | - | 356 | 63 | - | - | 20 | 31 |
| 2009 | - | - | 316 | - | 278 | 65 | - | - | 7 | 19 |
| 2010 | - | - | 443 | - | 331 | 75 | - | - | 8 | 17 |
| 2011 | - | - | 481 | - | 506 | 81 | - | 2 | 4 | 7 |
| 2012 | - | - | 299 | - | 332 | 76 | - | - | 43 | 101 |
| 2013 | - | - | 409 | - | 474 | 96 | - | - | 10 | 105 |
| 2014 | - | - | 423 | - | 557 | 68 | - | - | 28 | 79 |
| 2015 | - | - | 203 | - | 431 | 84 | - | - | 6 | 74 |
| 2016 | - | - | 502 | - | 671 | 76 | - | - | 75 | 116 |
| 2017 | - | - | 353 | - | 684 | 112 | - | - | 75 | 76 |
| 2018 | - | - | 403 | - | 549 | 92 | - | - | 44 | 91 |
| 2019 | - | - | 286 | - | 494 | 129 | - | - | 37 | 104 |
| 2020 | - | - | 186 | - | 389 | 99 | - | - | 32 | - |
| 2021 | - | - | 186 | - | 409 | 124 | - | - | - | 2 |
| 2022 | - | - | 289 | - | 455 | 80 | - | - | 23 | 16 |

Table 5. Recent age-proportion data used in the assessment for the U.S. Catcher-Processor fleet. Proportions are calculated from numbers of individuals in each age group. Age 15+ is an accumulator group.

| Year | Number of fish | Number of hauls | Age (\% of total for each year) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 2014 | 1,652 | 557 | 0.00 | 4.13 | 5.17 | 71.41 | 5.98 | 8.89 | 0.89 | 2.03 | 0.89 | 0.44 | 0.09 | 0.00 | 0.00 | 0.09 | 0.00 |
| 2015 | 1,263 | 431 | 3.49 | 1.66 | 7.55 | 3.45 | 76.45 | 3.20 | 2.16 | 0.33 | 0.77 | 0.52 | 0.00 | 0.12 | 0.12 | 0.00 | 0.15 |
| 2016 | 1,995 | 671 | 0.40 | 52.87 | 2.37 | 5.57 | 2.23 | 31.31 | 1.56 | 2.06 | 0.73 | 0.20 | 0.44 | 0.20 | 0.00 | 0.04 | 0.00 |
| 2017 | 2,026 | 684 | 1.75 | 0.87 | 50.75 | 2.36 | 4.99 | 3.08 | 28.79 | 3.01 | 2.11 | 1.17 | 0.25 | 0.58 | 0.17 | 0.00 | 0.12 |
| 2018 | 1,670 | 569 | 4.58 | 35.63 | 1.05 | 27.44 | 1.90 | 2.57 | 2.83 | 19.47 | 2.22 | 1.05 | 0.30 | 0.54 | 0.15 | 0.19 | 0.09 |
| 2019 | 1,685 | 566 | 0.00 | 6.45 | 26.06 | 1.43 | 38.29 | 1.60 | 4.00 | 1.54 | 17.34 | 1.20 | 1.10 | 0.28 | 0.14 | 0.25 | 0.32 |
| 2020 | 1,281 | 433 | 0.00 | 0.14 | 9.33 | 41.91 | 1.55 | 29.82 | 1.72 | 1.63 | 1.59 | 10.41 | 0.65 | 1.01 | 0.07 | 0.05 | 0.11 |
| 2021 | 1,206 | 409 | 3.88 | 0.62 | 2.82 | 13.37 | 36.29 | 1.66 | 22.87 | 1.90 | 1.99 | 1.64 | 10.94 | 1.37 | 0.43 | 0.16 | 0.07 |
| 2022 | 1,269 | 472 | 0.89 | 47.51 | 1.65 | 1.90 | 8.54 | 19.54 | 0.74 | 12.20 | 1.58 | 0.74 | 0.44 | 2.86 | 1.08 | 0.28 | 0.04 |
| 2023 | 1,277 | 391 | 0.69 | 51.27 | 24.03 | 0.78 | 0.93 | 3.56 | 8.87 | 1.21 | 4.97 | 0.59 | 0.35 | 0.50 | 1.91 | 0.27 | 0.06 |

Table 6. Recent age-proportion data used in the assessment for the U.S. Mothership fleet. Proportions are calculated from numbers of individuals in each age group. Age 15+ is an accumulator group.

| Year | Number of fish | Number of hauls | Age (\% of total for each year) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 2014 | 1,252 | 423 | 0.00 | 5.01 | 3.50 | 74.63 | 4.75 | 7.51 | 1.01 | 1.28 | 1.00 | 0.52 | 0.11 | 0.08 | 0.00 | 0.14 | 0.47 |
| 2015 | 601 | 203 | 1.81 | 0.65 | 10.41 | 4.77 | 71.42 | 4.00 | 4.13 | 1.07 | 0.63 | 0.83 | 0.29 | 0.00 | 0.00 | 0.00 | 0.00 |
| 2016 | 1,495 | 502 | 0.53 | 59.25 | 1.45 | 5.10 | 2.44 | 26.82 | 1.54 | 1.92 | 0.38 | 0.32 | 0.09 | 0.15 | 0.00 | 0.00 | 0.00 |
| 2017 | 1,054 | 353 | 7.78 | 0.77 | 51.20 | 2.21 | 3.41 | 1.28 | 27.73 | 1.88 | 1.96 | 0.49 | 0.08 | 0.81 | 0.19 | 0.16 | 0.06 |
| 2018 | 1,230 | 414 | 16.95 | 25.30 | 1.18 | 28.83 | 1.14 | 2.28 | 1.70 | 16.82 | 2.47 | 1.24 | 0.74 | 0.32 | 0.48 | 0.49 | 0.05 |
| 2019 | 903 | 307 | 0.00 | 14.98 | 20.59 | 0.97 | 36.30 | 1.33 | 4.12 | 1.53 | 16.62 | 1.47 | 1.04 | 0.42 | 0.48 | 0.14 | 0.01 |
| 2020 | 568 | 192 | 0.00 | 0.00 | 8.62 | 40.11 | 2.40 | 28.62 | 1.49 | 2.06 | 2.51 | 11.89 | 1.12 | 0.80 | 0.39 | 0.00 | 0.00 |
| 2021 | 545 | 186 | 0.00 | 0.43 | 1.78 | 11.57 | 37.92 | 2.18 | 22.34 | 1.27 | 1.98 | 2.77 | 13.83 | 2.40 | 0.67 | 0.21 | 0.67 |
| 2022 | 840 | 299 | 1.45 | 42.64 | 1.97 | 2.83 | 6.91 | 19.41 | 1.07 | 14.51 | 1.82 | 0.47 | 0.52 | 3.82 | 1.53 | 0.43 | 0.62 |
| 2023 | 448 | 127 | 2.28 | 39.60 | 33.17 | 1.15 | 1.21 | 5.46 | 7.49 | 0.67 | 5.19 | 0.70 | 0.25 | 0.49 | 2.06 | 0.29 | 0.00 |

Table 7. Recent age-proportion data used in the assessment for the U.S. Shore-based fleet. Proportions are calculated from numbers of individuals in each age group. Age $15+$ is an accumulator group.

| Year | Number of fish | Number of trips | Age (\% of total for each year) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 2014 | 1,355 | 68 | 0.00 | 2.14 | 3.38 | 63.99 | 8.26 | 15.10 | 1.30 | 2.40 | 1.67 | 0.63 | 0.23 | 0.00 | 0.20 | 0.20 | 0.50 |
| 2015 | 1,680 | 84 | 6.12 | 1.34 | 7.42 | 4.91 | 67.24 | 4.05 | 5.06 | 0.78 | 1.05 | 1.28 | 0.24 | 0.17 | 0.00 | 0.00 | 0.32 |
| 2016 | 1,518 | 76 | 0.11 | 65.44 | 1.41 | 3.25 | 1.55 | 22.03 | 1.60 | 2.70 | 0.72 | 0.29 | 0.31 | 0.26 | 0.14 | 0.10 | 0.08 |
| 2017 | 2,235 | 112 | 3.68 | 0.71 | 35.37 | 2.63 | 3.66 | 2.50 | 43.03 | 2.89 | 2.12 | 1.66 | 0.64 | 0.53 | 0.27 | 0.11 | 0.20 |
| 2018 | 1,834 | 92 | 7.72 | 27.85 | 1.75 | 31.45 | 1.24 | 2.40 | 2.61 | 19.08 | 2.65 | 1.32 | 0.86 | 0.49 | 0.40 | 0.15 | 0.05 |
| 2019 | 2,566 | 129 | 0.00 | 15.79 | 22.48 | 0.93 | 32.19 | 1.86 | 3.29 | 1.74 | 16.71 | 1.28 | 1.61 | 0.90 | 0.54 | 0.31 | 0.37 |
| 2020 | 1,974 | 99 | 0.00 | 0.02 | 8.34 | 34.50 | 1.35 | 32.07 | 1.24 | 2.29 | 1.57 | 15.88 | 1.06 | 0.88 | 0.43 | 0.06 | 0.32 |
| 2021 | 2,480 | 124 | 0.17 | 0.26 | 1.97 | 12.69 | 34.48 | 2.73 | 25.93 | 1.92 | 2.80 | 2.08 | 11.12 | 2.27 | 0.85 | 0.22 | 0.50 |
| 2022 | 1,800 | 90 | 0.41 | 10.55 | 1.19 | 1.86 | 11.71 | 34.34 | 1.74 | 20.59 | 2.37 | 1.49 | 1.30 | 9.22 | 1.83 | 0.85 | 0.55 |
| 2023 | 1,328 | 66 | 0.14 | 16.44 | 27.02 | 2.21 | 3.54 | 7.54 | 19.29 | 1.98 | 11.59 | 2.02 | 0.88 | 1.14 | 4.34 | 0.87 | 1.00 |

Table 8. Recent age-proportion data used in the assessment for the Canadian Shoreside fleet. Proportions are calculated from numbers of individuals in each age group. Age $15+$ is an accumulator group.

| Year | Number of fish | Number of trips | Age (\% of total for each year) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 2014 | 279 | 28 | 0.00 | 0.00 | 0.18 | 15.00 | 12.67 | 23.68 | 9.31 | 14.61 | 8.57 | 1.75 | 4.76 | 0.58 | 0.43 | 0.86 | 7.61 |
| 2015 | 296 | 6 | 2.71 | 0.00 | 1.15 | 2.68 | 63.57 | 8.08 | 11.53 | 1.32 | 5.61 | 1.82 | 0.00 | 0.52 | 0.00 | 0.34 | 0.69 |
| 2016 | 188 | 19 | 0.00 | 4.67 | 0.81 | 7.51 | 3.92 | 62.23 | 5.83 | 7.35 | 1.54 | 2.10 | 0.00 | 1.22 | 0.91 | 0.27 | 1.65 |
| 2017 | 680 | 68 | 6.94 | 0.33 | 7.83 | 1.72 | 3.00 | 7.29 | 48.07 | 13.25 | 6.94 | 1.32 | 1.25 | 1.19 | 0.14 | 0.15 | 0.55 |
| 2018 | 466 | 43 | 0.50 | 5.15 | 1.91 | 22.50 | 1.23 | 4.48 | 5.93 | 35.33 | 12.44 | 4.43 | 2.61 | 1.05 | 0.96 | 1.23 | 0.24 |
| 2019 | 296 | 33 | 0.00 | 13.24 | 11.41 | 2.87 | 30.27 | 1.90 | 4.36 | 2.70 | 26.37 | 2.28 | 3.26 | 0.83 | 0.51 | 0.00 | 0.00 |
| 2020 | 1,438 | 32 | 0.00 | 0.04 | 9.59 | 19.80 | 1.37 | 30.16 | 2.71 | 3.49 | 2.56 | 24.07 | 2.86 | 2.11 | 0.22 | 0.48 | 0.54 |
| 2022 | 596 | 22 | 0.00 | 0.00 | 0.13 | 1.42 | 13.76 | 22.91 | 6.59 | 17.47 | 4.75 | 4.29 | 4.52 | 13.98 | 5.88 | 2.41 | 1.88 |
| 2023 | 413 | 12 | 0.00 | 0.31 | 5.16 | 1.37 | 2.64 | 17.33 | 31.55 | 6.31 | 13.18 | 1.35 | 2.87 | 3.12 | 9.43 | 3.12 | 2.26 |

Table 9. Recent age-proportion data used in the assessment for the Canadian Freezer trawler fleet. Proportions are calculated from numbers of individuals in each age group. Age 15+ is an accumulator group.

| Year | Number of fish | Number of hauls | Age (\% of total for each year) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 2014 | 381 | 28 | 0.00 | 0.00 | 1.16 | 18.83 | 12.41 | 27.83 | 7.19 | 10.72 | 7.84 | 2.46 | 1.80 | 0.54 | 2.00 | 1.05 | 6.18 |
| 2015 | 215 | 21 | 0.00 | 0.00 | 4.46 | 1.85 | 55.54 | 12.22 | 15.64 | 2.84 | 2.82 | 3.24 | 1.13 | 0.28 | 0.00 | 0.00 | 0.00 |
| 2016 | 352 | 40 | 0.51 | 4.54 | 0.77 | 2.23 | 9.15 | 64.20 | 6.89 | 6.91 | 1.98 | 0.76 | 0.16 | 0.70 | 0.42 | 0.00 | 0.78 |
| 2017 | 760 | 76 | 0.00 | 0.52 | 7.41 | 2.45 | 5.46 | 5.04 | 50.03 | 12.19 | 9.69 | 2.40 | 2.51 | 1.38 | 0.22 | 0.19 | 0.50 |
| 2018 | 1,225 | 91 | 0.10 | 4.67 | 0.72 | 17.63 | 2.46 | 3.96 | 5.15 | 45.58 | 9.47 | 5.25 | 2.38 | 1.15 | 0.65 | 0.56 | 0.26 |
| 2019 | 901 | 103 | 0.04 | 18.04 | 15.07 | 3.66 | 19.21 | 2.75 | 3.93 | 4.56 | 23.12 | 5.38 | 2.37 | 1.15 | 0.37 | 0.36 | 0.00 |
| 2021 | 100 | 2 | 0.00 | 0.00 | 0.00 | 17.26 | 24.00 | 6.74 | 27.37 | 2.88 | 2.88 | 9.51 | 5.37 | 1.12 | 0.00 | 2.88 | 0.00 |
| 2022 | 421 | 16 | 0.00 | 0.00 | 0.00 | 0.24 | 6.20 | 22.15 | 8.02 | 17.76 | 7.55 | 4.38 | 6.08 | 16.52 | 7.45 | 2.87 | 0.76 |
| 2023 | 369 | 14 | 0.00 | 0.00 | 0.27 | 0.40 | 3.26 | 15.99 | 21.59 | 5.06 | 19.62 | 6.11 | 3.20 | 7.55 | 13.89 | 1.95 | 1.10 |

Table 10. Aggregated fishery age-proportion data used in the base model. Proportions are calculated from numbers of individuals in each age group where the contributions from each fleet are weighted by the catch in that fleet. Sample sizes are sum of hauls and trips from individual fleets (shown in preceding tables) as described in Section 2.1.2. Age 15+ is an accumulator group.

| Year | Number of samples | Age (\% of total for each year) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 1975 | 13 | 4.61 | 33.85 | 7.43 | 1.25 | 25.40 | 5.55 | 8.03 | 10.54 | 0.95 | 0.60 | 0.87 | 0.45 | 0.00 | 0.48 | 0.00 |
| 1976 | 142 | 0.09 | 1.34 | 14.47 | 6.74 | 4.10 | 24.58 | 9.77 | 8.90 | 12.10 | 5.43 | 4.30 | 4.08 | 1.07 | 2.36 | 0.69 |
| 1977 | 320 | 0.00 | 8.45 | 3.68 | 27.47 | 3.59 | 9.11 | 22.68 | 7.60 | 6.54 | 4.02 | 3.55 | 2.31 | 0.57 | 0.31 | 0.12 |
| 1978 | 341 | 0.47 | 1.11 | 6.51 | 6.31 | 26.42 | 6.09 | 8.87 | 21.50 | 9.78 | 4.71 | 4.68 | 2.34 | 0.52 | 0.35 | 0.34 |
| 1979 | 116 | 0.00 | 6.49 | 10.24 | 9.38 | 5.72 | 17.67 | 10.26 | 17.37 | 12.76 | 4.18 | 2.88 | 0.96 | 1.65 | 0.00 | 0.44 |
| 1980 | 221 | 0.15 | 0.54 | 30.09 | 1.85 | 4.49 | 8.16 | 11.23 | 5.01 | 8.94 | 11.08 | 9.46 | 2.63 | 3.79 | 1.52 | 1.07 |
| 1981 | 154 | 19.49 | 4.03 | 1.40 | 26.73 | 3.90 | 5.55 | 3.38 | 14.68 | 3.77 | 3.19 | 10.19 | 2.31 | 0.50 | 0.16 | 0.72 |
| 1982 | 170 | 0.00 | 32.05 | 3.52 | 0.49 | 27.35 | 1.53 | 3.68 | 3.89 | 11.76 | 3.27 | 3.61 | 7.64 | 0.24 | 0.30 | 0.66 |
| 1983 | 117 | 0.00 | 0.00 | 34.14 | 4.00 | 1.82 | 23.46 | 5.13 | 5.65 | 5.30 | 9.38 | 3.91 | 3.13 | 2.26 | 1.13 | 0.70 |
| 1984 | 123 | 0.00 | 0.00 | 1.39 | 61.90 | 3.62 | 3.85 | 16.78 | 2.85 | 1.51 | 1.24 | 3.34 | 0.92 | 0.59 | 1.44 | 0.56 |
| 1985 | 57 | 0.92 | 0.11 | 0.35 | 7.24 | 66.75 | 8.41 | 5.61 | 7.11 | 2.04 | 0.53 | 0.65 | 0.25 | 0.00 | 0.00 | 0.03 |
| 1986 | 120 | 0.00 | 15.34 | 5.38 | 0.53 | 0.76 | 43.63 | 6.90 | 8.15 | 8.26 | 2.19 | 2.82 | 1.83 | 3.13 | 0.46 | 0.61 |
| 1987 | 56 | 0.00 | 0.00 | 29.58 | 2.90 | 0.14 | 1.01 | 53.26 | 0.40 | 1.25 | 7.09 | 0.00 | 0.74 | 1.86 | 1.76 | 0.00 |
| 1988 | 84 | 0.00 | 0.65 | 0.07 | 32.28 | 0.98 | 1.45 | 0.66 | 46.05 | 1.35 | 0.84 | 10.48 | 0.79 | 0.05 | 0.06 | 4.28 |
| 1989 | 80 | 0.00 | 5.62 | 2.43 | 0.29 | 50.21 | 1.26 | 0.29 | 0.08 | 35.19 | 1.80 | 0.40 | 2.32 | 0.08 | 0.00 | 0.04 |
| 1990 | 163 | 0.00 | 5.19 | 20.56 | 1.89 | 0.59 | 31.35 | 0.51 | 0.20 | 0.04 | 31.90 | 0.30 | 0.07 | 6.41 | 0.00 | 0.99 |
| 1991 | 160 | 0.00 | 3.46 | 20.37 | 19.63 | 2.52 | 0.79 | 28.26 | 1.18 | 0.14 | 0.18 | 18.69 | 0.42 | 0.00 | 3.61 | 0.74 |
| 1992 | 243 | 0.46 | 4.24 | 4.30 | 13.05 | 18.59 | 2.27 | 1.04 | 33.93 | 0.77 | 0.08 | 0.34 | 18.05 | 0.41 | 0.04 | 2.43 |
| 1993 | 172 | 0.00 | 1.05 | 23.24 | 3.26 | 12.98 | 15.67 | 1.50 | 0.81 | 27.42 | 0.67 | 0.09 | 0.12 | 12.00 | 0.05 | 1.13 |
| 1994 | 235 | 0.00 | 0.04 | 2.83 | 21.39 | 1.26 | 12.63 | 18.69 | 1.57 | 0.57 | 29.91 | 0.26 | 0.28 | 0.02 | 9.63 | 0.91 |
| 1995 | 147 | 0.62 | 1.28 | 0.47 | 6.31 | 28.97 | 1.15 | 8.05 | 20.27 | 1.58 | 0.22 | 22.42 | 0.44 | 0.45 | 0.04 | 7.74 |
| 1996 | 186 | 0.00 | 18.28 | 16.24 | 1.51 | 7.74 | 18.14 | 1.00 | 4.91 | 10.98 | 0.58 | 0.35 | 15.72 | 0.01 | 0.11 | 4.44 |
| 1997 | 220 | 0.00 | 0.74 | 29.47 | 24.95 | 1.47 | 7.84 | 12.49 | 1.80 | 3.98 | 6.67 | 1.28 | 0.22 | 6.08 | 0.73 | 2.28 |
| 1998 | 243 | 0.01 | 4.78 | 20.34 | 20.29 | 26.60 | 2.87 | 5.41 | 9.31 | 0.92 | 1.56 | 3.90 | 0.35 | 0.09 | 2.94 | 0.63 |
| 1999 | 509 | 0.06 | 10.24 | 20.36 | 17.98 | 20.06 | 13.20 | 2.69 | 3.93 | 4.01 | 0.99 | 1.54 | 2.14 | 0.39 | 0.33 | 2.07 |
| 2000 | 530 | 1.00 | 4.22 | 10.94 | 14.29 | 12.88 | 21.06 | 13.12 | 6.55 | 4.65 | 2.51 | 2.07 | 2.31 | 1.29 | 0.72 | 2.41 |
| 2001 | 540 | 0.00 | 17.34 | 16.25 | 14.25 | 15.69 | 8.56 | 12.10 | 5.99 | 1.78 | 2.23 | 1.81 | 0.70 | 1.42 | 0.69 | 1.21 |
| 2002 | 449 | 0.00 | 0.03 | 50.64 | 14.93 | 9.69 | 5.72 | 4.44 | 6.58 | 3.55 | 0.87 | 0.84 | 1.04 | 0.24 | 0.48 | 0.95 |
| 2003 | 456 | 0.00 | 0.10 | 1.39 | 67.79 | 11.66 | 3.35 | 5.01 | 3.20 | 3.15 | 2.12 | 0.88 | 0.44 | 0.54 | 0.13 | 0.23 |
| 2004 | 501 | 0.00 | 0.02 | 5.34 | 6.13 | 68.29 | 8.12 | 2.18 | 4.13 | 2.51 | 1.27 | 1.07 | 0.35 | 0.27 | 0.16 | 0.17 |
| 2005 | 613 | 0.02 | 0.57 | 0.46 | 6.56 | 5.38 | 68.72 | 7.95 | 2.36 | 2.91 | 2.21 | 1.18 | 1.09 | 0.25 | 0.09 | 0.25 |
| 2006 | 720 | 0.33 | 2.81 | 10.44 | 1.67 | 8.57 | 4.88 | 59.04 | 5.28 | 1.72 | 2.38 | 1.13 | 1.01 | 0.43 | 0.14 | 0.19 |


| Year | Number of samples | Age (\% of total for each year) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 2007 | 629 | 0.78 | 11.52 | 3.81 | 15.70 | 1.59 | 6.89 | 3.81 | 43.95 | 5.08 | 1.71 | 2.20 | 1.66 | 0.48 | 0.19 | 0.64 |
| 2008 | 763 | 0.77 | 8.85 | 30.57 | 2.24 | 14.21 | 1.09 | 3.75 | 3.47 | 28.85 | 2.69 | 1.47 | 0.68 | 0.50 | 0.22 | 0.65 |
| 2009 | 664 | 0.79 | 0.62 | 37.45 | 30.09 | 2.76 | 9.07 | 0.70 | 2.00 | 1.30 | 12.41 | 1.48 | 0.36 | 0.60 | 0.17 | 0.19 |
| 2010 | 860 | 0.03 | 25.01 | 3.32 | 35.14 | 23.84 | 2.35 | 2.57 | 0.35 | 0.46 | 0.96 | 4.33 | 1.08 | 0.28 | 0.15 | 0.15 |
| 2011 | 1,075 | 2.67 | 8.69 | 71.58 | 2.60 | 6.05 | 4.26 | 0.99 | 0.81 | 0.28 | 0.33 | 0.07 | 1.33 | 0.14 | 0.08 | 0.12 |
| 2012 | 796 | 0.18 | 40.41 | 11.58 | 33.19 | 2.45 | 5.37 | 2.58 | 1.11 | 0.66 | 0.23 | 0.35 | 0.33 | 0.92 | 0.26 | 0.40 |
| 2013 | 1,044 | 0.03 | 0.54 | 69.85 | 5.92 | 10.51 | 1.19 | 3.54 | 2.08 | 0.98 | 1.44 | 0.28 | 0.32 | 0.56 | 2.28 | 0.48 |
| 2014 | 1,104 | 0.00 | 3.29 | 3.82 | 64.81 | 7.05 | 12.20 | 1.69 | 2.90 | 1.82 | 0.68 | 0.39 | 0.07 | 0.19 | 0.22 | 0.87 |
| 2015 | 745 | 3.62 | 1.10 | 7.06 | 3.85 | 69.58 | 4.95 | 5.56 | 0.93 | 1.45 | 1.20 | 0.24 | 0.17 | 0.04 | 0.03 | 0.21 |
| 2016 | 1,308 | 0.32 | 50.49 | 1.65 | 4.68 | 2.74 | 32.54 | 2.31 | 3.00 | 0.81 | 0.44 | 0.27 | 0.33 | 0.14 | 0.06 | 0.21 |
| 2017 | 1,293 | 3.77 | 0.72 | 38.47 | 2.38 | 4.12 | 3.10 | 36.81 | 4.38 | 3.08 | 1.33 | 0.61 | 0.72 | 0.21 | 0.09 | 0.20 |
| 2018 | 1,209 | 7.15 | 25.58 | 1.37 | 27.78 | 1.51 | 2.76 | 3.04 | 22.52 | 4.00 | 1.85 | 0.97 | 0.58 | 0.41 | 0.36 | 0.10 |
| 2019 | 1,138 | 0.01 | 13.13 | 21.21 | 1.61 | 32.56 | 1.84 | 3.78 | 2.12 | 18.63 | 1.91 | 1.65 | 0.69 | 0.40 | 0.25 | 0.22 |
| 2020 | 756 | 0.00 | 0.06 | 8.84 | 36.46 | 1.55 | 30.68 | 1.57 | 2.14 | 1.78 | 14.21 | 1.08 | 1.03 | 0.28 | 0.09 | 0.23 |
| 2021 | 721 | 1.39 | 0.36 | 1.96 | 13.45 | 33.99 | 2.87 | 24.69 | 1.98 | 2.44 | 3.06 | 10.54 | 1.81 | 0.57 | 0.57 | 0.30 |
| 2022 | 899 | 0.80 | 32.37 | 1.46 | 1.96 | 9.01 | 23.93 | 1.63 | 15.48 | 2.28 | 1.18 | 1.12 | 5.89 | 1.86 | 0.67 | 0.37 |
| 2023 | 610 | 0.69 | 35.25 | 24.90 | 1.28 | 1.95 | 5.88 | 12.93 | 1.63 | 7.97 | 1.36 | 0.67 | 1.09 | 3.40 | 0.58 | 0.43 |

Table 11. Acoustic age $2+$ survey age-proportion data used in the base model. Proportions are calculated from numbers of individuals in each age group. Age $15+$ is an accumulator group.

| Year | Number of samples | Age (\% of total for each year) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 1995 | 69 | 0.00 | 20.48 | 3.26 | 1.06 | 19.33 | 1.03 | 4.03 | 16.37 | 1.44 | 0.72 | 24.86 | 0.24 | 1.67 | 0.21 | 5.32 |
| 1998 | 105 | 0.00 | 6.83 | 8.03 | 17.03 | 17.25 | 1.77 | 11.37 | 10.79 | 1.73 | 4.19 | 7.60 | 1.27 | 0.34 | 9.74 | 2.06 |
| 2001 | 57 | 0.00 | 50.62 | 10.95 | 15.12 | 7.86 | 3.64 | 3.84 | 2.60 | 1.30 | 1.34 | 0.65 | 0.68 | 0.87 | 0.15 | 0.39 |
| 2003 | 71 | 0.00 | 23.06 | 1.63 | 43.40 | 13.07 | 2.71 | 5.14 | 3.43 | 1.82 | 2.44 | 1.44 | 0.49 | 0.43 | 0.42 | 0.52 |
| 2005 | 47 | 0.00 | 19.07 | 1.23 | 5.10 | 4.78 | 50.67 | 6.99 | 2.50 | 3.99 | 2.45 | 1.71 | 0.74 | 0.48 | 0.14 | 0.16 |
| 2007 | 69 | 0.00 | 28.29 | 2.16 | 11.64 | 1.38 | 5.01 | 3.25 | 38.64 | 3.92 | 1.94 | 1.70 | 0.83 | 0.77 | 0.34 | 0.12 |
| 2009 | 72 | 0.00 | 0.55 | 29.33 | 40.21 | 2.29 | 8.22 | 1.25 | 1.79 | 1.93 | 8.32 | 3.63 | 1.44 | 0.28 | 0.48 | 0.26 |
| 2011 | 46 | 0.00 | 27.62 | 56.32 | 3.71 | 2.64 | 2.94 | 0.70 | 0.78 | 0.38 | 0.66 | 0.97 | 2.10 | 0.76 | 0.31 | 0.11 |
| 2012 | 94 | 0.00 | 62.12 | 9.78 | 16.70 | 2.26 | 2.92 | 1.94 | 1.01 | 0.50 | 0.23 | 0.27 | 0.66 | 0.98 | 0.51 | 0.12 |
| 2013 | 67 | 0.00 | 2.17 | 74.97 | 5.63 | 8.68 | 0.95 | 2.20 | 2.59 | 0.71 | 0.35 | 0.10 | 0.13 | 0.36 | 0.77 | 0.38 |
| 2015 | 78 | 0.00 | 7.45 | 9.19 | 4.38 | 58.98 | 4.88 | 7.53 | 1.69 | 1.68 | 1.64 | 0.95 | 0.16 | 0.29 | 0.24 | 0.92 |
| 2017 | 58 | 0.00 | 0.49 | 52.73 | 2.80 | 3.70 | 3.31 | 26.02 | 4.13 | 2.91 | 1.14 | 0.91 | 0.87 | 0.42 | 0.33 | 0.25 |
| 2019 | 75 | 0.00 | 10.72 | 27.23 | 1.51 | 31.31 | 2.50 | 3.18 | 2.68 | 16.12 | 2.28 | 0.96 | 0.36 | 0.38 | 0.47 | 0.28 |
| 2021 | 65 | 0.00 | 8.03 | 5.78 | 14.04 | 28.24 | 3.49 | 20.90 | 3.06 | 2.05 | 1.95 | 9.92 | 1.50 | 0.31 | 0.22 | 0.50 |
| 2023 | 64 | 0.00 | 50.58 | 24.66 | 1.03 | 1.17 | 2.92 | 8.09 | 0.88 | 5.38 | 0.77 | 0.58 | 0.67 | 2.30 | 0.41 | 0.56 |

Table 12. Summary of the acoustic age $2+$ surveys from 1995 to 2023.

| Year | Start date | End date | Vessels | Age-2+ biomass index (million t ) | $\underset{\text { CV }}{\text { Sampling }}$ | $\begin{gathered} \hline \text { Number } \\ \text { of } \\ \text { hauls } \\ \text { with } \\ \text { age } \\ \text { samples } \\ \hline \end{gathered}$ | Age-1 index (billions of fish) | $\underset{\text { CV }}{\text { Sampling }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1995 | 1-Jul | 1-Sep | Miller Freeman Ricker | 1.318 | 0.086 | 69 | 0.232 | 0.500 |
| 1998 | 6-Jul | 27-Aug | $\begin{array}{c}\text { Miller Freeman } \\ \text { Ricker }\end{array}$ | 1.569 | 0.046 | 105 | 0.107 | 0.500 |
| 2001 | 15-Jun | 18-Aug | Miller Freeman Ricker | 0.862 | 0.102 | 57 | - | - |
| 2003 | 29-Jun | 1-Sep | Ricker | 2.138 | 0.062 | 71 | 0.024 | 0.500 |
| 2005 | 20-Jun | 19-Aug | Miller Freeman | 1.376 | 0.062 | 47 | 0.009 | 0.500 |
| 2007 | 20-Jun | 21-Aug | Miller Freeman | 0.943 | 0.074 | 69 | 1.029 | 0.500 |
| 2009 | 30-Jun | 7-Sep | $\begin{array}{c}\text { Miller Freeman } \\ \text { Ricker }\end{array}$ | 1.502 | 0.096 | 72 | 3.396 | 0.500 |
| 2011 | 26-Jun | 10-Sep | $\begin{gathered} \text { BellShimada } \\ \text { Ricker } \\ \text { Rallchima } \end{gathered}$ | 0.675 | 0.113 | 46 | 5.949 | 0.500 |
| 2012 | 23-Jun | 7-Sep | BellShimada Ricker F/V Forum Star | 1.279 | 0.065 | 94 | 0.064 | 0.500 |
| 2013 | 13-Jun | 11-Sep | BellShimada Ricker | 1.929 | 0.062 | 67 | 0.422 | 0.500 |
| 2015 | 15-Jun | 14-Sep | Bell Shimada Ricker | 2.156 | 0.081 | 78 | 4.665 | 0.500 |
| 2017 | 22-Jun | 13-Sep | Bell Shimada Nordic Pearl | 1.418 | 0.063 | 58 | 1.238 | 0.500 |
| 2019 | 13-Jun | 15-Sep | Bell Shimada Nordic Pearl | 1.718 | 0.062 | 75 | 0.734 | 0.500 |
| 2021 | 27-Jun | 24-Sep | Bell Shimada Nordic Pearl | 1.525 | 0.122 | 65 | 2.276 | 0.500 |
| 2023 | 23-Jun | 6-Sep | Bell Shimada John Franklin | 0.907 | 0.086 | 64 | 1.187 | 0.500 |

Table 13. Summary of the acoustic survey age- $2+$ biomass attributed to each country.

| Year | U.S. <br> Age-2+ <br> biomass <br> million t) | U.S. <br> sampling <br> CV <br> age-2+ | U.S. <br> prop. <br> of <br> biomass | Canada <br> Age-2+ <br> biomass <br> (million $\mathbf{t})$ | Canada <br> sampling <br> CV <br> age-2 + | Canada <br> prop. <br> of <br> biomass |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 1995 | 1.061 | 0.084 | 0.805 | 0.257 | 0.271 | 0.195 |
| 1998 | 0.606 | 0.093 | 0.386 | 0.963 | 0.047 | 0.614 |
| 2001 | 0.793 | 0.088 | 0.920 | 0.069 | 0.777 | 0.080 |
| 2003 | 1.678 | 0.063 | 0.785 | 0.459 | 0.174 | 0.215 |
| 2005 | 0.707 | 0.096 | 0.514 | 0.669 | 0.076 | 0.486 |
| 2007 | 0.683 | 0.085 | 0.724 | 0.260 | 0.149 | 0.276 |
| 2009 | 1.104 | 0.106 | 0.735 | 0.398 | 0.210 | 0.265 |
| 2011 | 0.602 | 0.104 | 0.93 | 0.072 | 0.607 | 0.107 |
| 2012 | 1.141 | 0.059 | 0.892 | 0.139 | 0.342 | 0.108 |
| 2013 | 1.805 | 0.054 | 0.936 | 0.124 | 0.568 | 0.064 |
| 2015 | 1.698 | 0.085 | 0.788 | 0.458 | 0.214 | 0.212 |
| 2017 | 1.028 | 0.073 | 0.725 | 0.390 | 0.126 | 0.275 |
| 2019 | 1.527 | 0.054 | 0.889 | 0.191 | 0.334 | 0.111 |
| 2021 | 1.459 | 0.103 | 0.957 | 0.066 | 1.641 | 0.043 |
| 2023 | 0.885 | 0.071 | 0.976 | 0.022 | 2.113 | 0.024 |

Table 14. Number of Pacific Hake ovaries collected for histological analysis and included in the estimates of time-varying maturity. Note that data from 2023 have yet to be analyzed.

| Year | NWFSC <br> Trawl <br> Survey | U.S. <br> Acoustic <br> Survey/ <br> Research | U.S. <br> At-Sea <br> Hake <br> Prserver <br> Program | Total |
| :---: | ---: | ---: | ---: | ---: |
| 2009 | 244 | 0 | 0 | $\mathbf{2 4 4}$ |
| 2012 | 64 | 181 | 0 | $\mathbf{2 4 5}$ |
| 2013 | 63 | 186 | 135 | $\mathbf{3 8 4}$ |
| 2014 | 197 | 0 | 196 | $\mathbf{3 9 3}$ |
| 2015 | 216 | 160 | 131 | 507 |
| 2016 | 66 | 131 | 194 | $\mathbf{3 9 1}$ |
| 2017 | 102 | 57 | 177 | $\mathbf{3 3 6}$ |
| 2018 | 109 | 54 | 0 | $\mathbf{1 6 3}$ |
| 2019 | 46 | 59 | 0 | $\mathbf{1 0 5}$ |
| 2020 | 0 | 0 | 0 | $\mathbf{0}$ |
| 2021 | 0 | 68 | 0 | $\mathbf{6 8}$ |
| 2022 | 0 | 0 | 0 | $\mathbf{0}$ |
| 2023 | 0 | 76 | 0 | $\mathbf{7 6}$ |
| Total | $\mathbf{1 , 1 0 7}$ | $\mathbf{9 7 2}$ | $\mathbf{8 3 3}$ | $\mathbf{2 , 9 1 2}$ |

Table 15. Summary of estimated model parameters and priors in the base model. The beta prior is parameterized with a mean and standard deviation (SD). The lognormal prior is parameterized with the median and SD in log space.

| Parameter | Number of parameters | Bounds (low, high) | Prior (Mean, SD) single value $=$ fixed |
| :---: | :---: | :---: | :---: |
| Stock Dynamics |  |  |  |
| $\overline{\log \left(R_{0}\right)}$ | 1 | $(13,17)$ | Uniform |
| Steepness (h) | 1 | $(0.2,1)$ | Beta (0.78, 0.11) |
| Recruitment variability ( $\sigma_{r}$ ) | - | - | 1.4 |
| Log recruitment deviations: 1946-2023 | 78 | $(-6,6)$ | Lognormal (0.00, $\sigma_{r}$ ) |
| Natural mortality ( $M$ ) | 1 | (0.05, 0.4) | Lognormal (-1.61, 0.10) |
| Data Source |  |  |  |
| Acoustic Survey |  |  |  |
| Additional variance for survey log (SE) | 1 | (0.05, 1.2) | Uniform |
| Non-parametric age-based selectivity: ages 3-6 | 4 | $(-5,9)$ | Uniform |
| Age-1 Survey |  |  |  |
| Additional variance for age-1 index log (SE) | 1 | (0.05, 1.2) | Uniform |
| Fishery Survey |  |  |  |
| Non-parametric age-based selectivity: ages 2-6 | 5 | $(-5,9)$ | Uniform |
| Selectivity deviations (1991-2023, ages 2-6) | 165 | $(-10,10)$ | Normal (0.00, 1.40) |
| Data Weighting |  |  |  |
| Dirichlet-multinomial fishery likelihood, $\log \left(\theta_{\text {fishery }}\right)$ | 2 | $(-5,20)$ | Normal (0.00, 1.81) |
| Dirichlet-multinomial survey likelihood, $\log \left(\theta_{\text {survey }}\right)$ | 2 | $(-5,20)$ | Normal (0.00, 1.81) |

Table 16. Annual changes in the modeling framework used to assess Pacific Hake since 2011. Methods used to weight the age-composition data (Comp Method), i.e., McAllister-Ianelli (M-I) and Dirichlet-multinomial (D-M) approaches, are explained in the main text. The Markov chain Monte Carlo column gives the number of Markov chain Monte Carlo samples used to describe model results and produce statistical inference.

| Year | Framework | Survey | Comp Method | Num. MCMC samples | Change |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 2011 | SS3 3-20, TINSS | yes | M-I (0.100, 0.890) | 999 | Increased compatibility of SS and TINSS, except for age-composition likelihood |
| 2012 | SS3 3-23b | yes | M-I (0.120, 0.940 ) | 999 | One framework for base model; TINSS changed to CCAM |
| 2013 | SS3 3-24j | no | M-I (0.120, 0.940 ) | 999 | Developed MSE |
| 2014 | SS3 3-24s | yes | M-I (0.120, 0.940) | 999 | Time-varying fishery selectivity |
| 2015 | SS3 3-24u | no | M-I (0.120, 0.940 ) | 999 | No major changes |
| 2016 | SS3 3-24u | yes | M-I (0.110, 0.510) | 999 | Re-analyzed 1998-2015 acoustic-survey data; Removed 1995 survey data |
| 2017 | SS3 3-24u | no | M-I (0.140, 0.410$)$ | 999 | Added 1995 survey data; Increased allowable selectivity variation to 0.20 |
| 2018 | SS3 3-30-10-00 | yes | D-M (0.450, 0.920) | 2,000 | Used D-M to weight age compositions; Updated maturity and fecundity; Stopped transforming selectivity parameters |
| 2019 | SS3 3-30-10-00 | no | D-M (0.363, 0.919$)$ | 2,000 | Change to time-varying fecundity |
| 2020 | SS3 3-30-14-08 | yes | D-M (0.364, 0.912) | 2,000 | Normal prior for D-M parameters; remove sum to zero constraint for recruitment deviations |
| 2021 | SS3 3-30-16-03 | no | D-M (0.361, 0.911$)$ | 8,250 | No U-turn MCMC Sampling (adnuts) |
| 2022 | SS3 3-30-16-03 | yes | D-M (0.363, 0.930) | 12,005 | Add relative age-1 index |
| 2023 | SS3 3-30-20-00 | no | D-M (0.348, 0.930) | 8,000 | No major changes |
| 2024 | SS3 3-30-22-00 | yes | D-M (0.348, 0.930) | 8,000 | Time-varying maturity |

Table 17. Estimated numbers-at-age at the beginning of the year from the base model (posterior medians; millions).

| Year | Age |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| 1966 | 1,633 | 1,466 | 907 | 496 | 293 | 187 | 142 | 106 | 91 | 75 | 61 | 53 | 43 | 36 | 29 | 24 | 20 | 16 | 13 | 11 | 32 |
| 1967 | 4,651 | 1,294 | 1,161 | 707 | 378 | 218 | 138 | 100 | 75 | 64 | 53 | 43 | 38 | 30 | 26 | 20 | 17 | 14 | 11 | 9 | 41 |
| 1968 | 3,128 | 3,692 | 1,020 | 897 | 525 | 272 | 154 | 90 | 66 | 49 | 42 | 35 | 28 | 25 | 20 | 17 | 13 | 11 | 9 | 7 | 41 |
| 1969 | 717 | 2,485 | 2,926 | 801 | 682 | 393 | 201 | 110 | 64 | 47 | 35 | 30 | 25 | 20 | 17 | 14 | 12 | 9 | 8 | 7 | 42 |
| 1970 | 9,385 | 568 | 1,976 | 2,280 | 602 | 501 | 283 | 137 | 74 | 44 | 32 | 24 | 20 | 17 | 14 | 12 | 10 | 8 | 6 | 5 | 39 |
| 1971 | 860 | 7,412 | 448 | 1,529 | 1,706 | 436 | 356 | 187 | 90 | 49 | 29 | 21 | 16 | 13 | 11 | 9 | 8 | 6 | 5 | 4 | 33 |
| 1972 | 554 | 684 | 5,854 | 350 | 1,167 | 1,279 | 324 | 253 | 133 | 64 | 35 | 21 | 15 | 11 | 10 | 8 | 6 | 6 | 5 | 4 | 30 |
| 1973 | 6,282 | 440 | 542 | 4,584 | 270 | 892 | 970 | 238 | 186 | 97 | 47 | 26 | 15 | 11 | 8 | 7 | 6 | 5 | 4 | 3 | 28 |
| 1974 | 353 | 4,974 | 349 | 425 | 3,525 | 204 | 671 | 705 | 173 | 135 | 71 | 34 | 19 | 11 | 8 | 6 | 5 | 4 | 3 | 3 | 25 |
| 1975 | 1,948 | 281 | 3,932 | 272 | 325 | 2,651 | 152 | 478 | 501 | 123 | 96 | 50 | 25 | 13 | 8 | 6 | 4 | 4 | 3 | 2 | 22 |
| 1976 | 215 | 1,542 | 222 | 3,079 | 210 | 247 | 1,990 | 110 | 346 | 363 | 89 | 70 | 36 | 18 | 10 | 6 | 4 | 3 | 3 | 2 | 20 |
| 1977 | 6,980 | 170 | 1,218 | 175 | 2,376 | 160 | 187 | 1,468 | 81 | 254 | 267 | 65 | 51 | 27 | 13 | 7 | 4 | 3 | 2 | 2 | 18 |
| 1978 | 138 | 5,521 | 135 | 956 | 137 | 1,840 | 124 | 142 | 1,112 | 61 | 192 | 202 | 49 | 39 | 20 | 10 | 5 | 3 | 2 | 2 | 16 |
| 1979 | 1,446 | 109 | 4,364 | 106 | 746 | 106 | 1,419 | 94 | 108 | 844 | 46 | 145 | 154 | 37 | 29 | 15 | 8 | 4 | 2 | 2 | 15 |
| 1980 | 17,672 | 1,143 | 87 | 3,425 | 83 | 577 | 82 | 1,069 | 70 | 81 | 636 | 35 | 109 | 116 | 28 | 22 | 12 | 6 | 3 | 2 | 13 |
| 1981 | 274 | 13,984 | 906 | 68 | 2,671 | 64 | 446 | 62 | 813 | 54 | 62 | 484 | 27 | 83 | 89 | 22 | 17 | 9 | 4 | 2 | 12 |
| 1982 | 325 | 218 | 11,049 | 710 | 53 | 2,047 | 49 | 331 | 46 | 604 | 40 | 46 | 359 | 20 | 62 | 66 | 16 | 12 | 7 | 3 | 12 |
| 1983 | 564 | 259 | 173 | 8,686 | 552 | 41 | 1,570 | 37 | 248 | 35 | 454 | 30 | 34 | 270 | 15 | 46 | 49 | 12 | 9 | 5 | 13 |
| 1984 | 14,415 | 447 | 205 | 136 | 6,771 | 427 | 31 | 1,190 | 28 | 188 | 26 | 344 | 23 | 26 | 204 | 11 | 35 | 37 | 9 | 7 | 15 |
| 1985 | 139 | 11,394 | 354 | 161 | 105 | 5,224 | 328 | 24 | 895 | 21 | 142 | 20 | 259 | 17 | 20 | 154 | 8 | 26 | 28 | 7 | 18 |
| 1986 | 191 | 110 | 9,019 | 280 | 126 | 82 | 4,043 | 250 | 18 | 682 | 16 | 108 | 15 | 197 | 13 | 15 | 117 | 6 | 20 | 21 | 21 |
| 1987 | 6,785 | 151 | 87 | 7,074 | 216 | 97 | 62 | 3,008 | 186 | 13 | 508 | 12 | 80 | 11 | 147 | 10 | 11 | 87 | 5 | 15 | 32 |
| 1988 | 2,144 | 5,369 | 120 | 68 | 5,447 | 165 | 73 | 45 | 2,202 | 136 | 10 | 371 | 9 | 59 | 8 | 107 | 7 | 8 | 64 | 3 | 35 |
| 1989 | 114 | 1,699 | 4,246 | 94 | 53 | 4,139 | 124 | 53 | 33 | 1,606 | 99 | 7 | 271 | 6 | 43 | 6 | 78 | 5 | 6 | 46 | 29 |
| 1990 | 4,395 | 90 | 1,344 | 3,304 | 71 | 39 | 3,058 | 88 | 38 | 23 | 1,136 | 70 | 5 | 191 | 4 | 30 | 4 | 55 | 4 | 4 | 54 |
| 1991 | 1,261 | 3,475 | 71 | 1,053 | 2,534 | 54 | 30 | 2,213 | 63 | 27 | 17 | 823 | 51 | 4 | 138 | 3 | 22 | 3 | 40 | 3 | 42 |
| 1992 | 127 | 999 | 2,745 | 53 | 720 | 1,872 | 38 | 21 | 1,578 | 45 | 19 | 12 | 586 | 36 | 3 | 99 | 2 | 16 | 2 | 28 | 32 |
| 1993 | 3,230 | 100 | 789 | 2,141 | 37 | 507 | 1,378 | 26 | 15 | 1,095 | 31 | 13 | 8 | 407 | 25 | 2 | 68 | 2 | 11 | 2 | 42 |
| 1994 | 3,331 | 2,556 | 79 | 619 | 1,603 | 26 | 362 | 992 | 19 | 11 | 787 | 22 | 10 | 6 | 292 | 18 | 1 | 49 | 1 | 8 | 32 |
| 1995 | 1,271 | 2,637 | 2,023 | 62 | 477 | 1,133 | 17 | 229 | 629 | 12 | 7 | 500 | 14 | 6 | 4 | 185 | 11 | 1 | 31 | 1 | 25 |
| 1996 | 1,864 | 1,006 | 2,084 | 1,592 | 48 | 362 | 789 | 11 | 152 | 418 | 8 | 4 | 332 | 9 | 4 | 3 | 123 | 8 | 1 | 21 | 17 |
| 1997 | 1,027 | 1,476 | 793 | 1,564 | 1,162 | 34 | 264 | 498 | 7 | 96 | 263 | 5 | 3 | 210 | 6 | 3 | 2 | 78 | 5 | 0 | 24 |
| 1998 | 2,013 | 812 | 1,168 | 621 | 1,094 | 786 | 23 | 166 | 313 | 4 | 60 | 165 | 3 | 2 | 132 | 4 | 2 | 1 | 49 | 3 | 15 |
| 1999 | 12,898 | 1,594 | 642 | 901 | 391 | 747 | 465 | 14 | 103 | 195 | 3 | 38 | 103 | 2 | 1 | 82 | 2 | 1 | 1 | 30 | 11 |
| 2000 | 315 | 10,204 | 1,259 | 465 | 590 | 224 | 473 | 285 | 9 | 63 | 119 | 2 | 23 | 63 | 1 | 1 | 50 | 1 | 1 | 0 | 26 |
| 2001 | 1,256 | 250 | 8,060 | 985 | 338 | 422 | 152 | 299 | 180 | 5 | 40 | 75 | 1 | 14 | 40 | 1 | 0 | 32 | 1 | 0 | 17 |
| 2002 | 44 | 994 | 198 | 6,319 | 724 | 226 | 283 | 99 | 196 | 118 | 4 | 26 | 49 | 1 | 9 | 26 | 0 | 0 | 21 | 1 | 11 |
| 2003 | 1,715 | 35 | 785 | 155 | 4,876 | 530 | 160 | 201 | 70 | 140 | 84 | 3 | 19 | 35 | 1 | 7 | 19 | 0 | 0 | 15 | 8 |

Continued on next page ...

| Year | Age |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| 2004 | 43 | 1,356 | 27 | 620 | 121 | 3,650 | 381 | 113 | 142 | 50 | 99 | 60 | 2 | 13 | 25 | 0 | 5 | 13 | 0 | 0 | 16 |
| 2005 | 2,857 | 34 | 1,071 | 21 | 460 | 73 | 2,528 | 253 | 75 | 94 | 33 | 66 | 40 | 1 | 9 | 17 | 0 | 3 | 9 | 0 | 11 |
| 2006 | 2,076 | 2,260 | 27 | 841 | 16 | 320 | 44 | 1,614 | 162 | 48 | 60 | 21 | 42 | 25 | 1 | 6 | 11 | 0 | 2 | 6 | 7 |
| 2007 | 25 | 1,642 | 1,784 | 19 | 596 | 10 | 196 | 27 | 972 | 97 | 29 | 36 | 13 | 25 | 15 | 0 | 3 | 6 | 0 | 1 | 8 |
| 2008 | 5,629 | 20 | 1,297 | 1,353 | 12 | 388 | 6 | 113 | 15 | 562 | 56 | 17 | 21 | 7 | 15 | 9 | 0 | 2 | 4 | 0 | 5 |
| 2009 | 1,371 | 4,449 | 15 | 990 | 908 | 8 | 235 | 3 | 62 | 8 | 306 | 31 | 9 | 11 | 4 | 8 | 5 | 0 | 1 | 2 | 3 |
| 2010 | 15,979 | 1,085 | 3,515 | 12 | 670 | 619 | 5 | 154 | 2 | 41 | 6 | 202 | 20 | 6 | 8 | 3 | 5 | 3 | 0 | 1 | 3 |
| 2011 | 384 | 12,618 | 857 | 2,674 | 8 | 369 | 379 | 4 | 106 | 2 | 28 | 4 | 138 | 14 | 4 | 5 | 2 | 4 | 2 | 0 | 3 |
| 2012 | 1,575 | 303 | 9,952 | 659 | 1,581 | 5 | 253 | 265 | 2 | 74 | 1 | 19 | 3 | 96 | 10 | 3 | 4 | 1 | 2 | 2 | 2 |
| 2013 | 371 | 1,247 | 240 | 7,665 | 478 | 1,063 | 3 | 178 | 187 | 2 | 52 | 1 | 14 | 2 | 68 | 7 | 2 | 3 | 1 | 2 | 2 |
| 2014 | 8,256 | 294 | 986 | 188 | 5,658 | 350 | 769 | 2 | 118 | 124 | 1 | 34 | 1 | 9 | 1 | 45 | 4 | 1 | 2 | 1 | 3 |
| 2015 | 34 | 6,522 | 232 | 765 | 134 | 4,149 | 251 | 530 | 2 | 81 | 85 | 1 | 24 | 0 | 6 | 1 | 31 | 3 | 1 | 1 | 2 |
| 2016 | 5,638 | 27 | 5,140 | 180 | 575 | 98 | 3,015 | 184 | 390 | 1 | 60 | 63 | 1 | 17 | 0 | 5 | 1 | 23 | 2 | 1 | 3 |
| 2017 | 1,565 | 4,458 | 21 | 3,660 | 133 | 411 | 68 | 2,122 | 130 | 274 | 1 | 42 | 44 | 0 | 12 | 0 | 3 | 0 | 16 | 2 | 2 |
| 2018 | 397 | 1,236 | 3,494 | 14 | 2,594 | 91 | 286 | 44 | 1,373 | 84 | 178 | 1 | 27 | 29 | 0 | 8 | 0 | 2 | 0 | 10 | 3 |
| 2019 | 273 | 315 | 943 | 2,555 | 10 | 1,849 | 66 | 189 | 29 | 907 | 55 | 117 | 0 | 18 | 19 | 0 | 5 | 0 | 1 | 0 | 9 |
| 2020 | 4,748 | 216 | 249 | 671 | 1,851 | 7 | 1,233 | 42 | 120 | 18 | 577 | 35 | 75 | 0 | 11 | 12 | 0 | 3 | 0 | 1 | 6 |
| 2021 | 10,187 | 3,749 | 171 | 195 | 506 | 1,232 | 4 | 797 | 27 | 77 | 12 | 374 | 23 | 48 | 0 | 7 | 8 | 0 | 2 | 0 | 4 |
| 2022 | 1,881 | 8,073 | 2,964 | 132 | 149 | 353 | 804 | 3 | 512 | 17 | 50 | 8 | 240 | 14 | 31 | 0 | 5 | 5 | 0 | 1 | 3 |
| 2023 | 979 | 1,486 | 6,371 | 2,189 | 101 | 111 | 246 | 510 | 2 | 324 | 11 | 31 | 5 | 152 | 9 | 20 | 0 | 3 | 3 | 0 | 3 |
| 2024 | 980 | 772 | 1,174 | 4,877 | 1,616 | 77 | 83 | 162 | 338 | 1 | 215 | 7 | 21 | 3 | 101 | 6 | 13 | 0 | 2 | 2 | 2 |

Table 18. Estimated total biomass-at-age at the beginning of the year from the base model (posterior medians; kilotonnes).

| Year | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | Age 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1966 | 33 | 138 | 234 | 197 | 142 | 103 | 84 | 67 | 63 | 56 | 48 | 45 | 39 | 34 | 29 | 26 | 21 | 17 | 14 | 11 | 34 |
| 1967 | 93 | 122 | 299 | 281 | 182 | 120 | 82 | 63 | 52 | 48 | 42 | 37 | 35 | 29 | 25 | 22 | 18 | 15 | 12 | 10 | 45 |
| 1968 | 63 | 347 | 263 | 357 | 253 | 150 | 92 | 57 | 46 | 37 | 33 | 29 | 26 | 23 | 20 | 18 | 14 | 12 | 10 | 8 | 44 |
| 1969 | 14 | 234 | 755 | 319 | 329 | 217 | 119 | 69 | 45 | 35 | 28 | 25 | 23 | 19 | 17 | 15 | 13 | 10 | 9 | 7 | 45 |
| 1970 | 188 | 53 | 509 | 907 | 291 | 276 | 168 | 86 | 52 | 33 | 25 | 20 | 19 | 16 | 13 | 13 | 10 | 9 | 7 | 6 | 42 |
| 1971 | 17 | 697 | 116 | 608 | 824 | 240 | 211 | 118 | 63 | 37 | 23 | 18 | 15 | 13 | 11 | 10 | 8 | 7 | 6 | 5 | 36 |
| 1972 | 11 | 64 | 1,510 | 139 | 564 | 705 | 192 | 160 | 93 | 48 | 28 | 17 | 14 | 11 | 10 | 9 | 7 | 6 | 5 | 4 | 33 |
| 1973 | 126 | 41 | 140 | 1,824 | 130 | 491 | 576 | 150 | 130 | 73 | 37 | 22 | 14 | 10 | 8 | 8 | 6 | 5 | 4 | 4 | 30 |
| 1974 | 7 | 468 | 90 | 169 | 1,702 | 113 | 399 | 445 | 121 | 102 | 56 | 29 | 17 | 10 | 8 | 6 | 6 | 5 | 4 | 3 | 27 |
| 1975 | 40 | 26 | 1,132 | 114 | 185 | 1,808 | 115 | 418 | 498 | 131 | 112 | 68 | 38 | 21 | 15 | 6 | 5 | 4 | 3 | 3 | 24 |
| 1976 | 4 | 170 | 63 | 1,557 | 122 | 182 | 1,651 | 101 | 378 | 441 | 112 | 98 | 60 | 32 | 18 | 13 | 10 | 7 | 6 | 5 | 45 |
| 1977 | 179 | 14 | 361 | 76 | 1,441 | 104 | 146 | 1,274 | 81 | 294 | 333 | 86 | 76 | 44 | 24 | 14 | 8 | 6 | 5 | 4 | 35 |
| 1978 | 2 | 603 | 28 | 398 | 65 | 1,156 | 78 | 106 | 963 | 60 | 210 | 243 | 63 | 53 | 31 | 18 | 10 | 6 | 4 | 3 | 29 |
| 1979 | 30 | 9 | 1,431 | 37 | 415 | 63 | 1,048 | 69 | 97 | 864 | 51 | 186 | 217 | 53 | 45 | 28 | 14 | 8 | 4 | 3 | 27 |
| 1980 | 341 | 92 | 17 | 1,453 | 30 | 306 | 44 | 702 | 48 | 66 | 567 | 34 | 125 | 140 | 35 | 31 | 16 | 8 | 4 | 3 | 18 |
| 1981 | 4 | 1,253 | 197 | 20 | 1,364 | 26 | 251 | 35 | 580 | 39 | 52 | 453 | 28 | 95 | 108 | 28 | 22 | 12 | 6 | 3 | 16 |
| 1982 | 4 | 13 | 2,453 | 217 | 17 | 1,080 | 19 | 179 | 26 | 420 | 27 | 37 | 325 | 19 | 66 | 79 | 19 | 15 | 8 | 4 | 14 |
| 1983 | 7 | 16 | 29 | 2,983 | 205 | 15 | 887 | 15 | 148 | 21 | 328 | 22 | 30 | 249 | 15 | 54 | 57 | 14 | 11 | 6 | 15 |
| 1984 | 270 | 29 | 35 | 36 | 2,951 | 189 | 13 | 744 | 13 | 126 | 17 | 276 | 18 | 24 | 201 | 13 | 39 | 42 | 10 | 8 | 17 |
| 1985 | 2 | 1,115 | 69 | 48 | 38 | 2,884 | 172 | 12 | 683 | 12 | 109 | 15 | 246 | 16 | 21 | 183 | 10 | 32 | 33 | 8 | 22 |
| 1986 | 4 | 9 | 2,348 | 82 | 44 | 33 | 2,326 | 136 | 10 | 546 | 9 | 86 | 12 | 186 | 12 | 17 | 130 | 7 | 22 | 24 | 23 |
| 1987 | 128 | 12 | 17 | 2,601 | 70 | 35 | 24 | 1,678 | 102 | 7 | 386 | 7 | 62 | 8 | 129 | 9 | 10 | 79 | 4 | 14 | 29 |
| 1988 | 43 | 498 | 27 | 22 | 2,554 | 64 | 30 | 20 | 1,414 | 84 | 6 | 314 | 5 | 49 | 7 | 108 | 7 | 8 | 64 | 3 | 36 |
| 1989 | 2 | 154 | 1,035 | 32 | 19 | 2,128 | 49 | 22 | 15 | 1,067 | 61 | 4 | 234 | 4 | 35 | 5 | 66 | 4 | 5 | 39 | 24 |
| 1990 | 88 | 8 | 335 | 1,256 | 30 | 17 | 1,697 | 37 | 17 | 12 | 791 | 46 | 3 | 170 | 3 | 27 | 4 | 50 | 3 | 4 | 48 |
| 1991 | 28 | 327 | 17 | 406 | 1,170 | 26 | 13 | 1,301 | 30 | 13 | 9 | 608 | 36 | 2 | 126 | 2 | 15 | 2 | 27 | 2 | 29 |
| 1992 | 3 | 103 | 708 | 20 | 338 | 989 | 19 | 10 | 1,023 | 23 | 10 | 7 | 462 | 26 | 2 | 98 | 2 | 16 | 2 | 28 | 32 |
| 1993 | 59 | 8 | 188 | 724 | 14 | 230 | 660 | 12 | 7 | 647 | 14 | 6 | 4 | 277 | 16 | 1 | 41 | 1 | 6 | 1 | 25 |
| 1994 | 71 | 239 | 19 | 248 | 717 | 12 | 191 | 548 | 10 | 6 | 526 | 12 | 5 | 3 | 222 | 13 | 1 | 36 | 1 | 6 | 23 |
| 1995 | 26 | 283 | 550 | 24 | 248 | 616 | 9 | 137 | 407 | 8 | 4 | 378 | 8 | 4 | 2 | 164 | 10 | 1 | 28 | 1 | 22 |
| 1996 | 36 | 93 | 584 | 639 | 22 | 205 | 438 | 6 | 95 | 278 | 5 | 3 | 255 | 5 | 2 | 2 | 78 | 5 | 0 | 13 | 11 |
| 1997 | 22 | 134 | 203 | 684 | 573 | 18 | 161 | 296 | 4 | 65 | 184 | 3 | 2 | 166 | 3 | 2 | 1 | 49 | 3 | 0 | 15 |
| 1998 | 37 | 77 | 270 | 228 | 540 | 410 | 12 | 100 | 190 | 3 | 40 | 114 | 2 | 1 | 99 | 2 | 1 | 1 | 29 | 2 | 9 |
| 1999 | 230 | 137 | 163 | 316 | 172 | 413 | 255 | 8 | 67 | 125 | 2 | 26 | 74 | 1 | 1 | 66 | 2 | 1 | 1 | 24 | 9 |
| 2000 | 8 | 1,104 | 382 | 236 | 326 | 145 | 362 | 214 | 7 | 57 | 103 | 1 | 22 | 60 | 1 | 1 | 43 | 1 | 1 | 0 | 22 |
| 2001 | 33 | 29 | 2,465 | 477 | 216 | 275 | 109 | 251 | 154 | 5 | 39 | 71 | 1 | 14 | 40 | 1 | 0 | 31 | 1 | 0 | 16 |
| 2002 | 1 | 124 | 65 | 3,044 | 435 | 168 | 201 | 77 | 185 | 111 | 3 | 27 | 51 | 1 | 10 | 29 | 1 | 0 | 23 | 1 | 12 |
| 2003 | 43 | 4 | 239 | 70 | 2,547 | 324 | 113 | 135 | 53 | 126 | 73 | 2 | 18 | 33 | 0 | 7 | 19 | 0 | 0 | 15 | 8 |

Continued on next page

| Year | Age |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| 2004 | 1 | 146 | 8 | 268 | 61 | 1,997 | 229 | 78 | 96 | 37 | 85 | 51 | 2 | 12 | 22 | 0 | 4 | 12 | 0 | 0 | 15 |
| 2005 | 53 | 3 | 308 | 9 | 237 | 41 | 1,450 | 158 | 56 | 67 | 25 | 59 | 35 | 1 | 8 | 16 | 0 | 3 | 8 | 0 | 10 |
| 2006 | 35 | 200 | 7 | 378 | 8 | 191 | 27 | 996 | 112 | 39 | 45 | 17 | 41 | 23 | 1 | 6 | 11 | 0 | 2 | 6 | 7 |
| 2007 | 0 | 120 | 406 | 8 | 306 | 6 | 117 | 16 | 618 | 68 | 23 | 27 | 10 | 23 | 13 | 0 | 3 | 6 | 0 | 1 | 7 |
| 2008 | 94 | 2 | 294 | 539 | 7 | 257 | 4 | 82 | 12 | 436 | 47 | 16 | 19 | 7 | 16 | 10 | 0 | 2 | 4 | 0 | 6 |
| 2009 | 24 | 342 | 3 | 341 | 432 | 5 | 163 | 2 | 48 | 7 | 242 | 26 | 9 | 10 | 4 | 9 | 5 | 0 | 1 | 2 | 3 |
| 2010 | 286 | 94 | 784 | 4 | 297 | 356 | 4 | 121 | 2 | 36 | 5 | 179 | 20 | 6 | 7 | 3 | 6 | 3 | 0 | 1 | 4 |
| 2011 | 7 | 1,038 | 199 | 902 | 3 | 183 | 228 | 3 | 89 | 1 | 25 | 3 | 128 | 13 | 4 | 5 | 2 | 4 | 2 | 0 | 3 |
| 2012 | 33 | 28 | 2,266 | 240 | 657 | 2 | 136 | 172 | 2 | 68 | 1 | 19 | 3 | 92 | 10 | 3 | 4 | 1 | 3 | 2 | 2 |
| 2013 | 8 | 131 | 65 | 2,889 | 227 | 539 | 2 | 109 | 142 | 2 | 53 | 1 | 15 | 2 | 72 | 8 | 2 | 3 | 1 | 2 | 3 |
| 2014 | 192 | 32 | 306 | 86 | 2,816 | 205 | 453 | 1 | 86 | 110 | 1 | 40 | 1 | 11 | 1 | 56 | 6 | 2 | 2 | 1 | 3 |
| 2015 | 1 | 578 | 56 | 298 | 60 | 1,907 | 127 | 269 | 1 | 51 | 64 | 1 | 24 | 0 | 6 | 1 | 33 | 3 | 1 | 1 | 2 |
| 2016 | 124 | 3 | 1,244 | 67 | 272 | 50 | 1,484 | 99 | 217 | 1 | 39 | 50 | 1 | 18 | 0 | 5 | 1 | 25 | 2 | 1 | 3 |
| 2017 | 41 | 504 | 6 | 1,501 | 66 | 243 | 41 | 1,214 | 84 | 180 | 1 | 32 | 41 | 0 | 14 | 0 | 4 | 1 | 20 | 2 | 3 |
| 2018 | 8 | 160 | 1,143 | 7 | 1,371 | 55 | 192 | 30 | 914 | 62 | 128 | 0 | 23 | 29 | 0 | 11 | 0 | 3 | 0 | 14 | 3 |
| 2019 | 6 | 28 | 298 | 1,151 | 5 | 992 | 38 | 119 | 19 | 578 | 38 | 80 | 0 | 14 | 17 | 0 | 6 | 0 | 2 | 0 | 10 |
| 2020 | 119 | 23 | 62 | 333 | 1,029 | 4 | 718 | 26 | 85 | 13 | 389 | 26 | 55 | 0 | 9 | 12 | 0 | 3 | 0 | 1 | 6 |
| 2021 | 212 | 461 | 52 | 78 | 318 | 815 | 3 | 514 | 19 | 61 | 9 | 278 | 19 | 38 | 0 | 7 | 7 | 0 | 2 | 0 | 4 |
| 2022 | 37 | 825 | 1,044 | 66 | 76 | 265 | 596 | 2 | 380 | 14 | 43 | 7 | 200 | 13 | 26 | 0 | 4 | 5 | 0 | 1 | 3 |
| 2023 | 19 | 130 | 1,693 | 1,134 | 58 | 62 | 188 | 382 | 1 | 246 | 8 | 27 | 4 | 122 | 8 | 17 | 0 | 3 | 3 | 0 | 2 |
| 2024 | 22 | 79 | 349 | 2,302 | 907 | 48 | 56 | 111 | 245 | 1 | 163 | 5 | 17 | 2 | 86 | 6 | 13 | 0 | 2 | 2 | 2 |

Table 19. Estimated exploitation-fraction-at-age (catch-at-age divided by biomass-at-age at the beginning of the year) presented as a percentage for each year from the base model (posterior medians; percentage of age class removed by fishing).

| Year | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | $\begin{aligned} & \text { Age } \\ & 10 \end{aligned}$ | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1966 | 0.00 | 0.62 | 5.03 | 7.88 | 9.67 | 10.07 | 15.23 | 14.33 | 12.97 | 12.00 | 11.43 | 10.63 | 9.86 | 9.55 | 9.12 | 8.36 | 8.36 | 8.36 | 8.36 | 8.36 | 8.36 |
| 1967 | 0.00 | 1.04 | 8.37 | 13.03 | 15.84 | 16.53 | 24.68 | 23.22 | 21.01 | 19.43 | 18.52 | 17.23 | 15.97 | 15.46 | 14.78 | 13.54 | 13.54 | 13.54 | 13.54 | 13.54 | 13.54 |
| 1968 | 0.00 | 0.61 | 4.89 | 7.67 | 9.39 | 9.86 | 14.90 | 14.02 | 12.69 | 11.73 | 11.18 | 10.40 | 9.64 | 9.34 | 8.92 | 8.17 | 8.17 | 8.17 | 8.17 | 8.17 | 8.17 |
| 1969 | 0.00 | 0.83 | 6.72 | 10.60 | 12.96 | 13.52 | 20.17 | 18.98 | 17.18 | 15.89 | 15.14 | 14.08 | 13.05 | 12.64 | 12.08 | 11.07 | 11.07 | 11.07 | 11.07 | 11.07 | 11.07 |
| 1970 | 0.00 | 0.99 | 7.94 | 12.45 | 15.17 | 15.88 | 23.61 | 22.21 | 20.10 | 18.59 | 17.72 | 16.48 | 15.28 | 14.80 | 14.14 | 12.95 | 12.95 | 12.95 | 12.95 | 12.95 | 12.95 |
| 1971 | 0.00 | 0.60 | 4.86 | 7.70 | 9.46 | 9.92 | 14.94 | 14.05 | 12.72 | 11.76 | 11.21 | 10.43 | 9.67 | 9.36 | 8.95 | 8.20 | 8.20 | 8.20 | 8.20 | 8.20 | 8.20 |
| 1972 | 0.00 | 0.41 | 3.30 | 5.22 | 6.43 | 6.75 | 10.22 | 9.62 | 8.70 | 8.05 | 7.67 | 7.14 | 6.61 | 6.40 | 6.12 | 5.61 | 5.61 | 5.61 | 5.61 | 5.61 | 5.61 |
| 1973 | 0.00 | 0.46 | 3.74 | 5.94 | 7.31 | 7.64 | 11.57 | 10.88 | 9.85 | 9.11 | 8.68 | 8.07 | 7.48 | 7.25 | 6.93 | 6.34 | 6.34 | 6.34 | 6.34 | 6.34 | 6.34 |
| 1974 | 0.00 | 0.58 | 4.66 | 7.40 | 9.07 | 9.50 | 14.28 | 13.44 | 12.16 | 11.25 | 10.72 | 9.97 | 9.24 | 8.95 | 8.55 | 7.84 | 7.84 | 7.84 | 7.84 | 7.84 | 7.84 |
| 1975 | 0.00 | 0.49 | 3.45 | 5.80 | 6.35 | 6.34 | 9.26 | 8.03 | 7.08 | 6.61 | 6.04 | 5.17 | 4.50 | 4.46 | 3.78 | 6.50 | 6.50 | 6.50 | 6.50 | 6.50 | 6.50 |
| 1976 | 0.00 | 0.34 | 2.97 | 4.07 | 5.30 | 4.97 | 7.23 | 6.54 | 5.49 | 4.93 | 4.77 | 4.27 | 3.64 | 3.32 | 3.26 | 2.60 | 2.60 | 2.60 | 2.60 | 2.60 | 2.60 |
| 1977 | 0.00 | 0.28 | 1.77 | 2.97 | 3.17 | 3.53 | 4.85 | 4.35 | 3.80 | 3.26 | 3.03 | 2.87 | 2.56 | 2.28 | 2.06 | 1.91 | 1.91 | 1.91 | 1.91 | 1.91 | 1.91 |
| 1978 | 0.00 | 0.20 | 2.39 | 2.91 | 3.79 | 3.44 | 5.61 | 4.73 | 4.10 | 3.66 | 3.25 | 2.96 | 2.79 | 2.60 | 2.31 | 1.96 | 1.96 | 1.96 | 1.96 | 1.96 | 1.96 |
| 1979 | 0.00 | 0.30 | 1.75 | 4.01 | 3.79 | 4.20 | 5.59 | 5.59 | 4.56 | 4.04 | 3.73 | 3.24 | 2.94 | 2.90 | 2.69 | 2.24 | 2.24 | 2.24 | 2.24 | 2.24 | 2.24 |
| 1980 | 0.00 | 0.26 | 2.42 | 2.74 | 4.85 | 3.91 | 6.37 | 5.20 | 5.03 | 4.19 | 3.83 | 3.47 | 3.00 | 2.85 | 2.79 | 2.44 | 2.44 | 2.44 | 2.44 | 2.44 | 2.44 |
| 1981 | 0.00 | 0.38 | 3.49 | 6.21 | 5.44 | 8.22 | 9.64 | 9.63 | 7.60 | 7.50 | 6.46 | 5.79 | 5.22 | 4.72 | 4.46 | 4.11 | 4.11 | 4.11 | 4.11 | 4.11 | 4.11 |
| 1982 | 0.00 | 0.46 | 2.82 | 4.97 | 6.95 | 5.16 | 11.44 | 8.27 | 7.98 | 6.43 | 6.57 | 5.54 | 4.95 | 4.67 | 4.19 | 3.73 | 3.73 | 3.73 | 3.73 | 3.73 | 3.73 |
| 1983 | 0.00 | 0.39 | 3.10 | 3.69 | 5.12 | 6.03 | 6.59 | 8.97 | 6.26 | 6.17 | 5.15 | 5.14 | 4.33 | 4.04 | 3.79 | 3.20 | 3.20 | 3.20 | 3.20 | 3.20 | 3.20 |
| 1984 | 0.00 | 0.42 | 3.47 | 5.39 | 5.00 | 5.88 | 10.17 | 6.82 | 8.97 | 6.39 | 6.52 | 5.32 | 5.30 | 4.66 | 4.33 | 3.82 | 3.82 | 3.82 | 3.82 | 3.82 | 3.82 |
| 1985 | 0.00 | 0.21 | 2.30 | 3.69 | 4.49 | 3.56 | 6.12 | 6.50 | 4.21 | 5.65 | 4.17 | 4.17 | 3.39 | 3.53 | 3.09 | 2.70 | 2.70 | 2.70 | 2.70 | 2.70 | 2.70 |
| 1986 | 0.00 | 0.43 | 2.84 | 6.18 | 7.71 | 8.01 | 9.21 | 9.76 | 10.01 | 6.62 | 9.20 | 6.65 | 6.62 | 5.63 | 5.83 | 4.79 | 4.79 | 4.79 | 4.79 | 4.79 | 4.79 |
| 1987 | 0.00 | 0.53 | 4.76 | 6.18 | 10.46 | 11.10 | 16.79 | 11.87 | 12.15 | 12.74 | 8.71 | 11.86 | 8.54 | 8.89 | 7.51 | 7.32 | 7.32 | 7.32 | 7.32 | 7.32 | 7.32 |
| 1988 | 0.00 | 0.48 | 4.31 | 7.58 | 7.66 | 11.08 | 17.14 | 15.95 | 10.90 | 11.39 | 12.35 | 8.28 | 11.22 | 8.45 | 8.74 | 6.96 | 6.96 | 6.96 | 6.96 | 6.96 | 6.96 |
| 1989 | 0.00 | 0.68 | 5.54 | 9.66 | 13.20 | 11.37 | 23.68 | 22.65 | 20.37 | 14.21 | 15.37 | 16.32 | 10.90 | 15.45 | 11.56 | 11.26 | 11.26 | 11.26 | 11.26 | 11.26 | 11.26 |
| 1990 | 0.00 | 0.55 | 4.29 | 6.87 | 9.33 | 10.90 | 13.69 | 17.80 | 16.45 | 15.10 | 10.90 | 11.55 | 12.22 | 8.53 | 12.02 | 8.47 | 8.47 | 8.47 | 8.47 | 8.47 | 8.47 |
| 1991 | 0.00 | 0.80 | 13.10 | 30.80 | 12.34 | 13.08 | 19.44 | 15.03 | 18.89 | 17.82 | 16.93 | 11.97 | 12.63 | 13.98 | 9.70 | 12.87 | 12.87 | 12.87 | 12.87 | 12.87 | 12.87 |
| 1992 | 0.00 | 0.47 | 4.14 | 15.04 | 20.50 | 11.51 | 21.38 | 22.52 | 16.83 | 21.58 | 21.08 | 19.61 | 13.81 | 15.25 | 16.76 | 10.95 | 10.95 | 10.95 | 10.95 | 10.95 | 10.95 |
| 1993 | 0.00 | 0.35 | 2.74 | 14.16 | 17.61 | 17.84 | 16.89 | 17.63 | 17.95 | 13.69 | 18.17 | 17.37 | 16.11 | 11.87 | 13.02 | 13.47 | 13.47 | 13.47 | 13.47 | 13.47 | 13.47 |
| 1994 | 0.00 | 0.30 | 2.67 | 4.87 | 21.10 | 20.21 | 33.44 | 31.95 | 32.25 | 33.50 | 26.44 | 34.37 | 32.75 | 31.77 | 23.24 | 24.00 | 24.00 | 24.00 | 24.00 | 24.00 | 24.00 |
| 1995 | 0.00 | 0.25 | 1.72 | 3.78 | 6.30 | 19.26 | 25.89 | 23.87 | 22.04 | 22.70 | 24.41 | 18.87 | 24.44 | 24.36 | 23.46 | 16.17 | 16.17 | 16.17 | 16.17 | 16.17 | 16.17 |
| 1996 | 0.00 | 1.27 | 16.05 | 16.46 | 13.65 | 11.86 | 32.55 | 32.39 | 28.85 | 27.20 | 28.99 | 30.52 | 23.51 | 31.86 | 31.54 | 28.61 | 28.61 | 28.61 | 28.61 | 28.61 | 28.61 |
| 1997 | 0.00 | 0.34 | 2.93 | 23.33 | 25.82 | 18.39 | 29.85 | 30.61 | 29.44 | 26.77 | 26.11 | 27.26 | 28.60 | 23.04 | 31.01 | 28.91 | 28.91 | 28.91 | 28.91 | 28.91 | 28.91 |
| 1998 | 0.00 | 0.75 | 9.08 | 48.76 | 24.08 | 42.55 | 35.94 | 31.05 | 30.77 | 30.20 | 28.42 | 27.15 | 28.24 | 30.99 | 24.80 | 31.43 | 31.43 | 31.43 | 31.43 | 31.43 | 31.43 |
| 1999 | 0.00 | 1.14 | 29.01 | 43.20 | 55.49 | 32.23 | 36.84 | 37.32 | 31.15 | 31.51 | 32.01 | 29.49 | 28.08 | 30.56 | 33.30 | 25.10 | 25.09 | 25.10 | 25.10 | 25.10 | 25.10 |
| 2000 | 0.00 | 0.24 | 3.27 | 13.97 | 15.41 | 19.19 | 23.34 | 23.72 | 23.22 | 19.79 | 20.71 | 20.60 | 18.92 | 18.84 | 20.36 | 20.90 | 20.90 | 20.90 | 20.90 | 20.90 | 20.90 |
| 2001 | 0.00 | 0.25 | 2.29 | 12.91 | 21.78 | 21.07 | 20.97 | 17.88 | 17.56 | 17.55 | 15.47 | 15.86 | 15.72 | 15.10 | 14.94 | 15.20 | 15.20 | 15.20 | 15.20 | 15.20 | 15.20 |
| 2002 | 0.00 | 0.11 | 1.04 | 4.55 | 11.23 | 12.97 | 12.62 | 11.57 | 9.53 | 9.56 | 9.88 | 8.53 | 8.72 | 9.04 | 8.62 | 8.03 | 8.03 | 8.03 | 8.03 | 8.03 | 8.03 |
| 2003 | 0.00 | 0.07 | 0.62 | 2.52 | 9.08 | 13.18 | 12.79 | 13.49 | 11.95 | 10.05 | 10.43 | 10.56 | 9.09 | 9.71 | 10.00 | 8.98 | 8.98 | 8.98 | 8.98 | 8.98 | 8.98 |
| 2004 | 0.00 | 0.38 | 4.29 | 12.65 | 38.25 | 20.29 | 23.38 | 20.31 | 20.71 | 18.72 | 16.29 | 16.56 | 16.71 | 15.04 | 15.96 | 15.48 | 15.48 | 15.48 | 15.48 | 15.48 | 15.48 |
| 2005 | 0.00 | 0.19 | 1.62 | 6.04 | 20.22 | 31.16 | 30.08 | 27.65 | 23.22 | 24.17 | 22.61 | 19.27 | 19.51 | 20.60 | 18.41 | 18.40 | 18.40 | 18.40 | 18.40 | 18.40 | 18.40 |
| 2006 | 0.00 | 1.23 | 16.39 | 20.18 | 24.29 | 33.42 | 35.01 | 34.63 | 30.76 | 26.36 | 28.39 | 26.01 | 22.09 | 23.40 | 24.53 | 20.65 | 20.65 | 20.65 | 20.65 | 20.65 | 20.65 |


| Year | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | $\begin{aligned} & \text { Age } \\ & 10 \end{aligned}$ | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2007 | 0.00 | 1.13 | 16.11 | 29.93 | 30.68 | 24.45 | 40.16 | 39.53 | 37.77 | 34.25 | 30.38 | 32.04 | 29.25 | 25.99 | 27.34 | 26.99 | 26.99 | 26.99 | 26.99 | 26.99 | 26.99 |
| 2008 | 0.00 | 1.75 | 13.60 | 33.60 | 23.66 | 31.38 | 41.08 | 38.97 | 37.07 | 36.16 | 33.93 | 29.47 | 30.98 | 29.58 | 26.10 | 25.86 | 25.86 | 25.86 | 25.86 | 25.86 | 25.86 |
| 2009 | 0.00 | 0.78 | 9.36 | 37.28 | 26.16 | 15.37 | 21.24 | 20.84 | 19.10 | 18.55 | 18.72 | 17.20 | 14.89 | 16.37 | 15.53 | 12.90 | 12.90 | 12.90 | 12.90 | 12.90 | 12.90 |
| 2010 | 0.00 | 0.55 | 15.18 | 27.74 | 61.00 | 34.65 | 17.23 | 15.31 | 14.52 | 13.58 | 13.65 | 13.49 | 12.35 | 11.19 | 12.21 | 10.91 | 10.91 | 10.91 | 10.91 | 10.91 | 10.91 |
| 2011 | 0.00 | 1.78 | 10.26 | 67.18 | 31.70 | 24.07 | 16.90 | 14.06 | 12.07 | 11.68 | 11.31 | 11.13 | 10.97 | 10.50 | 9.44 | 9.71 | 9.71 | 9.71 | 9.71 | 9.71 | 9.71 |
| 2012 | 0.00 | 0.99 | 9.68 | 20.19 | 32.18 | 17.57 | 17.83 | 14.78 | 11.88 | 10.41 | 10.43 | 9.89 | 9.70 | 9.99 | 9.51 | 8.05 | 8.05 | 8.05 | 8.05 | 8.05 | 8.05 |
| 2013 | 0.00 | 0.26 | 2.90 | 15.48 | 13.89 | 14.64 | 25.63 | 23.29 | 18.66 | 15.32 | 13.89 | 13.62 | 12.87 | 13.20 | 13.51 | 12.11 | 12.11 | 12.11 | 12.11 | 12.11 | 12.11 |
| 2014 | 0.00 | 0.51 | 5.75 | 18.25 | 12.78 | 13.72 | 19.38 | 17.89 | 15.71 | 12.85 | 10.91 | 9.69 | 9.47 | 9.36 | 9.54 | 9.19 | 9.19 | 9.19 | 9.19 | 9.19 | 9.19 |
| 2015 | 0.00 | 2.35 | 5.77 | 10.95 | 13.65 | 15.69 | 12.21 | 12.24 | 10.92 | 9.79 | 8.28 | 6.89 | 6.10 | 6.23 | 6.12 | 5.87 | 5.87 | 5.87 | 5.87 | 5.87 | 5.87 |
| 2016 | 0.00 | 4.19 | 36.01 | 15.50 | 17.95 | 19.13 | 19.78 | 18.04 | 17.48 | 15.91 | 14.76 | 12.24 | 10.14 | 9.39 | 9.53 | 8.81 | 8.81 | 8.81 | 8.81 | 8.81 | 8.81 |
| 2017 | 0.00 | 6.52 | 27.54 | 22.42 | 21.67 | 17.70 | 26.86 | 28.16 | 24.82 | 24.56 | 23.13 | 21.01 | 17.36 | 15.05 | 13.84 | 13.23 | 13.23 | 13.23 | 13.23 | 13.23 | 13.23 |
| 2018 | 0.00 | 22.85 | 20.38 | 15.28 | 16.51 | 10.45 | 21.85 | 21.72 | 22.00 | 19.79 | 20.27 | 18.70 | 16.93 | 14.62 | 12.59 | 10.90 | 10.90 | 10.90 | 10.90 | 10.90 | 10.90 |
| 2019 | 0.00 | 1.78 | 28.20 | 16.20 | 17.65 | 26.08 | 30.01 | 27.17 | 26.10 | 26.99 | 25.13 | 25.20 | 23.17 | 21.94 | 18.82 | 15.26 | 15.26 | 15.26 | 15.26 | 15.26 | 15.26 |
| 2020 | 0.00 | 0.20 | 1.98 | 7.71 | 25.42 | 16.97 | 27.87 | 26.27 | 22.99 | 22.54 | 24.12 | 22.00 | 21.98 | 21.14 | 19.88 | 16.06 | 16.06 | 16.06 | 16.06 | 16.06 | 16.06 |
| 2021 | 0.00 | 1.06 | 3.92 | 7.24 | 16.12 | 23.48 | 24.16 | 25.96 | 23.65 | 21.12 | 21.43 | 22.46 | 20.41 | 21.33 | 20.37 | 18.04 | 18.04 | 18.04 | 18.04 | 18.04 | 18.04 |
| 2022 | 0.00 | 0.23 | 16.57 | 4.42 | 7.87 | 13.60 | 23.81 | 22.91 | 23.78 | 22.12 | 20.45 | 20.32 | 21.22 | 20.17 | 20.93 | 18.83 | 18.83 | 18.83 | 18.83 | 18.83 | 18.83 |
| 2023 | 0.00 | 0.58 | 10.58 | 11.07 | 4.69 | 7.30 | 18.54 | 18.94 | 17.61 | 18.66 | 17.96 | 16.26 | 16.10 | 17.59 | 16.60 | 16.23 | 16.23 | 16.23 | 16.23 | 16.23 | 16.23 |

Table 20. Estimated catch-at-age in numbers for each year from the base model (posterior medians; thousands).

| Year | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | Age <br> 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1966 | 0 | 804 | 11,444 | 15,264 | 13,766 | 10,440 | 12,805 | 9,606 | 8,410 | 6,772 | 5,523 | 4,871 | 3,975 | 3,291 | 2,607 | 2,181 | 1,771 | 1,458 | 1,178 | 939 | 2,878 |
| 1967 | 0 | 1,208 | 24,741 | 36,876 | 28,807 | 19,925 | 20,452 | 14,547 | 10,796 | 9,520 | 7,701 | 6,228 | 5,488 | 4,502 | 3,725 | 2,958 | 2,481 | 2,014 | 1,653 | 1,332 | 5,911 |
| 1968 | 0 | 2,016 | 12,766 | 27,862 | 23,867 | 14,612 | 13,782 | 8,049 | 5,744 | 4,258 | 3,739 | 3,015 | 2,441 | 2,142 | 1,755 | 1,471 | 1,149 | 972 | 788 | 651 | 3,536 |
| 1969 | 0 | 1,891 | 52,058 | 33,851 | 43,462 | 29,203 | 24,230 | 13,123 | 7,656 | 5,520 | 4,054 | 3,601 | 2,869 | 2,361 | 2,051 | 1,683 | 1,402 | 1,103 | 927 | 753 | 4,911 |
| 1970 | 0 | 507 | 40,530 | 115,976 | 44,282 | 45,222 | 40,249 | 18,922 | 10,328 | 6,027 | 4,402 | 3,227 | 2,827 | 2,293 | 1,854 | 1,615 | 1,350 | 1,105 | 874 | 743 | 5,276 |
| 1971 | 0 | 4,328 | 5,524 | 47,485 | 79,753 | 24,011 | 32,547 | 16,759 | 8,001 | 4,335 | 2,541 | 1,837 | 1,354 | 1,190 | 955 | 777 | 674 | 561 | 466 | 366 | 2,887 |
| 1972 | 0 | 254 | 50,302 | 7,175 | 36,920 | 49,421 | 19,862 | 15,631 | 8,110 | 3,858 | 2,102 | 1,229 | 895 | 658 | 577 | 465 | 377 | 328 | 273 | 228 | 1,812 |
| 1973 | 0 | 188 | 5,291 | 109,024 | 9,534 | 38,116 | 69,149 | 16,307 | 12,984 | 6,761 | 3,229 | 1,752 | 1,022 | 738 | 551 | 480 | 387 | 314 | 276 | 228 | 1,885 |
| 1974 | 0 | 2,776 | 4,235 | 12,680 | 155,594 | 10,577 | 57,894 | 62,185 | 14,853 | 11,725 | 6,087 | 2,928 | 1,569 | 915 | 666 | 492 | 430 | 347 | 283 | 247 | 2,132 |
| 1975 | 0 | 124 | 39,268 | 6,655 | 11,892 | 115,648 | 10,587 | 34,659 | 36,746 | 8,747 | 6,926 | 3,601 | 1,731 | 935 | 545 | 393 | 293 | 254 | 207 | 168 | 1,552 |
| 1976 | 0 | 591 | 1,899 | 63,565 | 6,534 | 9,181 | 120,263 | 6,547 | 21,256 | 22,733 | 5,417 | 4,295 | 2,225 | 1,071 | 576 | 335 | 244 | 180 | 159 | 128 | 1,160 |
| 1977 | 0 | 40 | 6,436 | 2,277 | 45,892 | 3,714 | 7,161 | 55,854 | 3,038 | 9,865 | 10,573 | 2,520 | 1,997 | 1,032 | 499 | 267 | 155 | 113 | 84 | 74 | 653 |
| 1978 | 0 | 1,262 | 662 | 11,586 | 2,510 | 40,036 | 4,456 | 5,091 | 39,729 | 2,176 | 7,032 | 7,510 | 1,793 | 1,421 | 735 | 355 | 191 | 111 | 81 | 60 | 558 |
| 1979 | 0 | 28 | 25,248 | 1,485 | 15,824 | 2,704 | 58,745 | 3,927 | 4,502 | 35,001 | 1,901 | 6,180 | 6,628 | 1,581 | 1,253 | 645 | 312 | 168 | 97 | 71 | 587 |
| 1980 | 0 | 244 | 415 | 39,978 | 1,427 | 12,039 | 2,819 | 36,544 | 2,437 | 2,803 | 21,798 | 1,185 | 3,838 | 4,118 | 982 | 782 | 403 | 194 | 104 | 61 | 445 |
| 1981 | 0 | 4,952 | 6,914 | 1,282 | 74,649 | 2,107 | 24,174 | 3,395 | 44,136 | 2,940 | 3,395 | 26,296 | 1,427 | 4,632 | 4,981 | 1,189 | 942 | 484 | 235 | 126 | 664 |
| 1982 | 0 | 60 | 69,759 | 10,877 | 1,211 | 56,252 | 2,165 | 14,801 | 2,076 | 27,003 | 1,795 | 2,067 | 16,023 | 874 | 2,838 | 3,029 | 723 | 570 | 300 | 143 | 533 |
| 1983 | 0 | 59 | 887 | 110,788 | 10,486 | 929 | 58,559 | 1,352 | 9,236 | 1,301 | 16,825 | 1,116 | 1,292 | 10,011 | 542 | 1,759 | 1,881 | 450 | 354 | 185 | 467 |
| 1984 | 0 | 116 | 1,208 | 1,939 | 148,338 | 11,069 | 1,350 | 50,644 | 1,175 | 7,988 | 1,124 | 14,582 | 962 | 1,114 | 8,655 | 469 | 1,522 | 1,630 | 392 | 310 | 627 |
| 1985 | 0 | 2,353 | 1,569 | 1,742 | 1,700 | 102,956 | 10,582 | 765 | 28,807 | 663 | 4,537 | 640 | 8,277 | 548 | 631 | 4,910 | 267 | 862 | 915 | 223 | 578 |
| 1986 | 0 | 36 | 67,447 | 5,006 | 3,376 | 2,586 | 214,204 | 13,211 | 955 | 36,027 | 831 | 5,687 | 801 | 10,354 | 682 | 790 | 6,153 | 335 | 1,070 | 1,147 | 1,104 |
| 1987 | 0 | 62 | 806 | 162,134 | 7,302 | 3,879 | 4,049 | 199,463 | 12,259 | 887 | 33,541 | 775 | 5,291 | 747 | 9,642 | 634 | 738 | 5,725 | 310 | 996 | 2,114 |
| 1988 | 0 | 2,436 | 1,169 | 1,636 | 197,086 | 7,050 | 5,038 | 3,117 | 154,354 | 9,496 | 685 | 25,900 | 600 | 4,087 | 574 | 7,458 | 492 | 571 | 4,421 | 239 | 2,440 |
| 1989 | 0 | 1,048 | 57,588 | 3,025 | 2,550 | 242,708 | 11,555 | 4,974 | 3,092 | 151,932 | 9,349 | 676 | 25,576 | 590 | 4,026 | 566 | 7,342 | 485 | 560 | 4,361 | 2,726 |
| 1990 | 0 | 43 | 14,419 | 86,481 | 2,773 | 1,803 | 232,179 | 6,589 | 2,827 | 1,743 | 86,090 | 5,273 | 381 | 14,444 | 334 | 2,281 | 321 | 4,169 | 275 | 317 | 4,046 |
| 1991 | 0 | 2,660 | 2,156 | 124,968 | 145,239 | 3,176 | 2,602 | 197,197 | 5,540 | 2,376 | 1,467 | 72,812 | 4,447 | 324 | 12,240 | 280 | 1,921 | 270 | 3,525 | 232 | 3,710 |
| 1992 |  | 476 | 29,979 | 2,851 | 69,769 | 115,068 | 4,121 | 2,297 | 173,148 | 4,897 | 2,092 | 1,297 | 64,055 | 3,907 | 285 | 10,762 | 248 | 1,691 | 237 | 3,102 | 3,482 |
| 1993 | 0 | 26 | 5,108 | 103,304 | 2,491 | 41,554 | 111,538 | 2,105 | 1,176 | 89,140 | 2,497 | 1,075 | 668 | 32,977 | 2,010 | 144 | 5,512 | 127 | 871 | 122 | 3,394 |
| 1994 | 0 | 716 | 477 | 11,990 | 151,229 | 2,458 | 63,787 | 175,125 | 3,306 | 1,838 | 139,554 | 3,951 | 1,692 | 1,044 | 51,754 | 3,152 | 229 | 8,677 | 201 | 1,366 | 5,541 |
| 1995 | 0 | 726 | 9,608 | 881 | 15,662 | 119,535 | 2,401 | 32,450 | 89,864 | 1,693 | 939 | 71,605 | 2,005 | 865 | 536 | 26,474 | 1,620 | 117 | 4,447 | 102 | 3,549 |
| 1996 | 0 | 1,188 | 94,583 | 105,816 | 2,912 | 23,979 | 142,294 | 2,020 | 27,377 | 75,336 | 1,417 | 792 | 60,098 | 1,697 | 730 | 451 | 22,280 | 1,356 | 98 | 3,724 | 3,087 |
| 1997 | 0 | 467 | 6,003 | 160,752 | 149,473 | 3,101 | 47,324 | 90,430 | 1,286 | 17,301 | 47,799 | 903 | 500 | 38,058 | 1,065 | 462 | 283 | 14,121 | 857 | 62 | 4,344 |
| 1998 |  | 582 | 24,646 | 112,252 | 130,732 | 175,544 | 4,179 | 30,575 | 58,223 | 827 | 11,041 | 30,889 | 581 | 320 | 24,640 | 686 | 299 | 183 | 9,121 | 555 | 2,855 |
| 1999 | 0 | 1,570 | 47,983 | 135,926 | 95,361 | 134,181 | 94,112 | 2,800 | 20,686 | 39,293 | 555 | 7,514 | 20,746 | 389 | 217 | 16,522 | 466 | 199 | 124 | 6,119 | 2,301 |
| 2000 | 0 | 2,694 | 12,736 | 33,225 | 50,606 | 27,998 | 84,167 | 50,742 | 1,532 | 11,191 | 21,173 | 303 | 4,044 | 11,244 | 213 | 118 | 8,938 | 254 | 108 | 67 | 4,555 |
| 2001 | 0 | 71 | 57,433 | 61,987 | 47,037 | 58,459 | 22,421 | 44,719 | 26,981 | 810 | 5,914 | 11,270 | 160 | 2,161 | 5,956 | 112 | 63 | 4,751 | 134 | 58 | 2,465 |
| 2002 | 0 | 143 | 650 | 139,826 | 49,136 | 21,660 | 25,252 | 8,884 | 17,626 | 10,609 | 316 | 2,327 | 4,436 | 63 | 845 | 2,350 | 44 | 24 | 1,878 | 52 | 1,001 |
| 2003 | 0 | 3 | 1,488 | 1,720 | 232,838 | 42,777 | 14,273 | 18,075 | 6,307 | 12,580 | 7,610 | 228 | 1,663 | 3,170 | 45 | 605 | 1,686 | 32 | 18 | 1,345 | 759 |
| 2004 | 0 | 571 | 327 | 34,584 | 23,499 | 407,662 | 53,254 | 15,606 | 19,794 | 6,953 | 13,845 | 8,349 | 249 | 1,830 | 3,473 | 50 | 665 | 1,848 | 35 | 19 | 2,315 |
| 2005 | 0 | 6 | 5,099 | 539 | 48,164 | 12,872 | 436,374 | 43,463 | 12,779 | 16,153 | 5,662 | 11,284 | 6,803 | 204 | 1,498 | 2,839 | 41 | 542 | 1,505 | 28 | 1,901 |
| 2006 | 0 | 2,540 | 1,196 | 77,098 | 1,994 | 64,002 | 9,368 | 345,386 | 34,377 | 10,143 | 12,760 | 4,490 | 8,917 | 5,391 | 161 | 1,187 | 2,248 | 32 | 429 | 1,191 | 1,528 |
| 2007 | 0 | 1,382 | 66,146 | 2,234 | 94,667 | 1,354 | 46,919 | 6,322 | 232,992 | 23,175 | 6,851 | 8,606 | 3,027 | 6,013 | 3,634 | 109 | 800 | 1,517 | 22 | 290 | 1,839 |
| 2008 | 0 | 27 | 40,540 | 181,756 | 1,599 | 81,013 | 1,768 | 31,806 | 4,267 | 157,874 | 15,683 | 4,620 | 5,826 | 2,044 | 4,067 | 2,460 | 74 | 541 | 1,023 | 15 | 1,446 |
| 2009 | 0 | 2,768 | 318 | 127,186 | 113,952 | 742 | 34,624 | 504 | 9,113 | 1,224 | 45,320 | 4,507 | 1,319 | 1,673 | 587 | 1,169 | 705 | 21 | 154 | 294 | 420 |
| 2010 | 0 | 524 | 119,722 | 1,182 | 181,730 | 123,709 | 618 | 18,510 | 266 | 4,871 | 652 | 24,349 | 2,407 | 703 | 891 | 315 | 623 | 378 | 11 | 83 | 384 |
| 2011 | 0 | 18,875 | 21,228 | 607,908 | 1,053 | 44,239 | 38,602 | 355 | 10,708 | 154 | 2,803 | 379 | 14,027 | 1,387 | 405 | 514 | 179 | 360 | 217 | 7 | 270 |
| 2012 | 0 | 277 | 220,913 | 49,163 | 211,980 | 421 | 23,931 | 25,324 | 231 | 7,008 | 102 | 1,845 | 249 | 9,204 | 912 | 266 | 338 | 119 | 237 | 143 | 183 |
| 2013 | 0 | 345 | 1,912 | 448,996 | 31,879 | 79,391 | 460 | 25,211 | 26,571 | 244 | 7,368 | 107 | 1,931 | 261 | 9,704 | 962 | 279 | 355 | 125 | 249 | 344 |
| 2014 | 0 | 165 | 18,117 | 16,001 | 361,417 | 28,506 | 87,913 | 249 | 13,370 | 14,119 | 129 | 3,912 | 57 | 1,025 | 138 | 5,127 | 508 | 149 | 188 | 66 | 315 |
| 2015 | 0 | 14,026 | 3,274 | 33,191 | 8,463 | 299,742 | 15,473 | 32,943 | 93 | 4,999 | 5,272 | 48 | 1,463 | 21 | 383 | 51 | 1,914 | 190 | 56 | 70 | 143 |
| 2016 | 0 | 113 | 448,494 | 10,600 | 49,434 | 9,916 | 294,038 | 17,804 | 37,667 | 106 | 5,729 | 6,037 | 56 | 1,673 | 24 | 438 | 59 | 2,201 | 218 | 64 | 246 |
| 2017 | 0 | 33,785 | 1,797 | 338,054 | 14,715 | 43,665 | 10,637 | 342,040 | 20,740 | 43,850 | 123 | 6,665 | 7,027 | 65 | 1,947 | 28 | 510 | 69 | 2,561 | 253 | 363 |
| 2018 | 0 | 37,766 | 233,090 | 1,083 | 227,091 | 5,840 | 41,772 | 6,306 | 201,691 | 12,232 | 25,869 | 73 | 3,932 | 4,143 | 38 | 1,148 | 17 | 301 | 41 | 1,508 | 366 |


| Year | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | ge 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2019 | 0 | 516 | 85,169 | 187,606 | 927 | 259,774 | 11,345 | 32,439 | 4,914 | 157,002 | 9,495 | 20,146 | 57 | 3,057 | 3,234 | 30 | 896 | 13 | 233 | 32 | 1,458 |
| 2020 | 0 | 44 | 1,264 | 26,736 | 263,752 | 686 | 200,508 | 6,772 | 19,325 | 2,926 | 93,776 | 5,665 | 12,043 | 34 | 1,827 | 1,927 | 18 | 534 | 8 | 140 | 893 |
| 2021 | 0 | 5,177 | 2,058 | 5,830 | 52,219 | 192,222 | 710 | 133,708 | 4,505 | 12,923 | 1,945 | 62,541 | 3,790 | 8,030 | 23 | 1,218 | 1,283 | 12 | 356 | 5 | 690 |
| 2022 | 0 | 1,919 | 172,880 | 2,896 | 6,164 | 36,855 | 142,112 | 476 | 90,492 | 3,042 | 8,722 | 1,317 | 42,275 | 2,555 | 5,411 | 15 | 822 | 871 | 8 | 240 | 470 |
| 2023 | 0 | 763 | 179,428 | 125,930 | 2,721 | 4,673 | 34,996 | 72,774 | 244 | 46,311 | 1,554 | 4,463 | 676 | 21,653 | 1,302 | 2,781 | 8 | 420 | 445 | 4 | 365 |

Table 21. Estimated catch-at-age in total biomass for each year from the base model (posterior medians; tonnes).

| Year |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| 1966 | 0 | 73 | 3,294 | 6,400 | 7,839 | 7,120 | 9,727 | 8,406 | 8,355 | 7,204 | 6,426 | 6,618 | 6,207 | 5,185 | 4,850 | 2,361 | 1,917 | 1,577 | 1,275 | 1,017 | 3,115 |
| 1967 | 0 | 110 | 7,120 | 15,462 | 16,405 | 13,588 | 15,536 | 12,731 | 10,725 | 10,128 | 8,960 | 8,463 | 8,571 | 7,094 | 6,932 | 3,201 | 2,685 | 2,179 | 1,789 | 1,442 | 6,397 |
| 1968 | 0 | 184 | 3,674 | 11,683 | 13,592 | 9,965 | 10,469 | 7,044 | 5,706 | 4,530 | 4,351 | 4,097 | 3,813 | 3,375 | 3,266 | 1,592 | 1,244 | 1,052 | 852 | 704 | 3,827 |
| 1969 | 0 | 173 | 14,982 | 14,194 | 24,751 | 19,915 | 18,406 | 11,484 | 7,606 | 5,872 | 4,718 | 4,893 | 4,481 | 3,720 | 3,817 | 1,821 | 1,518 | 1,194 | 1,003 | 815 | 5,315 |
| 1970 | 0 | 46 | 11,665 | 48,628 | 25,218 | 30,839 | 30,575 | 16,559 | 10,261 | 6,411 | 5,122 | 4,385 | 4,415 | 3,612 | 3,451 | 1,747 | 1,461 | 1,196 | 945 | 805 | 5,710 |
| 1971 | 0 | 395 | 1,590 | 19,910 | 45,418 | 16,374 | 24,724 | 14,666 | 7,949 | 4,612 | 2,956 | 2,496 | 2,115 | 1,876 | 1,776 | 841 | 729 | 607 | 504 | 396 | 3,124 |
| 1972 | 0 | 23 | 14,477 | 3,008 | 21,026 | 33,703 | 15,087 | 13,680 | 8,057 | 4,104 | 2,445 | 1,670 | 1,398 | 1,036 | 1,074 | 504 | 408 | 355 | 295 | 246 | 1,961 |
| 1973 | 0 | 17 | 1,523 | 45,714 | 5,430 | 25,993 | 52,528 | 14,271 | 12,899 | 7,193 | 3,758 | 2,381 | 1,596 | 1,162 | 1,026 | 519 | 419 | 339 | 298 | 247 | 2,040 |
| 1974 | 0 | 254 | 1,219 | 5,317 | 88,610 | 7,213 | 43,978 | 54,421 | 14,756 | 12,473 | 7,082 | 3,979 | 2,449 | 1,442 | 1,239 | 533 | 466 | 376 | 306 | 268 | 2,307 |
| 1975 | 0 | 11 | 11,302 | 2,790 | 6,772 | 78,866 | 8,042 | 30,331 | 36,506 | 9,306 | 8,059 | 4,893 | 2,703 | 1,473 | 1,014 | 426 | 317 | 275 | 224 | 181 | 1,680 |
| 1976 | 0 | 65 | 539 | 32,146 | 3,788 | 6,774 | 99,751 | 5,999 | 23,219 | 27,616 | 6,809 | 6,029 | 3,660 | 1,935 | 1,058 | 772 | 562 | 416 | 365 | 295 | 2,671 |
| 1977 | 0 | 3 | 1,908 | - 984 | 27,835 | 2,420 | 5,574 | 48,468 | 3,013 | 11,430 | 13,164 | 3,315 | 2,944 | 1,705 | 911 | 526 | 305 | 223 | 166 | 146 | 1,286 |
| 1978 | 0 | 138 | 137 | 4,825 | 1,199 | 25,149 | 2,823 | 3,821 | 34,403 | 2,110 | 7,696 | 9,020 | 2,283 | 1,939 | 1,131 | 642 | 345 | 200 | 146 | 109 | 1,009 |
| 1979 | 0 | 2 | 8,279 | 521 | 8,809 | 1,618 | 43,392 | 2,901 | 4,078 | 35,831 | 2,106 | 7,890 | 9,319 | 2,253 | 1,926 | 1,189 | 575 | 309 | 179 | 131 | 1,080 |
| 1980 | 0 | 20 | 81 | 16,961 | 510 | 6,393 | 1,510 | 23,973 | 1,655 | 2,286 | 19,415 | 1,166 | 4,369 | 4,934 | 1,201 | 1,096 | 565 | 273 | 146 | 85 | 623 |
| 1981 | 0 | 444 | 1,505 | 383 | 38,124 | 849 | 13,593 | 1,910 | 31,464 | 2,125 | 2,848 | 24,603 | 1,480 | 5,313 | 6,057 | 1,567 | 1,241 | 638 | 309 | 166 | 875 |
| 1982 | 0 | 4 | 15,489 | 3,320 | 399 | 29,693 | 847 | 8,009 | 1,163 | 18,781 | 1,223 | 1,669 | 14,483 | 838 | 3,028 | 3,637 | 869 | 684 | 360 | 172 | 640 |
| 1983 | 0 | 4 | 148 | 38,045 | 3,895 | 349 | 33,090 | 561 | 5,491 | 785 | 12,169 | 808 | 1,112 | 9,218 | 533 | 2,043 | 2,186 | 523 | 411 | 215 | 542 |
| 1984 | 0 | 7 | 207 | 521 | 64,638 | 4,888 | 566 | 31,677 | 559 | 5,329 | 735 | 11,679 | 774 | 1,018 | 8,525 | 524 | 1,700 | 1,820 | 437 | 346 | 700 |
| 1985 | 0 | 230 | 304 | 514 | 618 | 56,834 | 5,558 | 378 | 21,994 | 377 | 3,497 | 494 | 7,849 | 498 | 657 | 5,854 | 318 | 1,028 | 1,091 | 266 | 689 |
| 1986 | 0 | 3 | 17,559 | 1,460 | 1,180 | 1,043 | 123,226 | 7,172 | 505 | 28,818 | 479 | 4,532 | 641 | 9,739 | 620 | 873 | 6,799 | 370 | 1,182 | 1,267 | 1,220 |
| 1987 | 0 | 5 | 157 | 59,622 | 2,367 | 1,412 | 1,597 | 111,257 | 6,677 | 461 | 25,485 | 433 | 4,104 | 556 | 8,497 | 574 | 668 | 5,177 | 280 | 900 | 1,912 |
| 1988 | 0 | 226 | 267 | 517 | 92,414 | 2,730 | 2,057 | 1,367 | 99,106 | 5,832 | 388 | 21,897 | 374 | 3,384 | 460 | 7,502 | 494 | 574 | 4,447 | 240 | 2,454 |
| 1989 | 0 | 95 | 14,036 | 1,029 | 944 | 124,772 | 4,606 | 2,073 | 1,432 | 100,951 | 5,742 | 391 | 22,158 | 360 | 3,288 | 474 | 6,157 | 407 | 469 | 3,658 | 2,286 |
| 1990 | 0 | 4 | 3,596 | 32,875 | 1,157 | 767 | 128,851 | 2,812 | 1,305 | 876 | 59,993 | 3,468 | 237 | 12,856 | 211 | 2,047 | 288 | 3,740 | 247 | 284 | 3,630 |
| 1991 | 0 | 250 | 525 | 48,119 | 67,082 | 1,509 | 1,184 | 115,994 | 2,593 | 1,179 | 766 | 53,797 | 3,112 | 205 | 11,158 | 192 | 1,320 | 186 | 2,422 | 159 | 2,550 |
| 1992 | 0 | 49 | 7,738 | 1,077 | 32,785 | 60,768 | 2,101 | 1,112 | 112,205 | 2,474 | 1,082 | 721 | 50,578 | 2,794 | 186 | 10,717 | 247 | 1,684 | 236 | 3,089 | 3,468 |
| 1993 | 0 |  | 1,217 | 34,950 | 971 | 18,871 | 53,445 | 966 | 530 | 52,693 | 1,112 | 501 | 335 | 22,484 | 1,249 | 87 | 3,310 | 76 | 523 | 73 | 2,038 |
| 1994 | , | 67 | 113 | 4,798 | 67,646 | 1,187 | 33,689 | 96,783 | 1,810 | 969 | 93,210 | 2,030 | 913 | 580 | 39,327 | 2,319 | 169 | 6,384 | 148 | 1,005 | 4,077 |
| 1995 | 0 | 78 | 2,615 | 345 | 8,134 | 65,020 | 1,323 | 19,405 | 58,190 | 1,064 | 549 | 54,167 | 1,171 | 507 | 326 | 23,371 | 1,430 | 104 | 3,926 | 90 | 3,133 |
| 1996 | 0 | 110 | 26,487 | 42,474 | 1,323 | 13,542 | 79,018 | 1,127 | 17,152 | 50,069 | 884 | 469 | 46,204 | 963 | 419 | 285 | 14,078 | 857 | 62 | 2,353 | 1,951 |
| 1997 | 0 | 43 | 1,536 | 70,281 | 73,708 | 1,621 | 28,882 | 53,817 | 796 | 11,776 | 33,352 | 603 | 318 | 30,092 | 626 | 291 | 178 | 8,900 | 540 | 39 | 2,738 |
| 1998 | 0 | 55 | 5,697 | 41,246 | 64,580 | 91,638 | 2,171 | 18,392 | 35,343 | 511 | 7,257 | 21,250 | 384 | 193 | 18,557 | 407 | 177 | 109 | 5,420 | 330 | 1,697 |
| 1999 | 0 | 135 | 12,183 | 47,651 | 41,814 | 74,109 | 51,571 | 1,515 | 13,405 | 25,172 | 350 | 5,143 | 14,915 | 257 | 131 | 13,291 | 375 | 160 | 100 | 4,922 | 1,851 |
| 2000 | 0 | 292 | 3,860 | 16,885 | 27,940 | 18,115 | 64,401 | 38,201 | 1,178 | 10,101 | 18,256 | 263 | 3,817 | 10,658 | 186 | 101 | 7,638 | 217 | 92 | 58 | 3,893 |
| 2001 | 0 | 8 | 17,562 | 30,042 | 30,076 | 38,047 | 16,056 | 37,556 | 23,071 | 693 | 5,739 | 10,669 | 153 | 2,148 | 5,988 | 111 | 62 | 4,693 | 133 | 57 | 2,435 |
| 2002 | 0 | 18 | 213 | 67,364 | 29,523 | 16,087 | 17,923 | 6,880 | 16,567 | 9,946 |  | 2,443 | 4,559 | 62 | 878 | 2,621 | 50 | 27 | 2,096 | 58 | 1,116 |
| 2003 | 0 | 1 | 453 | \%75 | 121,650 | 26,115 | 10,113 | 12,136 | 4,781 | 11,339 | 6,612 | 195 | 1,658 | 2,958 | 41 | 610 | 1,700 | 32 | 18 | 1,357 | 765 |
| 2004 | 0 | 61 | 92 | 14,959 | 11,838 | 223,024 | 31,961 | 10,780 | 13,409 | 5,210 | 11,920 | 7,075 | 209 | 1,707 | 3,053 | 45 | 603 | 1,675 | 31 | 18 | 2,099 |
| 2005 | 0 | 1 | 1,465 | 230 | 24,820 | 7,238 | 250,202 | 27,111 | 9,494 | 11,530 | 4,320 | 10,101 | 6,015 | 171 | 1,403 | 2,662 | 38 | 508 | 1,411 | 27 | 1,782 |
| 2006 | 0 | 225 | 328 | 34,695 | 1,048 | 38,100 | 5,713 | 212,993 | 23,868 | 8,216 | 9,597 | 3,686 | 8,618 | 4,919 | 140 | 1,228 | 2,324 | 33 | 444 | 1,231 | 1,580 |
| 2007 | 0 | 101 | 15,050 | 890 | 48,666 | 762 | 28,066 | 3,842 | 148,174 | 16,256 | 5,418 | 6,452 | 2,486 | 5,557 | 3,194 | 97 | 712 | 1,350 | 19 | 258 | 1,637 |
| 2008 | 0 | 2 | 9,197 | 72,451 | 878 | 53,790 | 1,207 | 22,880 | 3,228 | 122,401 | 12,960 | 4,396 | 5,273 | 1,937 | 4,368 | 2,668 | 80 | 587 | 1,110 | 16 | 1,568 |
| 2009 | 0 | 213 | 68 | 43,752 | 54,194 | 456 | 24,127 | -358 | 7,061 | 976 | 35,827 | 3,877 | 1,311 | 1,513 | 560 | 1,341 | 808 | 24 | 177 | 337 | 482 |
| 2010 | 0 | 46 | 26,698 | 416 | 80,697 | 71,156 | 431 | 14,524 | 220 | 4,308 | 574 | 21,682 | 2,341 | 755 | 876 | 347 | 686 | 416 | 12 | 91 | 423 |
| 2011 | 0 | 1,553 | 4,930 | 205,058 | 441 | 21,906 | 23,247 | 257 | 9,028 | 134 | 2,522 | 346 | 13,019 | 1,344 | 436 | 538 | 188 | 377 | 228 | 7 | 283 |
| 2012 | - | 26 | 50,305 | 17,882 | 88,120 | 204 | 12,846 | 16,395 | 186 | 6,440 | 94 | 1,785 | 246 | 8,814 | 918 | 316 | 402 | 142 | 282 | 170 | 218 |
| 2013 | 0 | 36 | 519 | 169,222 | 15,102 | 40,270 | 255 | 15,375 | 20,225 | 227 | 7,535 | 111 | 2,131 | 281 | 10,201 | 1,128 | 327 | 416 | 146 | 292 | 404 |
| 2014 | 0 | 18 | 5,627 | 7,284 | 179,884 | 16,708 | 51,802 | 159 | 9,721 | 12,552 | 135 | 4,610 | 69 | 1,251 | 166 | 6,371 | 632 | 186 | 233 | 82 | 391 |
| 2015 | 0 | 1,242 | 792 | 12,912 | 3,797 | 137,758 | 7,862 | 16,689 | 53 | 3,168 | 3,947 | 43 | 1,488 | 21 | 388 | 54 | 2,023 | 200 | 59 | 74 | 151 |
| 2016 | 0 | 11 | 108,582 | 3,965 | 23,410 | 5,074 | 144,706 | 9,607 | 20,970 | 65 | 3,777 | 4,801 | 53 | 1,734 | 25 | 484 | 65 | 2,431 | 241 | 71 | 272 |
| 2017 | 0 | 3,818 | 543 | 138,594 | 7,335 | 25,815 | 6,379 | 195,664 | 13,462 | 28,767 | 86 | 5,109 | 6,521 | 69 | 2,267 | 35 | 622 | 84 | 3,119 | 309 | 443 |
| 2018 | 0 | 4,901 | 76,251 | 536 | 120,000 | 3,516 | 28,000 | 4,252 | 134,248 | 9,050 | 18,693 | 57 | 3,402 | 4,149 | 44 | 1,543 | 22 | 404 | 55 | 2,025 | 491 |


| Year |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20+ |
| 2019 | 0 | 46 | 26,902 | 84,516 | 497 | 139,366 | 6,510 | 20,554 | 3,242 | 100,146 | 6,506 | 13,766 | 42 | 2,399 | 2,958 | 33 | 1,010 | 15 | 263 | 36 | 1,645 |
| 2020 | 0 | 5 | 313 | 13,251 | 146,704 | 426 | 116,830 | 4,185 | 13,648 | 2,108 | 63,115 | 4,182 | 8,897 | 26 | 1,493 | 1,948 | 18 | 540 | 8 | 142 | 903 |
| 2021 | 0 | 637 | 630 | 2,330 | 32,843 | 127,122 | 492 | 86,246 | 3,189 | 10,242 | 1,519 | 46,613 | 3,109 | 6,303 | 19 | 1,130 | 1,191 | 11 | 331 | 5 | 641 |
| 2022 | 0 | 196 | 60,860 | 1,437 | 3,139 | 27,669 | 105,326 | 367 | 67,133 | 2,426 | 7,527 | 1,144 | 35,153 | 2,235 | 4,561 | 14 | 770 | 817 | 7 | 225 | 440 |
| 2023 | 0 | 67 | 47,681 | 65,253 | 1,563 | 2,582 | 26,768 | 54,495 | 197 | 35,191 | 1,227 | 3,892 | 595 | 17,458 | 1,112 | 2,430 | 7 | 367 | 389 | 4 | 319 |

Table 22. Calculations showing changes in biomass at each age due to natural mortality and fishing for recent strong cohorts. Start Biomass is the biomass at the beginning of the year, Catch Weight is the catch for the cohort for the year, Natural Mortality is the biomass attributed to natural mortality, and Surviving Biomass is what survives to the end of the year. Surviving Biomass does not equal the Start Biomass in the following year because the empirical weights-at-age change between years. Estimated quantities are posterior medians.

| Age | Start Biomass (kt) | Catch Weight (kt) | Natural Mortality (kt) | Surviving Biomass (kt) |
| :---: | :---: | :---: | :---: | :---: |
| 2021 cohort |  |  |  |  |
| 0 | 211.8 | 0.0 | 43.9 | 167.8 |
| 1 | 824.8 | 0.2 | 173.7 | 650.9 |
| 2 | 1,693.1 |  |  |  |
| 2020 cohort |  |  |  |  |
| 0 | 119.3 | 0.0 | 25.1 | 94.2 |
| 1 | 461.3 | 0.6 | 95.9 | 364.7 |
| 2 | 1,043.6 | 60.9 | 212.0 | 770.7 |
| 3 | 1,134.5 |  |  |  |
| 2016 cohort |  |  |  |  |
| 0 | 124.0 | 0.0 | 26.0 | 98.0 |
| 1 | 503.7 | 3.8 | 105.0 | 394.9 |
| 2 | 1,143.1 | 76.3 | 230.9 | 836.0 |
| 3 | 1,151.2 | 84.5 | 233.0 | 833.7 |
| 4 | 1,029.4 | 146.7 | 197.6 | 685.1 |
| 5 | 814.6 | 127.1 | 155.5 | 532.0 |
| 6 | 596.2 | 105.3 | 112.8 | 378.0 |
| 7 | 381.9 |  |  |  |
| 2014 cohort |  |  |  |  |
| 0 | 192.3 | 0.0 | 40.4 | 151.9 |
| 1 | 577.5 | 1.2 | 121.1 | 455.2 |
| 2 | 1,244.4 | 108.6 | 249.7 | 886.1 |
| 3 | 1,500.6 | 138.6 | 298.5 | 1,063.5 |
| 4 | 1,370.7 | 120.0 | 273.6 | 977.1 |
| 5 | 992.0 | 139.4 | 191.4 | 661.3 |
| 6 | 718.2 | 116.8 | 137.2 | 464.2 |
| 7 | 513.9 | 86.2 | 97.6 | 330.0 |
| 8 | 379.6 | 67.1 | 72.0 | 240.5 |
| 9 | 246.3 |  |  |  |
| 2010 cohort |  |  |  |  |
| 0 | 286.3 | 0.0 | 60.2 | 226.1 |
| 1 | 1,038.1 | 1.6 | 217.8 | 818.7 |
| 2 | 2,266.1 | 50.3 | 470.4 | 1,745.5 |
| 3 | 2,888.9 | 169.2 | 587.3 | 2,132.4 |
| 4 | 2,816.0 | 179.9 | 571.2 | 2,064.9 |
| 5 | 1,906.7 | 137.8 | 383.4 | 1,385.6 |
| 6 | 1,483.7 | 144.7 | 294.9 | 1,044.2 |
| 7 | 1,213.7 | 195.7 | 232.5 | 785.6 |
| 8 | 914.1 | 134.2 | 176.2 | 603.6 |
| 9 | 578.4 | 100.1 | 109.9 | 368.4 |
| 10 | 388.7 | 63.1 | 74.1 | 251.4 |
| 11 | 278.4 | 46.6 | 52.9 | 178.9 |
| 12 | 199.6 | 35.2 | 38.2 | 126.3 |
| 13 | 122.4 |  |  |  |
| 1999 cohort |  |  |  |  |
| 0 | 229.7 | 0.0 | 48.0 | 181.8 |
| 1 | 1,104.2 | 0.3 | 231.7 | 872.2 |


| ... Continued from previous page |  |  |  |  |
| ---: | ---: | ---: | ---: | ---: |
| Age | Start <br> Biomass <br> $(\mathbf{k t})$ | Catch <br> Weight <br> $(\mathbf{k t})$ | Natural <br> Mortality <br> $(\mathbf{k t})$ | Surviving <br> Biomass <br> $(\mathbf{k t})$ |
| 2 | $2,464.7$ | 17.6 | 515.0 | $1,932.2$ |
| 3 | $3,044.3$ | 67.4 | 628.0 | $2,348.9$ |
| 4 | $2,547.3$ | 121.6 | 518.7 | $1,907.0$ |
| 5 | $1,996.9$ | 223.0 | 390.6 | $1,383.2$ |
| 6 | $1,449.7$ | 250.2 | 273.9 | 925.6 |
| 7 | 995.5 | 213.0 | 182.9 | 599.7 |
| 8 | 618.4 | 148.2 | 112.8 | 357.5 |
| 9 | 435.8 | 122.4 | 76.4 | 237.0 |
| 10 | 241.7 | 35.8 | 46.5 | 159.3 |
| 11 | 179.5 | 21.7 | 34.9 | 122.9 |
| 12 | 128.1 | 13.0 | 25.6 | 89.5 |
| 13 | 92.3 | 8.8 | 18.4 | 65.2 |
| 14 | 71.5 | 10.2 | 13.9 | 47.4 |
| 15 | 56.1 | 6.4 | 11.2 | 38.5 |
| 16 | 32.8 | 2.0 | 6.6 | 24.1 |
| 17 | 25.2 | 2.4 | 5.0 | 17.8 |
| 18 | 19.6 | 3.1 | 3.8 | 12.7 |
| 19 | 14.0 | 2.0 | 0.5 | 11.4 |
| 20 | 9.6 |  |  |  |

Table 23. Time series of median posterior population estimates from the base model. Relative spawning biomass is spawning biomass relative to the unfished equilibrium $\left(B_{0}\right)$. Total biomass includes females and males of ages 0 and above. Age-2+ biomass includes females and males ages 2 and above. Exploitation fraction is total catch divided by total age- $2+$ biomass. Relative fishing intensity is ( $1-\mathrm{SPR}$ )/( $1-\mathrm{SPR}_{40 \%}$ ) such that values below $100 \%$ represent fishing below $F_{40 \%}$. In the last row, dashes (-) indicate quantities requiring 2024 catch which has not taken place yet.

| Year | Female spawning biomass (kt) | Relative spawning biomass (\%) | Total biomass (kt) | Age-2+ biomass (kt) | $\begin{gathered} \text { Age-0 } \\ \text { recruits } \\ \text { (millions) } \end{gathered}$ | Relative fishing intensity (\%) | Exploitation fraction (\%) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1966 | 954 | 49.9 | 2,389 | 2,171 | 1,633 | 47.2 | 6.3 |
| 1967 | 959 | 50.8 | 2,504 | 2,230 | 4,651 | 64.6 | 9.6 |
| 1968 | 967 | 51.1 | 2,674 | 2,245 | 3,128 | 46.5 | 5.4 |
| 1969 | 1,097 | 57.8 | 3,022 | 2,762 | 717 | 57.0 | 6.5 |
| 1970 | 1,255 | 66.4 | 3,286 | 3,028 | 9,385 | 62.7 | 7.7 |
| 1971 | 1,296 | 68.7 | 3,525 | 2,812 | 860 | 46.4 | 5.5 |
| 1972 | 1,462 | 77.2 | 3,985 | 3,909 | 554 | 35.0 | 3.0 |
| 1973 | 1,747 | 92.5 | 4,136 | 3,962 | 6,282 | 38.5 | 4.1 |
| 1974 | 1,712 | 90.4 | 4,043 | 3,568 | 353 | 44.9 | 5.9 |
| 1975 | 2,123 | 112.0 | 5,056 | 4,985 | 1,948 | 40.9 | 4.4 |
| 1976 | 2,415 | 127.4 | 5,358 | 5,181 | 215 | 40.8 | 4.6 |
| 1977 | 2,165 | 113.9 | 4,827 | 4,634 | 6,980 | 27.6 | 2.9 |
| 1978 | 1,809 | 95.2 | 4,297 | 3,690 | 138 | 26.6 | 2.8 |
| 1979 | 1,952 | 102.9 | 4,857 | 4,815 | 1,446 | 29.9 | 2.8 |
| 1980 | 1,725 | 90.8 | 4,193 | 3,755 | 17,672 | 24.8 | 2.4 |
| 1981 | 1,663 | 87.5 | 4,687 | 3,433 | 274 | 36.1 | 4.1 |
| 1982 | 1,772 | 93.4 | 5,106 | 5,086 | 325 | 30.9 | 2.1 |
| 1983 | 2,265 | 119.6 | 5,180 | 5,155 | 564 | 26.2 | 2.2 |
| 1984 | 2,327 | 122.5 | 5,122 | 4,823 | 14,415 | 28.9 | 2.9 |
| 1985 | 2,325 | 122.3 | 5,783 | 4,665 | 139 | 22.9 | 2.4 |
| 1986 | 2,316 | 122.1 | 6,114 | 6,100 | 191 | 34.0 | 3.5 |
| 1987 | 2,388 | 125.6 | 5,454 | 5,313 | 6,785 | 38.8 | 4.4 |

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| Year | Female <br> spawning <br> biomass <br> (kt) | Relative <br> spawning <br> biomass <br> $(\%)$ | Total <br> biomass <br> $(\mathbf{k t})$ | Age-2+ <br> biomass <br> $(\mathbf{k t})$ | Age-0 <br> recruits <br> (millions) | Relative <br> fishing <br> intensity <br> $(\%)$ | Exploitation <br> fraction <br> $(\%)$ |
| :---: | :---: | :---: | :---: | :---: | ---: | ---: | ---: |
| 1988 | 2,375 | 124.8 | 5,403 | 4,858 | 2,144 | 40.1 | 5.1 |
| 1989 | 2,111 | 110.8 | 5,031 | 4,876 | 114 | 47.4 | 6.1 |
| 1990 | 2,056 | 108.0 | 4,666 | 4,570 | 4,395 | 39.8 | 5.7 |
| 1991 | 1,864 | 98.0 | 4,206 | 3,853 | 1,261 | 58.7 | 8.3 |
| 1992 | 1,687 | 88.6 | 3,944 | 3,838 | 127 | 56.7 | 7.8 |
| 1993 | 1,314 | 68.9 | 2,961 | 2,895 | 3,230 | 46.4 | 6.9 |
| 1994 | 1,267 | 66.5 | 2,926 | 2,617 | 3,331 | 59.8 | 13.8 |
| 1995 | 1,147 | 60.2 | 2,946 | 2,634 | 1,271 | 50.8 | 9.5 |
| 1996 | 1,087 | 57.1 | 2,783 | 2,654 | 1,864 | 63.7 | 11.5 |
| 1997 | 1,081 | 56.8 | 2,596 | 2,438 | 1,027 | 68.4 | 13.3 |
| 1998 | 912 | 48.0 | 2,175 | 2,059 | 2,013 | 81.5 | 15.6 |
| 1999 | 783 | 41.1 | 2,097 | 1,730 | 12,898 | 92.2 | 18.0 |
| 2000 | 886 | 46.5 | 3,104 | 1,993 | 315 | 66.1 | 11.5 |
| 2001 | 1,284 | 67.4 | 4,236 | 4,170 | 1,256 | 67.6 | 5.5 |
| 2002 | 1,896 | 99.8 | 4,574 | 4,448 | 44 | 47.7 | 4.1 |
| 2003 | 1,754 | 92.4 | 3,833 | 3,786 | 1,715 | 43.7 | 5.4 |
| 2004 | 1,454 | 76.5 | 3,127 | 2,981 | 43 | 70.9 | 11.5 |
| 2005 | 1,145 | 60.3 | 2,553 | 2,496 | 2,857 | 68.3 | 14.5 |
| 2006 | 926 | 48.8 | 2,159 | 1,923 | 2,076 | 85.1 | 18.8 |
| 2007 | 705 | 37.1 | 1,787 | 1,666 | 25 | 86.9 | 17.5 |
| 2008 | 736 | 38.9 | 1,858 | 1,762 | 5,629 | 91.5 | 18.3 |
| 2009 | 616 | 32.6 | 1,686 | 1,316 | 1,371 | 79.7 | 13.6 |
| 2010 | 746 | 39.4 | 2,224 | 1,841 | 15,979 | 87.6 | 12.4 |
| 2011 | 729 | 38.5 | 2,849 | 1,803 | 384 | 87.0 | 15.9 |
| 2012 | 890 | 47.1 | 3,744 | 3,684 | 1,575 | 71.2 | 5.6 |
| 2013 | 1,662 | 88.0 | 4,277 | 4,138 | 371 | 66.8 | 6.9 |
| 2014 | 1,977 | 104.9 | 4,415 | 4,189 | 8,256 | 62.3 | 7.1 |
| 2015 | 1,448 | 76.9 | 3,487 | 2,908 | 34 | 45.5 | 6.7 |
| 2016 | 1,223 | 65.0 | 3,708 | 3,582 | 5,638 | 73.6 | 9.3 |
| 2017 | 1,646 | 87.5 | 3,998 | 3,452 | 1,565 | 79.3 | 12.8 |
| 2018 | 1,711 | 90.9 | 4,164 | 3,987 | 397 | 72.2 | 10.4 |
| 2019 | 1,402 | 74.5 | 3,417 | 3,378 | 273 | 80.3 | 12.2 |
| 2020 | 1,350 | 71.5 | 2,940 | 2,788 | 4,748 | 62.5 | 13.6 |
| 2021 | 1,118 | 58.9 | 2,939 | 2,251 | 10,187 | 62.4 | 14.5 |
| 2022 | 1,116 | 58.6 | 3,661 | 2,786 | 1,881 | 61.8 | 11.6 |
| 2023 | 1,335 | 69.9 | 4,246 | 4,009 | 979 | 55.1 | 6.6 |
| 2024 | 1,885 | 98.7 | 4,758 | 4,460 | 980 | - | - |
|  |  |  |  |  |  |  |  |

Table 24. Time-series of $95 \%$ posterior credibility intervals for the quantities shown in Table 23. In the last row, dashes (-) indicate quantities requiring 2024 catch which has not taken place yet.

| Year | Female <br> spawning <br> biomass <br> $(\mathbf{k t})$ | Relative <br> spawning <br> biomass | Total <br> biomass <br> $(\mathbf{k t})$ | Age-2+ <br> biomass <br> $(\mathbf{k t})$ | Age-0 <br> recruits <br> (millions) | Relative <br> fishing <br> intensity | Exploitation <br> fraction |
| :---: | :---: | :---: | :---: | :---: | ---: | ---: | ---: |
| 1966 | $576-1,816$ | $29.2-89.5 \%$ | $1,501-4,529$ | $1,284-4,196$ | $57-10,657$ | $25.2-71.9 \%$ | $3.3-10.7 \%$ |
| 1967 | $587-1,833$ | $29.7-89.7 \%$ | $1,589-4,908$ | $1,354-4,337$ | $214-14,910$ | $36.9-91.2 \%$ | $4.9-15.8 \%$ |
| 1968 | $590-1,921$ | $29.9-91.8 \%$ | $1,672-5,380$ | $1,338-4,588$ | $209-10,277$ | $23.6-71.7 \%$ | $2.7-9.1 \%$ |
| 1969 | $685-2,191$ | $34.0-104.8 \%$ | $1,878-6,113$ | $1,703-5,558$ | $39-4,067$ | $30.2-83.1 \%$ | $3.2-10.6 \%$ |
| 1970 | $773-2,529$ | $38.7-122.3 \%$ | $2,018-6,698$ | $1,875-6,065$ | $4,722-22,796$ | $33.5-90.2 \%$ | $3.9-12.5 \%$ |
| 1971 | $785-2,623$ | $39.5-126.6 \%$ | $2,110-7,269$ | $1,698-5,683$ | $83-3,161$ | $22.6-72.9 \%$ | $2.7-9.1 \%$ |
| 1972 | $879-2,960$ | $44.3-143.1 \%$ | $2,378-8,208$ | $2,325-8,053$ | $67-1,969$ | $16.1-59.2 \%$ | $1.5-5.1 \%$ |
| 1973 | $1,054-3,509$ | $52.6-170.3 \%$ | $2,478-8,371$ | $2,390-8,002$ | $3,235-14,699$ | $18.2-63.3 \%$ | $2.0-6.8 \%$ |
| 1974 | $1,036-3,412$ | $51.5-164.9 \%$ | $2,417-8,132$ | $2,151-7,092$ | $37-1,425$ | $22.1-71.7 \%$ | $3.0-9.8 \%$ |
| 1975 | $1,261-4,207$ | $63.1-204.6 \%$ | $2,994-10,097$ | $2,956-9,952$ | $934-4,575$ | $20.0-67.0 \%$ | $2.2-7.5 \%$ |
| 1976 | $1,436-4,754$ | $71.8-231.9 \%$ | $3,183-10,618$ | $3,084-10,258$ | $24-920$ | $19.6-67.8 \%$ | $2.3-7.7 \%$ |
| 1977 | $1,286-4,231$ | $64.5-206.6 \%$ | $2,864-9,463$ | $2,748-9,060$ | $3,706-14,871$ | $12.8-49.2 \%$ | $1.5-4.8 \%$ |

Continued on next page ...

| Year | Female spawning biomass (kt) | Relative spawning biomass | Total biomass (kt) | Age-2+ biomass (kt) | $\begin{aligned} & \text { Age-0 } \\ & \text { recruits } \\ & \text { (millions) } \end{aligned}$ | Relative fishing intensity | Exploitation fraction |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1978 | 1,089-3,449 | 54.6-169.9\% | 2,585-8,235 | 2,224-7,057 | 18-718 | 12.5-47.2\% | 1.5-4.7\% |
| 1979 | 1,196-3,640 | 59.8-181.3\% | 2,966-9,108 | 2,940-9,023 | 547-3,485 | 14.5-51.3\% | 1.5-4.7\% |
| 1980 | 1,074-3,153 | 53.1-157.1\% | 2,601-7,643 | 2,330-6,878 | 10,315-34,580 | 12.1-43.3\% | 1.3-3.9\% |
| 1981 | 1,053-2,952 | 51.5-149.0\% | 2,991-8,310 | 2,177-6,071 | 32-1,121 | 18.7-58.8\% | 2.3-6.4\% |
| 1982 | 1,151-3,057 | 55.7-155.7\% | 3,324-8,880 | 3,310-8,841 | 54-1,059 | 16.0-51.0\% | 1.2-3.3\% |
| 1983 | 1,510-3,816 | 72.6-196.9\% | 3,453-8,742 | 3,438-8,707 | 96-1,601 | 13.7-43.2\% | 1.3-3.3\% |
| 1984 | 1,593-3,805 | 75.3-199.2\% | 3,500-8,416 | 3,301-7,898 | 9,172-25,635 | 15.6-46.6\% | 1.8-4.2\% |
| 1985 | 1,630-3,696 | 76.1-197.1\% | 4,048-9,253 | 3,268-7,415 | 19-575 | 12.4-37.1\% | 1.5-3.4\% |
| 1986 | 1,671-3,583 | 77.4-192.8\% | 4,385-9,491 | 4,379-9,472 | 25-692 | 19.7-51.5\% | 2.2-4.8\% |
| 1987 | 1,758-3,609 | 80.6-196.5\% | 4,004-8,270 | 3,904-8,045 | 4,345-11,598 | 23.2-56.7\% | 2.9-6.0\% |
| 1988 | 1,784-3,503 | 80.8-193.8\% | 4,036-8,004 | 3,651-7,174 | 1,128-3,995 | 24.4-58.1\% | 3.5-6.8\% |
| 1989 | 1,610-3,051 | 72.1-171.2\% | 3,827-7,287 | 3,693-7,062 | 17-439 | 29.9-66.1\% | 4.2-8.1\% |
| 1990 | 1,588-2,914 | 70.0-166.1\% | 3,609-6,633 | 3,538-6,480 | 2,871-7,189 | 24.8-56.4\% | 4.0-7.4\% |
| 1991 | 1,470-2,584 | 63.7-149.8\% | 3,306-5,891 | 3,041-5,349 | 598-2,398 | 37.5-89.4\% | 6.0-10.5\% |
| 1992 | 1,345-2,321 | 57.8-134.6\% | 3,125-5,461 | 3,040-5,311 | 16-524 | 36.3-88.6\% | 5.6-9.9\% |
| 1993 | 1,055-1,786 | 45.1-104.2\% | 2,367-4,045 | 2,319-3,933 | 2,200-5,169 | 28.8-74.7\% | 5.1-8.6\% |
| 1994 | 1,033-1,693 | 43.5-100.4\% | 2,366-3,955 | 2,129-3,486 | 2,273-5,264 | 40.7-82.2\% | 10.4-17.0\% |
| 1995 | 930-1,539 | 39.3-91.3\% | 2,367-4,002 | 2,125-3,558 | 770-2,150 | 34.3-69.0\% | 7.0-11.7\% |
| 1996 | 886-1,454 | 37.1-86.6\% | 2,255-3,754 | 2,153-3,569 | 1,199-3,064 | 44.9-86.5\% | 8.6-14.2\% |
| 1997 | 883-1,441 | 37.1-86.0\% | 2,111-3,489 | 1,991-3,264 | 565-1,845 | 49.5-88.4\% | 10.0-16.3\% |
| 1998 | 745-1,219 | 31.2-72.4\% | 1,764-2,931 | 1,678-2,764 | 1,276-3,417 | 61.9-99.3\% | 11.6-19.1\% |
| 1999 | 633-1,060 | 26.7-62.3\% | 1,663-2,909 | 1,394-2,353 | 8,949-20,678 | 71.2-110.1\% | 13.3-22.4\% |
| 2000 | 698-1,227 | 30.3-70.3\% | 2,387-4,460 | 1,564-2,766 | 95-714 | 47.3-84.2\% | 8.3-14.6\% |
| 2001 | 1,001-1,811 | 43.9-102.2\% | 3,274-6,056 | 3,227-5,961 | 849-2,062 | 47.9-86.5\% | 3.8-7.1\% |
| 2002 | 1,499-2,643 | 65.3-150.9\% | 3,610-6,406 | 3,513-6,208 | 13-127 | 31.7-65.1\% | 2.9-5.1\% |
| 2003 | 1,422-2,382 | 60.5-139.1\% | 3,102-5,244 | 3,067-5,161 | 1,173-2,804 | 28.5-60.0\% | 4.0-6.7\% |
| 2004 | 1,207-1,930 | 50.2-115.0\% | 2,583-4,183 | 2,473-3,958 | 12-144 | 48.4-96.2\% | 8.6-13.8\% |
| 2005 | 957-1,511 | 39.6-90.7\% | 2,114-3,424 | 2,073-3,329 | 1,987-4,786 | 47.4-91.2\% | 10.9-17.5\% |
| 2006 | 770-1,240 | 32.1-73.4\% | 1,775-2,951 | 1,597-2,578 | 1,422-3,416 | 60.3-117.1\% | 14.0-22.7\% |
| 2007 | 575-976 | 24.4-56.0\% | 1,445-2,527 | 1,352-2,335 | 6-95 | 61.2-117.8\% | 12.5-21.6\% |
| 2008 | 587-1,056 | 25.4-58.6\% | 1,474-2,698 | 1,405-2,543 | 3,966-9,328 | 67.4-114.0\% | 12.7-23.0\% |
| 2009 | 479-914 | 21.1-49.8\% | 1,303-2,520 | 1,026-1,940 | 770-2,706 | 55.0-103.3\% | 9.2-17.4\% |
| 2010 | 579-1,114 | 25.6-60.3\% | 1,690-3,415 | 1,427-2,755 | 10,711-28,567 | 61.0-117.2\% | 8.3-16.0\% |
| 2011 | 560-1,105 | 25.1-59.3\% | 2,124-4,501 | 1,390-2,737 | 150-898 | 59.5-116.0\% | 10.5-20.7\% |
| 2012 | 665-1,403 | 30.5-73.4\% | 2,761-6,026 | 2,715-5,929 | 1,006-2,973 | 46.0-99.0\% | 3.5-7.6\% |
| 2013 | 1,246-2,641 | 57.0-137.2\% | 3,197-6,839 | 3,099-6,585 | 133-881 | 42.9-88.2\% | 4.3-9.2\% |
| 2014 | 1,496-3,143 | 68.1-163.1\% | 3,322-7,081 | 3,172-6,686 | 5,667-14,926 | 39.3-85.8\% | 4.5-9.4\% |
| 2015 | 1,105-2,310 | 50.0-119.8\% | 2,632-5,615 | 2,218-4,629 | 9-122 | 26.4-66.6\% | 4.2-8.7\% |
| 2016 | 940-1,954 | 42.2-101.5\% | 2,805-6,043 | 2,721-5,814 | 3,715-10,991 | 47.7-98.3\% | 5.7-12.2\% |
| 2017 | 1,248-2,721 | 56.4-139.1\% | 2,990-6,728 | 2,618-5,704 | 849-3,553 | 50.4-113.5\% | 7.7-16.8\% |
| 2018 | 1,259-2,948 | 57.8-148.5\% | 3,023-7,297 | 2,917-6,920 | 112-1,287 | 44.1-103.0\% | 6.0-14.2\% |
| 2019 | 1,004-2,501 | 46.5-125.4\% | 2,405-6,183 | 2,387-6,115 | 47-1,015 | 49.7-107.8\% | 6.7-17.3\% |
| 2020 | 911-2,566 | 43.3-125.4\% | 1,954-5,628 | 1,874-5,317 | 2,063-12,728 | 36.3-87.3\% | 7.2-20.3\% |
| 2021 | 699-2,269 | 34.3-110.0\% | 1,770-6,247 | 1,406-4,561 | 4,085-29,499 | 35.3-87.7\% | 7.2-23.2\% |
| 2022 | 627-2,453 | 31.7-118.5\% | 1,954-8,518 | 1,517-6,238 | 289-8,859 | 33.0-91.4\% | 5.2-21.3\% |
| 2023 | 652-3,225 | 34.2-154.7\% | 2,056-10,368 | 1,915-10,002 | 43-20,272 | 26.7-87.2\% | 2.6-13.8\% |
| 2024 | 853-4,828 | 45.0-229.8\% | 2,162-11,986 | 2,022-11,288 | 47-20,193 | - | - |

Table 25. Select parameters, derived quantities, and reference point posterior median estimates for the (2024) base model compared to the previous assessment's (2023) base model. Dashes (-) in column for the previous assessment indicate quantities that were not available in that assessment

| Parameter, Quantity, or Reference point | Base <br> model | 2023 <br> Base <br> model |
| :---: | :---: | :---: |

## Parameters

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| Parameter, Quantity, or Reference point | Base model | $\begin{gathered} 2023 \\ \begin{array}{c} \text { Base } \\ \text { model } \end{array} \end{gathered}$ |
| :---: | :---: | :---: |
| Natural mortality ( $M$ ) | 0.235 | 0.233 |
| Unfished recruitment ( $R_{0}$, millions) | 2,600 | 2,547 |
| Steepness ( $h$ ) | 0.812 | 0.808 |
| Additional biomass index SD | 0.322 | 0.286 |
| Catchability: biomass index ( $q_{b}$ ) | 0.838 | 0.833 |
| Additional age-1 index SD | 0.381 | 0.375 |
| Catchability: age- 1 index ( $q_{1}$ ) | 0.490 | 0.398 |
| Dirichlet-multinomial fishery ( $\log \theta_{\text {fish }}$ ) | -0.663 | -0.629 |
| Dirichlet-multinomial survey ( $\left.\log \theta_{\text {surv }}\right)$ | 2.770 | 2.595 |
| Derived Quantities |  |  |
| 2014 recruitment (millions) | 8,256 | 9,165 |
| 2016 recruitment (millions) | 5,638 | 6,374 |
| 2020 recruitment (millions) | 4,748 | 11,409 |
| Unfished female spawning biomass ( $B_{0}, \mathrm{kt}$ ) | 1,919 | 1,815 |
| 2009 relative spawning biomass | 32.6\% | 34.8\% |
| 2023 relative spawning biomass | 69.9\% | 104\% |
| 2024 relative spawning biomass | 98.7\% | - |
| 2023 rel. fishing intensity: $(1-\mathrm{SPR}) /\left(1-\mathrm{SPR}_{40 \%}\right)$ | 55.1\% | - |
| Reference Points based on $F_{\text {SPR }=40 \%}$ |  |  |
| Female spawning biomass at $F_{\text {SPR }=40} \%\left(B_{\text {SPR }=40 \%}\right.$, , kt $)$ | 681 | 642 |
| SPR at $F_{\text {SPR }=40 \%}$ (kt) | 40.0\% | 40.0\% |
| Exploitation fraction corresponding to SPR | 19.1\% | 18.6\% |
| Yield at $B_{\text {SPR }=40 \%}(\mathrm{kt})$ | 317 | 309 |

Table 26. Summary of median and $95 \%$ credibility intervals of equilibrium conceptual reference points for the base assessment model. Equilibrium reference points were computed using 1975-2023 averages for mean weight-at-age and baseline selectivity-at-age (1966-1990; prior to time-varying deviations). Dashes (-) indicate values that are static at one value and do not have a credible interval associated with them.

| Quantity | 2.5\% | Median | 97.5\% |
| :---: | :---: | :---: | :---: |
| Unfished female spawning biomass ( $B_{0}, \mathrm{kt}$ ) | 1,235 | 1,919 | 3,132 |
| Unfished recruitment ( $R_{0}$, millions) | 1,394 | 2,600 | 5,383 |
| Reference points (equilibrium) based on $F_{S P R=40 \%}$ |  |  |  |
| Female spawning biomass at $F_{\text {SPR }=40 \%}\left(B_{\text {SPR }}=40 \%, \mathrm{k} \mathrm{t}\right)$ | 409 | 681 | 1,127 |
| SPR at $F_{\text {SPR }=40 \%}$ | - | 40\% | - |
| Exploitation fraction corresponding to $F_{\text {SPR }}=40 \%$ | 16.3\% | 19.1\% | 22.0\% |
| Yield associated with $F_{\text {SPR }=40 \%}(\mathrm{kt})$ | 180 | 317 | 594 |
| Reference points (equilibrium) based on $B_{40 \%}\left(40 \%\right.$ of $\left.B_{0}\right)$ |  |  |  |
| Female spawning biomass ( $B_{40 \%}$, kt) | 494 | 767 | 1,253 |
| SPR at $B_{40} \%$ | 40.7\% | 43.5\% | 50.8\% |
| Exploitation fraction resulting in $B_{40 \%}$ | 12.9\% | 16.8\% | 20.2\% |
| Yield at $B_{40 \%}$ (kt) | 177 | 309 | 580 |
| Reference points (equilibrium) based on estimated MSY |  |  |  |
| Female spawning biomass ( $B_{\text {MSY }}, \mathrm{kt}$ ) | 297 | 490 | 867 |
| SPR at MSY | 22.8\% | 29.6\% | 45.1\% |
| Exploitation fraction corresponding to SPR at MSY | 15.8\% | 27.0\% | 36.5\% |
| MSY (kt) | 188 | 336 | 639 |

Table 27. Forecast quantiles of Pacific Hake relative spawning biomass at the beginning of the year. Catch alternatives are defined by letters a-o and are a constant value across all forecasted years unless otherwise defined in the first column. Acronyms are defined in the glossary (Appendix C).

| Catch alternative |  |  | Biomass at start of year | Relative spawning biomass |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Catch year | Catch (t) |  | 5\% | 50\% | 95\% |
|  |  |  | Start of 2024 | 0.51 | 0.99 | 2.01 |
| a: | 2024 | 0 | Start of 2025 | 0.57 | 1.11 | 2.23 |
|  | 2025 | 0 | Start of 2026 | 0.59 | 1.13 | 2.35 |
|  | 2026 | 0 | Start of 2027 | 0.57 | 1.12 | 2.45 |
| b: | 2024 | 180,000 | Start of 2025 | 0.53 | 1.06 | 2.18 |
|  | 2025 | 180,000 | Start of 2026 | 0.50 | 1.04 | 2.26 |
|  | 2026 | 180,000 | Start of 2027 | 0.46 | 1.00 | 2.32 |
| c: | 2024 | 225,000 | Start of 2025 | 0.52 | 1.05 | 2.16 |
|  | 2025 | 225,000 | Start of 2026 | 0.48 | 1.02 | 2.23 |
|  | 2026 | 225,000 | Start of 2027 | 0.43 | 0.97 | 2.29 |
| d: | 2024 | 320,000 | Start of 2025 | 0.50 | 1.02 | 2.14 |
| 10\% reduction | 2025 | 288,000 | Start of 2026 | 0.45 | 0.98 | 2.20 |
| each year | 2026 | 259,200 | Start of 2027 | 0.39 | 0.93 | 2.24 |
| e: | 2024 | 264,000 | Start of 2025 | 0.51 | 1.04 | 2.15 |
| 2023 catch | 2025 | 264,000 | Start of 2026 | 0.47 | 1.00 | 2.21 |
|  | 2026 | 264,000 | Start of 2027 | 0.41 | 0.94 | 2.26 |
| f: | 2024 | 350,000 | Start of 2025 | 0.49 | 1.01 | 2.13 |
|  | 2025 | 350,000 | Start of 2026 | 0.42 | 0.96 | 2.17 |
|  | 2026 | 350,000 | Start of 2027 | 0.35 | 0.88 | 2.20 |
| g: | 2024 | 350,000 | Start of 2025 | 0.49 | 1.01 | 2.13 |
| $10 \%$ reduction | 2025 | 315,000 | Start of 2026 | 0.43 | 0.97 | 2.18 |
| each year | 2026 | 283,500 | Start of 2027 | 0.37 | 0.91 | 2.23 |
| h : | 2024 | 380,000 | Start of 2025 | 0.49 | 1.01 | 2.12 |
|  | 2025 | 380,000 | Start of 2026 | 0.41 | 0.94 | 2.16 |
|  | 2026 | 380,000 | Start of 2027 | 0.33 | 0.86 | 2.17 |
| i: | 2024 | 380,000 | Start of 2025 | 0.49 | 1.01 | 2.12 |
| 10\% reduction | 2025 | 342,000 | Start of 2026 | 0.42 | 0.95 | 2.17 |
| each year | 2026 | 307,800 | Start of 2027 | 0.36 | 0.89 | 2.21 |
| j: | 2024 | 430,000 | Start of 2025 | 0.47 | 0.99 | 2.11 |
|  | 2025 | 430,000 | Start of 2026 | 0.39 | 0.92 | 2.14 |
|  | 2026 | 430,000 | Start of 2027 | 0.30 | 0.83 | 2.13 |
| k: | 2024 | 545,000 | Start of 2025 | 0.45 | 0.96 | 2.08 |
| 2022 TAC | 2025 | 545,000 | Start of 2026 | 0.33 | 0.86 | 2.08 |
|  | 2026 | 545,000 | Start of 2027 | 0.22 | 0.75 | 2.05 |
| 1: | 2024 | 625,000 | Start of 2025 | 0.43 | 0.94 | 2.06 |
| 2023 TAC | 2025 | 625,000 | Start of 2026 | 0.30 | 0.83 | 2.03 |
|  | 2026 | 625,000 | Start of 2027 | 0.18 | 0.70 | 1.99 |
| m: | 2024 | 875,262 | Start of 2025 | 0.37 | 0.88 | 1.99 |
| Fishing intensity | 2025 | 861,614 | Start of 2026 | 0.22 | 0.71 | 1.91 |
| at 100\% | 2026 | 782,426 | Start of 2027 | 0.13 | 0.57 | 1.86 |
| n: | 2024 | 747,588 | Start of 2025 | 0.40 | 0.91 | 2.02 |
| Default HR | 2025 | 772,111 | Start of 2026 | 0.24 | 0.76 | 1.97 |
| $\left(F_{\text {SPR }=40 \%}-40: 10\right)$ | 2026 | 717,464 | Start of 2027 | 0.14 | 0.62 | 1.91 |
| o: | 2024 | 767,382 | Start of 2025 | 0.39 | 0.90 | 2.02 |
| Equal catch | 2025 | 767,382 | Start of 2026 | 0.24 | 0.76 | 1.96 |
| $\left(\mathrm{C}_{2024} \approx \mathrm{C}_{2025}\right)$ | 2026 | 712,782 | Start of 2027 | 0.14 | 0.62 | 1.91 |

Table 28. Forecast quantiles of Pacific Hake relative fishing intensity (1-SPR)/(1-SPR ${ }_{40 \%}$ ), expressed as a proportion. Catch alternatives are defined by letters a-o and are a constant value across all forecasted years unless otherwise defined in the first column. Acronyms are defined in the glossary (Appendix C).

| Catch alternative |  |  | Relative fishing intensity |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Catch year | Catch (t) | 5\% | 50\% | 95\% |
| a: | 2024 | 0 | 0.00 | 0.00 | 0.00 |
|  | 2025 | 0 | 0.00 | 0.00 | 0.00 |
|  | 2026 | 0 | 0.00 | 0.00 | 0.00 |
| b: | 2024 | 180,000 | 0.22 | 0.43 | 0.69 |
|  | 2025 | 180,000 | 0.18 | 0.37 | 0.63 |
|  | 2026 | 180,000 | 0.16 | 0.33 | 0.59 |
| c: | 2024 | 225,000 | 0.27 | 0.50 | 0.78 |
|  | 2025 | 225,000 | 0.22 | 0.44 | 0.72 |
|  | 2026 | 225,000 | 0.20 | 0.40 | 0.69 |
| d: <br> $10 \%$ reduction each year | 2024 | 320,000 | 0.35 | 0.62 | 0.91 |
|  | 2025 | 288,000 | 0.27 | 0.53 | 0.83 |
|  | 2026 | 259,200 | 0.23 | 0.46 | 0.78 |
| e: <br> 2023 catch | 2024 | 264,000 | 0.30 | 0.55 | 0.84 |
|  | 2025 | 264,000 | 0.25 | 0.49 | 0.79 |
|  | 2026 | 264,000 | 0.23 | 0.46 | 0.77 |
| f: | 2024 | 350,000 | 0.38 | 0.66 | 0.94 |
|  | 2025 | 350,000 | 0.32 | 0.60 | 0.92 |
|  | 2026 | 350,000 | 0.29 | 0.57 | 0.92 |
| g: $10 \%$ reduction each year | 2024 | 350,000 | 0.38 | 0.66 | 0.94 |
|  | 2025 | 315,000 | 0.30 | 0.56 | 0.88 |
|  | 2026 | 283,500 | 0.25 | 0.49 | 0.83 |
| h: | 2024 | 380,000 | 0.40 | 0.69 | 0.97 |
|  | 2025 | 380,000 | 0.34 | 0.63 | 0.96 |
|  | 2026 | 380,000 | 0.31 | 0.60 | 0.97 |
| i: $10 \%$ reduction each year | 2024 | 380,000 | 0.40 | 0.69 | 0.97 |
|  | 2025 | 342,000 | 0.32 | 0.59 | 0.92 |
|  | 2026 | 307,800 | 0.26 | 0.52 | 0.88 |
| j: |  | $430,000$ | 0.44 | 0.73 | 1.02 |
|  | 2025 | 430,000 | 0.38 | 0.68 | 1.02 |
|  | 2026 | 430,000 | 0.35 | 0.66 | 1.05 |
| $\begin{gathered} \text { k: } \\ 2022 \text { TAC } \end{gathered}$ | 2024 | 545,000 | 0.51 | 0.82 | 1.11 |
|  | 2025 | 545,000 | 0.45 | 0.78 | 1.13 |
|  | 2026 | 545,000 | 0.42 | 0.78 | 1.20 |
| $\begin{gathered} \text { 1: } \\ 2023 \text { TAC } \end{gathered}$ | 2024 | 625,000 | 0.56 | 0.87 | 1.16 |
|  | 2025 | 625,000 | 0.50 | 0.85 | 1.20 |
|  | 2026 | 625,000 | 0.47 | 0.85 | 1.26 |
| m: <br> Fishing intensity at $100 \%$ | 2024 | 875,262 | 0.68 | 1.00 | 1.27 |
|  | 2025 | 861,614 | 0.62 | 1.00 | 1.30 |
|  | 2026 | 782,426 | 0.57 | 1.00 | 1.31 |
| $\mathbf{n}:$Default HR$\left(F_{\text {SPR }=40 \%}-40: 10\right)$ | 2024 | 747,588 | 0.62 | 0.94 | 1.22 |
|  | 2025 | 772,111 | 0.58 | 0.94 | 1.28 |
|  | 2026 | 717,464 | 0.53 | 0.94 | 1.30 |
| o: | 2024 | 767,382 | 0.63 | 0.95 | 1.23 |
| Equal catch | 2025 | 767,382 | 0.58 | 0.94 | 1.28 |
| $\left(\mathrm{C}_{2024} \approx \mathrm{C}_{2025}\right)$ | 2026 | 712,782 | 0.53 | 0.94 | 1.30 |

Table 29. Probabilities related to spawning biomass, relative fishing intensity, and the 2025 default harvest policy catch for alternative 2024 catch options (catch options explained in Table 27).

|  | $\begin{aligned} & \text { Catch }(t) \text { ) } \\ & \text { in } 2020 \end{aligned}$ | $\begin{gathered} \mathrm{B}_{2025} \\ <\mathrm{B}_{2024} \end{gathered}$ | $\begin{aligned} & \mathrm{B}_{2025} \\ & <B_{40 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2025} \\ & <B_{25 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2025} \\ & <B_{10 \%} \end{aligned}$ | 2024 <br> Fishing intensity $>100 \%$ | $\begin{gathered} 2025 \\ \text { Default HR } \\ \text { catch } \\ >2024 \\ \text { catch } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| a: | 0 | 0.03 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| b: | 180,000 | 0.22 | 0.01 | 0.00 | 0.00 | 0.00 | 0.00 |
| c: | 225,000 | 0.29 | 0.01 | 0.00 | 0.00 | 0.00 | 0.00 |
| d: | 320,000 | 0.44 | 0.02 | 0.00 | 0.00 | 0.02 | 0.03 |
| e: | 264,000 | 0.36 | 0.01 | 0.00 | 0.00 | 0.00 | 0.01 |
| f: | 350,000 | 0.49 | 0.02 | 0.00 | 0.00 | 0.03 | 0.04 |
| g : | 350,000 | 0.49 | 0.02 | 0.00 | 0.00 | 0.03 | 0.04 |
| h : | 380,000 | 0.53 | 0.02 | 0.00 | 0.00 | 0.04 | 0.06 |
| i: | 380,000 | 0.53 | 0.02 | 0.00 | 0.00 | 0.04 | 0.06 |
| j: | 430,000 | 0.60 | 0.02 | 0.00 | 0.00 | 0.07 | 0.10 |
| k: | 545,000 | 0.71 | 0.03 | 0.00 | 0.00 | 0.16 | 0.23 |
| $1:$ | 625,000 | 0.76 | 0.04 | 0.00 | 0.00 | 0.24 | 0.33 |
| m: | 875,262 | 0.87 | 0.07 | 0.01 | 0.00 | 0.50 | 0.61 |
| n: | 747,588 | 0.83 | 0.05 | 0.01 | 0.00 | 0.37 | 0.48 |
| o: | 767,382 | 0.83 | 0.05 | 0.01 | 0.00 | 0.39 | 0.50 |

Table 30. Probabilities related to spawning biomass, relative fishing intensity, and the 2026 default harvest policy catch for alternative 2025 catch options, given the 2024 catch level shown in Table 29 (catch options explained in Table 27).

|  | $\begin{aligned} & \text { Catch }(t) \\ & \text { in } 2025 \end{aligned}$ | $\begin{gathered} \mathrm{B}_{2026} \\ <\mathrm{B}_{2025} \end{gathered}$ | $\begin{aligned} & \mathrm{B}_{2026} \\ & <B_{40 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2026} \\ & <B_{25 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2026} \\ & <B_{10 \%} \end{aligned}$ | 2025 <br> Fishing intensity $>100 \%$ | $\begin{gathered} 2026 \\ \text { Default HR } \\ \text { catch } \\ >2025 \\ \text { catch } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| a: | 0 | 0.59 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| b: | 180,000 | 0.70 | 0.02 | 0.00 | 0.00 | 0.00 | 0.00 |
| c: | 225,000 | 0.72 | 0.02 | 0.00 | 0.00 | 0.00 | 0.00 |
| d: | 288,000 | 0.75 | 0.03 | 0.00 | 0.00 | 0.01 | 0.01 |
| e: | 264,000 | 0.74 | 0.03 | 0.00 | 0.00 | 0.00 | 0.01 |
| f: | 350,000 | 0.77 | 0.04 | 0.00 | 0.00 | 0.02 | 0.04 |
| g: | 315,000 | 0.76 | 0.03 | 0.00 | 0.00 | 0.01 | 0.03 |
| h: | 380,000 | 0.78 | 0.05 | 0.01 | 0.00 | 0.03 | 0.06 |
| i: | 342,000 | 0.76 | 0.04 | 0.01 | 0.00 | 0.02 | 0.04 |
| j: | 430,000 | 0.79 | 0.06 | 0.01 | 0.00 | 0.06 | 0.10 |
| k: | 545,000 | 0.82 | 0.09 | 0.02 | 0.00 | 0.15 | 0.25 |
| 1 : | 625,000 | 0.84 | 0.11 | 0.03 | 0.00 | 0.23 | 0.35 |
| m : | 861,614 | 0.87 | 0.20 | 0.07 | 0.00 | 0.50 | 0.64 |
| n: | 772,111 | 0.86 | 0.16 | 0.06 | 0.00 | 0.39 | 0.54 |
| 0: | 767,382 | 0.86 | 0.16 | 0.06 | 0.00 | 0.39 | 0.54 |

Table 31. Probabilities related to spawning biomass, relative fishing intensity, and the 2027 default harvest policy catch for alternative 2026 catch options, given the 2024 and 2025 catch levels shown in Tables 29 and 30 (catch options explained in Table 27).

|  | $\begin{gathered} \text { Catch }(t) \\ \text { in } 2026 \end{gathered}$ | $\begin{gathered} \mathrm{B}_{2027} \\ <\mathrm{B}_{2026} \end{gathered}$ | $\begin{aligned} & \mathrm{B}_{2027} \\ & <B_{40 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2027} \\ & <B_{25 \%} \end{aligned}$ | $\begin{aligned} & \mathrm{B}_{2027} \\ & <B_{10 \%} \end{aligned}$ | 2026 Fishing intensity $>100 \%$ | $\begin{gathered} 2027 \\ \text { Default HR } \\ \text { catch } \\ >2026 \\ \text { catch } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| a: | 0 | 0.66 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| b: | 180,000 | 0.73 | 0.03 | 0.00 | 0.00 | 0.00 | 0.00 |
| c: | 225,000 | 0.74 | 0.04 | 0.00 | 0.00 | 0.00 | 0.00 |
| d: | 259,200 | 0.75 | 0.05 | 0.01 | 0.00 | 0.01 | 0.01 |
| e: | 264,000 | 0.75 | 0.05 | 0.01 | 0.00 | 0.01 | 0.01 |
| f: | 350,000 | 0.78 | 0.07 | 0.02 | 0.00 | 0.03 | 0.05 |
| g: | 283,500 | 0.75 | 0.06 | 0.01 | 0.00 | 0.01 | 0.02 |
| h : | 380,000 | 0.78 | 0.09 | 0.02 | 0.00 | 0.04 | 0.07 |
| i: | 307,800 | 0.76 | 0.07 | 0.01 | 0.00 | 0.02 | 0.03 |
| j: | 430,000 | 0.79 | 0.11 | 0.03 | 0.00 | 0.07 | 0.12 |
| k: | 545,000 | 0.82 | 0.17 | 0.06 | 0.01 | 0.18 | 0.29 |
| 1 : | 625,000 | 0.83 | 0.21 | 0.09 | 0.01 | 0.27 | 0.41 |
| m: | 782,426 | 0.84 | 0.35 | 0.19 | 0.03 | 0.50 | 0.65 |
| n : | 717,464 | 0.84 | 0.29 | 0.14 | 0.02 | 0.41 | 0.56 |
| o: | 712,782 | 0.84 | 0.29 | 0.15 | 0.02 | 0.41 | 0.56 |

Table 32. Posterior medians for select parameters, derived quantities, reference points, and negative log likelihoods for the base model and some sensitivity runs (described in Section 3.8). A dash (-) indicates that the parameter or derived quantity was not estimated in the model.

| Parameter, Quantity, or Reference point | Base model | Steepness Mean Prior Low (0.5) | $\begin{gathered} \text { Steepness } \\ \text { Fix } \\ 1.0 \end{gathered}$ | $\begin{gathered} \text { Sigma } \\ \mathbf{R} \\ 1.0 \end{gathered}$ | $\begin{gathered} \text { Sigma } \\ \mathbf{R} \\ 1.6 \end{gathered}$ | Natural Mortality (SD=0.2) | Natural Mortality (SD=0.3) | Natural Mortality (Hamel Cope prior) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Parameters |  |  |  |  |  |  |  |  |
| Natural mortality ( $M$ ) | 0.235 | 0.239 | 0.234 | 0.231 | 0.237 | 0.290 | 0.312 | 0.315 |
| Unfished recruitment ( $R_{0}$, millions) | 2,600 | 2,772 | 2,535 | 1,870 | 3,201 | 5,477 | 7,873 | 8,476 |
| Steepness ( $h$ ) | 0.812 | 0.541 | - | 0.813 | 0.808 | 0.797 | 0.793 | 0.787 |
| Additional biomass index SD | 0.322 | 0.324 | 0.319 | 0.313 | 0.324 | 0.338 | 0.347 | 0.350 |
| Catchability: biomass index ( $q_{b}$ ) | 0.838 | 0.819 | 0.844 | 0.860 | 0.828 | 0.575 | 0.468 | 0.443 |
| Additional age-1 index SD | 0.381 | 0.380 | 0.376 | 0.394 | 0.383 | 0.354 | 0.346 | 0.342 |
| Catchability: age-1 index ( $q_{1}$ ) | 0.490 | 0.474 | 0.495 | 0.472 | 0.496 | 0.294 | 0.228 | 0.216 |
| Dirichlet-multinomial fishery ( $\left.\log \theta_{\text {fish }}\right)$ | -0.663 | -0.667 | -0.660 | -0.716 | -0.647 | -0.662 | -0.663 | -0.659 |
| Dirichlet-multinomial survey ( $\left.\log \theta_{\text {surv }}\right)$ | 2.770 | 2.783 | 2.795 | 2.736 | 2.758 | 2.775 | 2.782 | 2.782 |
| Derived Quantities |  |  |  |  |  |  |  |  |
| 2014 recruitment (millions) | 8,256 | 8,610 | 8,177 | 7,875 | 8,462 | 14,408 | 19,228 | 20,470 |
| 2016 recruitment (millions) | 5,638 | 5,875 | 5,578 | 5,363 | 5,784 | 9,934 | 13,212 | 14,096 |
| 2020 recruitment (millions) | 4,748 | 5,006 | 4,672 | 4,254 | 4,964 | 8,886 | 12,136 | 12,939 |
| Unfished female spawning biomass ( $B_{0}, \mathrm{kt}$ ) | 1,919 | 1,992 | 1,886 | 1,426 | 2,325 | 2,716 | 3,373 | 3,519 |
| 2009 relative spawning biomass | 32.6\% | 32.1\% | 32.9\% | 42.5\% | 27.2\% | 35.4\% | 36.1\% | 36.1\% |
| 2023 relative spawning biomass | 69.9\% | 70.7\% | 70.4\% | 86.8\% | 59.8\% | 80.9\% | 84.4\% | 85.3\% |
| 2024 relative spawning biomass | 98.7\% | 99.7\% | 99.3\% | 118.8\% | 85.3\% | 114.1\% | 119.0\% | 120.0\% |
| 2023 rel. fishing intensity: (1-SPR)/(1-SPR ${ }_{40 \%}$ ) | 55.1\% | 53.0\% | 55.7\% | 57.7\% | 53.9\% | 32.9\% | 25.0\% | 23.6\% |
| Reference Points based on $F_{\text {SPR }=40 \%}$ |  |  |  |  |  |  |  |  |
| Female spawning biomass at $F_{\text {SPR }=40 \%} \%\left(B_{\text {SPR }}=40 \%, \mathrm{kt}\right)$ | 681 | 450 | 754 | 510 | 826 | 948 | 1,172 | 1,195 |
| SPR at $F_{\text {SPR }=40 \%}(\mathrm{kt})$ | 40.0\% | 40.0\% | 40.0\% | 40.0\% | 40.0\% | 40.0\% | 40.0\% | 40.0\% |
| Exploitation fraction corresponding to SPR | 19.1\% | 19.3\% | 19.0\% | 18.8\% | 19.2\% | 22.6\% | 23.9\% | 24.1\% |
| Negative log likelihoods |  |  |  |  |  |  |  |  |
| Yield at $B_{\text {SPR }=40 \%}$ (kt) | 317 | 212 | 351 | 233 | 389 | 551 | 736 | 761 |
| Total | 2,225.16 | 2,226.82 | 2,234.82 | 2,228.62 | 2,227.44 | 2,225.16 | 2,225.15 | 2,225.19 |
| Survey index | -4.40 | -4.39 | -4.40 | -3.64 | -4.52 | -4.40 | -4.40 | -4.39 |
| Survey age compositions | 305.08 | 305.07 | 305.08 | 305.70 | 304.96 | 305.07 | 305.07 | 305.08 |
| Fishery age compositions | 1,840.11 | 1,840.15 | 1,840.08 | 1,846.84 | 1,838.33 | 1,840.10 | 1,840.09 | 1,840.12 |
| Recruitment | 63.85 | 65.02 | 63.53 | 58.22 | 68.38 | 63.87 | 63.89 | 63.83 |
| Parameter priors | 1.13 | 1.60 | 11.14 | 1.11 | 1.15 | 1.14 | 1.14 | 1.15 |
| Parameter deviations | 19.38 | 19.38 | 19.39 | 20.40 | 19.14 | 19.37 | 19.37 | 19.40 |

Table 33. Posterior medians for select parameters, derived quantities, reference points, and negative log likelihoods for the base model and further sensitivity runs (described in Section 3.8). A dash (-) indicates that the parameter or derived quantity was not estimated in the model.

| Parameter, Quantity, or Reference point | Base model | Remove Age Index | Downweight Fishery Comps |
| :---: | :---: | :---: | :---: |
| Parameters |  |  |  |
| Natural mortality ( $M$ ) | 0.235 | 0.233 | 0.236 |
| Unfished recruitment ( $R_{0}$, millions) | 2,600 | 2,384 | 2,748 |
| Steepness ( $h$ ) | 0.812 | 0.805 | 0.806 |
| Additional biomass index SD | 0.322 | 0.299 | 0.316 |
| Catchability: biomass index ( $q_{b}$ ) | 0.838 | 0.889 | 0.875 |
| Additional age-1 index SD | 0.381 | - | 0.351 |
| Catchability: age-1 index ( $q_{1}$ ) | 0.490 | - | 0.474 |
| Dirichlet-multinomial fishery $\left(\log \theta_{\text {fish }}\right)$ | -0.663 | -0.658 | - |
| Dirichlet-multinomial survey ( $\log \theta_{\text {surv }}$ ) | 2.770 | 2.787 | - |
| Derived Quantities |  |  |  |
| 2014 recruitment (millions) | 8,256 | 7,611 | 8,117 |
| 2016 recruitment (millions) | 5,638 | 5,053 | 5,613 |
| 2020 recruitment (millions) | 4,748 | 3,659 | 4,451 |
| Unfished female spawning biomass ( $B_{0}, \mathrm{kt}$ ) | 1,919 | 1,801 | 2,021 |
| 2009 relative spawning biomass | 32.6\% | 33.6\% | 29.3\% |
| 2023 relative spawning biomass | 69.9\% | 57.8\% | 61.8\% |
| 2024 relative spawning biomass | 98.7\% | 78.4\% | 84.9\% |
| 2023 rel. fishing intensity: $(1-\mathrm{SPR}) /\left(1-\mathrm{SPR}_{40 \%}\right)$ | 55.1\% | 65.1\% | 59.5\% |
| Reference Points based on $F_{\text {SPR }=40 \%}$ |  |  |  |
| Female spawning biomass at $F_{\text {SPR }=40 \%}\left(B_{\text {SPR }}=40 \%\right.$, kt$)$ | 681 | 635 | 713 |
| SPR at $F_{\text {SPR }=40 \%}$ (kt) | 40.0\% | 40.0\% | 40.0\% |
| Exploitation fraction corresponding to SPR | 19.1\% | 18.9\% | 19.2\% |
| Negative log likelihoods |  |  |  |
| Yield at $B_{\text {SPR }}=40 \%$ (kt) | 317 | 294 | 335 |
| Total | 2,225.16 | 2,219.13 | 216.94 |
| Survey index | -4.40 | -8.94 | -5.62 |
| Survey age compositions | 305.08 | 305.25 | 38.64 |
| Fishery age compositions | 1,840.11 | 1,839.04 | 117.06 |
| Recruitment | 63.85 | 63.42 | 56.25 |
| Parameter priors | 1.13 | 1.14 | -0.02 |
| Parameter deviations | 19.38 | 19.22 | 10.63 |

Table 34. Posterior medians for select parameters, derived quantities, reference points, and negative log likelihoods for the base model and further sensitivity runs (described in Section 3.8). A dash (-) indicates that the parameter or derived quantity was not estimated in the model.

| Parameter, Quantity, or Reference point | Base model | $\qquad$ | $\begin{gathered} \text { Phi } \\ \text { t.v. } \\ \text { selectivity } \\ (0.70) \end{gathered}$ | Phi t.v. selectivity (2.10) |
| :---: | :---: | :---: | :---: | :---: |
| Parameters |  |  |  |  |
| Natural mortality ( $M$ ) | 0.235 | 0.221 | 0.229 | 0.239 |
| Unfished recruitment ( $R_{0}$, millions) | 2,600 | 2,433 | 2,461 | 2,717 |
| Steepness ( $h$ ) | 0.812 | 0.810 | 0.807 | 0.807 |
| Additional biomass index SD | 0.322 | 0.369 | 0.329 | 0.316 |
| Catchability: biomass index ( $q_{b}$ ) | 0.838 | 0.857 | 0.859 | 0.823 |
| Additional age-1 index SD | 0.381 | 0.441 | 0.440 | 0.352 |
| Catchability: age-1 index ( $q_{1}$ ) | 0.490 | 0.495 | 0.508 | 0.477 |
| Dirichlet-multinomial fishery ( $\left.\log \theta_{\text {fish }}\right)$ | -0.663 | -0.976 | -0.724 | -0.640 |
| Dirichlet-multinomial survey ( $\log \theta_{\text {surv }}$ ) | 2.770 | 2.833 | 2.707 | 2.791 |
| Derived Quantities |  |  |  |  |
| 2014 recruitment (millions) | 8,256 | 8,126 | 8,012 | 8,484 |
| 2016 recruitment (millions) | 5,638 | 6,128 | 5,500 | 5,799 |
| 2020 recruitment (millions) | 4,748 | 6,605 | 5,288 | 4,470 |
| Unfished female spawning biomass ( $B_{0}, \mathrm{kt}$ ) | 1,919 | 2,009 | 1,928 | 1,952 |
| 2009 relative spawning biomass | 32.6\% | 28.2\% | 31.3\% | 32.7\% |
| 2023 relative spawning biomass | 69.9\% | 85.8\% | 72.2\% | 68.9\% |
| 2024 relative spawning biomass | 98.7\% | 120.7\% | 99.4\% | 98.9\% |
| 2023 rel. fishing intensity: $(1-$ SPR $) /\left(1-\right.$ SPR $\left._{40 \%}\right)$ | 55.1\% | 54.3\% | 58.2\% | 53.0\% |
| Reference Points based on $F_{\text {SPR }=40 \%}$ |  |  |  |  |
| Female spawning biomass at $F_{\text {SPR }=40 \%}\left(B_{\text {SPR }=40 \%}, \mathrm{kt}\right)$ | 681 | 714 | 670 | 689 |
| SPR at $F_{\text {SPR }=40 \%}(\mathrm{kt})$ | 40.0\% | 40.0\% | 40.0\% | 40.0\% |
| Exploitation fraction corresponding to SPR | 19.1\% | 18.1\% | 18.7\% | 19.3\% |
| Negative log likelihoods |  |  |  |  |
| Yield at $B_{\text {SPR }=40 \%}$ (kt) | 317 | 312 | 306 | 326 |
| Total | 2,225.16 | 2,356.78 | 2,259.11 | 2,210.95 |
| Survey index | -4.40 | -1.35 | -2.95 | -4.77 |
| Survey age compositions | 305.08 | 304.79 | 306.21 | 304.95 |
| Fishery age compositions | 1,840.11 | 1,940.60 | 1,859.39 | 1,831.04 |
| Recruitment | 63.85 | 63.44 | 64.69 | 63.06 |
| Parameter priors | 1.13 | 1.33 | 1.12 | 1.13 |
| Parameter deviations | 19.38 | 47.97 | 30.65 | 15.54 |

Table 35. Posterior medians from the base model for select parameters, derived quantities, reference point estimates, and negative log likelihoods for retrospective analyses. Some values are implied since they occur after the ending year of the respective retrospective analysis. A dash $(-)$ indicates that the parameter or derived quantity was not output by the model.

| Parameter, Quantity, or Reference point | Base model | year | years | years | years | years |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Parameters |  |  |  |  |  |  |
| Natural mortality ( $M$ ) | 0.235 | 0.233 | 0.231 | 0.231 | 0.231 | 0.229 |
| Unfished recruitment ( $R_{0}$, millions) | 2,600 | 2,520 | 2,477 | 2,434 | 2,448 | 2,384 |
| Steepness ( $h$ ) | 0.812 | 0.805 | 0.804 | 0.805 | 0.805 | 0.804 |
| Additional biomass index SD | 0.322 | 0.292 | 0.296 | 0.321 | 0.320 | 0.333 |
| Catchability: biomass index ( $q_{b}$ ) | 0.838 | 0.863 | 0.864 | 0.860 | 0.897 | 0.938 |
| Additional age-1 index SD | 0.381 | 0.343 | 0.338 | 0.292 | 0.267 | 0.281 |
| Catchability: age- 1 index ( $q_{1}$ ) | 0.490 | 0.413 | 0.440 | 0.408 | 0.407 | 0.400 |
| Dirichlet-multinomial fishery ( $\left.\log \theta_{\text {fish }}\right)$ | -0.663 | -0.645 | -0.595 | -0.599 | -0.573 | -0.554 |
| Dirichlet-multinomial survey ( $\log \theta_{\text {surv }}$ ) | 2.770 | 2.620 | 2.634 | 2.448 | 2.444 | 2.198 |
| Derived Quantities |  |  |  |  |  |  |
| 2014 recruitment (millions) | 8,256 | 8,761 | 8,838 | 9,727 | 10,098 | 9,922 |
| 2016 recruitment (millions) | 5,638 | 6,180 | 6,073 | 5,287 | 4,829 | 4,527 |
| 2020 recruitment (millions) | 4,748 | 11,230 | 6,001 | 930 | 950 | 946 |
| Unfished female spawning biomass ( $B_{0}, \mathrm{kt}$ ) | 1,919 | 1,901 | 1,886 | 1,859 | 1,870 | 1,838 |
| 2009 relative spawning biomass | 32.6\% | 33.2\% | 33.3\% | 34.4\% | 33.3\% | 32.9\% |
| 2023 relative spawning biomass | 69.9\% | 107.9\% | - | - | - | - |
| 2024 relative spawning biomass | 98.7\% | - | - | - | - | - |
| 2023 rel. fishing intensity: $(1-\mathrm{SPR}) /\left(1-\mathrm{SPR}_{40 \%}\right)$ | 55.1\% | - | - | - | - | - |
| Reference Points based on $F_{\text {SPR }=40 \%}$ |  |  |  |  |  |  |
| Female spawning biomass at $F_{\text {SPR }=40} \%\left(B_{\text {SPR }=40 \%,}\right.$, kt) | 681 | 669 | 663 | 658 | 660 | 651 |
| SPR at $F_{\text {SPR }=40 \%}$ (kt) | 40.0\% | 40.0\% | 40.0\% | 40.0\% | 40.0\% | 40.0\% |
| Exploitation fraction corresponding to SPR | 19.1\% | 18.9\% | 18.8\% | 18.8\% | 18.8\% | 18.7\% |
| Negative log likelihoods |  |  |  |  |  |  |
| Yield at $B_{\text {SPR }}=40 \%$ (kt) | 317 | 308 | 305 | 301 | 302 | 295 |
| Total | 2,225.16 | 2,156.46 | 2,098.34 | 2,035.58 | 1,987.54 | 1,921.37 |
| Survey index | -4.40 | -6.05 | -6.15 | -6.21 | -6.68 | -5.06 |
| Survey age compositions | 305.08 | 288.23 | 288.38 | 269.94 | 269.51 | 251.84 |
| Fishery age compositions | 1,840.11 | 1,793.52 | 1,736.78 | 1,694.15 | 1,648.11 | 1,599.10 |
| Recruitment | 63.85 | 62.96 | 62.45 | 61.20 | 60.97 | 60.28 |
| Parameter priors | 1.13 | 0.99 | 0.97 | 0.82 | 0.84 | 0.68 |
| Parameter deviations | 19.38 | 16.82 | 15.91 | 15.67 | 14.79 | 14.53 |

## 7 FIGURES



Figure 1. Overview map of the area in the Northeast Pacific Ocean occupied by Pacific Hake. Ports and areas of interest referred to in this document or past assessment documents are shown.


Figure 2. Spatial distribution of acoustic backscatter attributable to age-2 and older Pacific Hake from the Joint U.S. and Canadian Integrated Acoustic and Trawl Survey (1995-2023). Area of the circle is roughly proportional to observed backscatter. Bar plots show survey-estimated biomass for ages 2 to 20, with major cohorts highlighted in color. Figure produced by Julia Clemons (NOAA).


Figure 3. Spatial distribution of acoustic backscatter attributable to aggregations of age-1 Pacific Hake from the Joint U.S. and Canadian Integrated Acoustic and Trawl Survey 2003-2023 (spatial details are not available for survey years 1995, 1998, and 2001). Age-1 Pacific Hake are not fully sampled during the acoustic survey and were not explicitly considered during establishment of the survey sampling design. Additional backscatter from age-1 fish intermixed with older fish is not shown. Area of the circle is roughly proportional to observed backscatter. Figure produced by Julia Clemons (NOAA).


Figure 4. Total Pacific Hake catch used in the assessment by sector, 1966-2023. U.S. tribal catches are included in the sectors where they are represented.


Figure 5. Distribution of fishing depths (left) and bottom depths (right), in meters, of hauls targeting Pacific Hake in the U.S. Catcher-Processor and Mothership sectors from 2019-2023. Horizontal lines in each box represent the median depth and boxes encompass the middle $50 \%$ of the data. Whiskers encompass the $95 \%$ quantiles.


Figure 6. Distribution of fishing depths (left) and bottom depths (right), in meters, of hauls targeting Pacific Hake in the Canadian fleets from 2019-2023. Horizontal lines in each box represent the median depth and boxes encompass the middle $50 \%$ of the data. Whiskers encompass the $95 \%$ quantiles.


Figure 7. Overview of data used in this assessment. Circle areas are proportional to total catch for the fishery data, precision for the indices, and total sample size for the age compositions (and cannot be compared across data types). Additionally, mean weight-at-age data (1975-2023; not depicted here but see Figure 13 for sample sizes) are used to account for time-varying growth.


Proportion $0.0 \bigcirc 0.2 \bigcirc 0.4 \bigcirc 0.6$

Figure 8. Age compositions for the aggregate fishery (top, all sectors combined) and acoustic survey (bottom) for the years 1975-2023. Proportions in each year sum to 1.0 and area of the bubbles are proportional to the proportion and consistent in both panels (see key at top). The largest bubble in the fishery data is 0.72 for age 3 in 2011 and in the survey data is 0.75 for age 3 in 2013. Green lines track large cohorts.


Figure 9. Acoustic survey biomass index of age-2+ fish (Mt, Table 12). Approximate 95\% confidence intervals are based on sampling variability (intervals without the additional squid/hake apportionment uncertainty included in 2009, black line).


Figure 10. Relative index of age-1 fish (numbers of fish, Table 12) and approximate $95 \%$ confidence intervals based on sampling variability. The index is relative because the survey does not attempt to sample all available age- 1 fish and the analysis does not include kriging as is done to estimate age- $2+$ biomass.


Figure 11. Maturity ogives by year used in the assessment. The thick black line shows the equilibrium ogive which is an average of all years; the thick red line shows the forecast ogive which is an average of the last five years (2019-2023). The colors of the year lines move from orange in 2009 through the spectrum to dark blue in 2023.


Figure 12. Empirical weight-at-age (kg) values used for the base model as predicted from the timevarying model. Colors correspond to the values, with red being the lightest fish (across all years and ages) and blue being the heaviest fish. For each age, the most transparent cells indicate the lightest fish of that age. Data are only available from 1975-2023. Values based on assumptions for the pre-1975 and forecast years are shown outside the blue lines. Bold values between 1975-2023 represent unavailable data such that weights were predicted from the time-varying model. The bottom row (mean) is the mean weight-at-age over all years of data.

| 2028- | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2027 - | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2026 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2025 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2024 - | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| $2023-$ | 0 | 289 | 1,528 | 1,345 | 92 | 147 | 495 | 1,071 | 141 | 660 | 99 | 77 | 113 | 321 | 67 | 49 | 6,494 |
| $2022-$ | 0 | 17 | 740 | 61 | 103 | 574 | 1,389 | 138 | 923 | 145 | 100 | 104 | 428 | 123 | 42 | 38 | 4,925 |
| 2021 - | 0 | 770 | 132 | 177 | 743 | 1,875 | 157 | 1,375 | 132 | 149 | 135 | 666 | 118 | 42 | 13 | 25 | 6,509 |
| $2020-$ | 0 | 0 | 3 | 463 | 1,675 | 85 | 1,623 | 101 | 131 | 110 | 876 | 85 | 66 | 17 | 9 | 17 | 5,261 |
| 2019 | 1 | 363 | 965 | 1,916 | 118 | 2,765 | 178 | 304 | 189 | 1,566 | 153 | 121 | 54 | 37 | 29 | 27 | 8,786 |
| 2018 | 0 | 264 | 1,301 | 75 | 1,618 | 90 | 165 | 161 | 1,323 | 214 | 104 | 48 | 37 | 26 | 19 | 7 | 5,452 |
| 2017 - | 41 | 605 | 45 | 3,127 | 205 | 325 | 268 | 2,919 | 336 | 251 | 103 | 58 | 69 | 26 | 16 | 21 | 8,415 |
| 2016 | 0 | 12 | 2,375 | 106 | 274 | 132 | 1,826 | 117 | 144 | 40 | 20 | 22 | 16 | 3 | 5 | 10 | 5,102 |
| 2015 | 0 | 1,640 | 227 | 445 | 231 | 3,416 | 210 | 268 | 56 | 70 | 71 | 25 | 12 | 8 | 11 | 60 | 6,750 |
| 2014 | 0 | 0 | 136 | 173 | 2,879 | 282 | 499 | 52 | 102 | 61 | 27 | 10 | 2 | 4 | 9 | 22 | 4,258 |
| $2013-$ | 0 | 241 | 74 | 5,010 | 458 | 844 | 95 | 186 | 187 | 63 | 37 | 20 | 22 | 39 | 88 | 45 | 7,409 |
| 2012 | 0 | 39 | 2,654 | 864 | 2,052 | 205 | 329 | 224 | 127 | 58 | 36 | 38 | 63 | 112 | 55 | 28 | 6,884 |
| 2011 - | 0 | 688 | 564 | 3,915 | 180 | 383 | 300 | 62 | 53 | 28 | 32 | 39 | 132 | 39 | 16 | 9 | 6,440 |
| $2010-$ | 0 | 1 | 921 | 102 | 1,443 | 730 | 67 | 110 | 14 | 21 | 40 | 155 | 25 | 3 | 3 | 6 | 3,641 |
| 2009 | 0 | 488 | 33 | 2,002 | 2,265 | 160 | 538 | 55 | 118 | 114 | 770 | 129 | 41 | 27 | 20 | 15 | 6,775 |
| 2008 - | 0 | 44 | 330 | 1,374 | 124 | 756 | 58 | 180 | 169 | 1,200 | 90 | 29 | 31 | 11 | 11 | 9 | 4,416 |
| 2007 - | 9 | 432 | 1,175 | 240 | 972 | 103 | 448 | 279 | 2,894 | 291 | 139 | 112 | 60 | 37 | 18 | 14 | 7,223 |
| 2006 | 0 | 10 | 120 | 524 | 93 | 424 | 254 | 2,514 | 167 | 47 | 66 | 34 | 15 | 11 | 4 | 4 | 4,287 |
| $2005-$ | 0 | 6 | 329 | 49 | 420 | 317 | 3,739 | 394 | 128 | 166 | 106 | 74 | 50 | 13 | 6 | 12 | 5,809 |
| 2004 - | 0 | 0 | 1 | 230 | 270 | 2,492 | 235 | 59 | 134 | 75 | 31 | 33 | 12 | 8 | 4 | 5 | 3,589 |
| $2003-$ | 0 | 40 | 121 | 106 | 4,049 | 639 | 191 | 347 | 257 | 164 | 136 | 54 | 31 | 31 | 19 | 29 | 6,214 |
| $2002-$ | 0 | 0 | 1 | 1,786 | 598 | 374 | 248 | 170 | 240 | 138 | 37 | 35 | 40 | 11 | 17 | 33 | 3,728 |
| 2001 - | 0 | 6 | 1,245 | 987 | 905 | 1,111 | 650 | 710 | 408 | 124 | 138 | 101 | 69 | 57 | 34 | 73 | 6,618 |
| $2000-$ | 0 | 20 | 105 | 220 | 375 | 357 | 538 | 327 | 189 | 117 | 66 | 58 | 61 | 30 | 26 | 73 | 2,562 |
| 1999 | 0 | 0 | 204 | 618 | 481 | 430 | 252 | 70 | 95 | 110 | 19 | 38 | 59 | 11 | 14 | 58 | 2,459 |
| 1998 | 0 | 125 | 728 | 1,332 | 1,161 | 1,421 | 149 | 376 | 576 | 78 | 101 | 310 | 29 | 15 | 291 | 74 | 6,766 |
| $1997-$ | 0 | 0 | 28 | 833 | 799 | 21 | 174 | 304 | 20 | 82 | 139 | 3 | 6 | 143 | 4 | 72 | 2,628 |
| 1996 | 0 | 0 | 484 | 443 | 41 | 174 | 505 | 12 | 88 | 276 | 5 | 3 | 315 | 0 | 5 | 103 | 2,454 |
| 1995 | 0 | 162 | 425 | 77 | 152 | 1,128 | 74 | 321 | 889 | 46 | 8 | 1,066 | 17 | 36 | 1 | 339 | 4,741 |
| 1994 - | 0 | 0 | 1 | 94 | 789 | 24 | 481 | 624 | 18 | 9 | 825 | 2 | 12 | 2 | 221 | 27 | 3,129 |
| 1993 | 0 | 0 | 23 | 511 | 57 | 250 | 314 | 21 | 15 | 512 | 7 | 4 | 2 | 168 | 2 | 26 | 1,912 |
| 1992 | 0 | 153 | 454 | 164 | 635 | 1,040 | 119 | 54 | 1,761 | 65 | 21 | 16 | 788 | 27 | 2 | 76 | 5,375 |
| 1991 - | 0 | 0 | 35 | 408 | 529 | 75 | 22 | 692 | 38 | 5 | 8 | 337 | 10 | 0 | 59 | 10 | 2,228 |
| 1990 | 0 | 0 | 179 | 971 | 84 | 10 | 887 | 18 | 9 | 4 | 941 | 8 | 1 | 190 | 0 | 25 | 3,327 |
| 1989 | 49 | 0 | 168 | 154 | 89 | 2,420 | 109 | 89 | 61 | 1,639 | 104 | 26 | 188 | 14 | 6 | 27 | 5,143 |
| 1988 | 0 | 1 | 71 | 23 | 1,574 | 146 | 73 | 67 | 1,213 | 69 | 16 | 212 | 5 | 2 | 0 | 46 | 3,518 |
| 1987 | 0 | 0 | 32 | 1,180 | 427 | 236 | 129 | 1,722 | 114 | 54 | 263 | 10 | 2 | 7 | 36 | 7 | 4,219 |
| 1986 | 0 | 7 | 566 | 466 | 353 | 270 | 1,748 | 388 | 251 | 361 | 97 | 75 | 44 | 82 | 13 | 27 | 4,748 |
| 1985 | 0 | 27 | 37 | 94 | 302 | 1,876 | 225 | 165 | 203 | 42 | 17 | 15 | 11 | 4 | 1 | 3 | 3,022 |
| 1984 | 0 | 1 | 57 | 79 | 2,345 | 270 | 352 | 639 | 158 | 80 | 47 | 67 | 19 | 12 | 26 | 5 | 4,157 |
| 1983 | 0 | 109 | 22 | 1,295 | 176 | 453 | 1,179 | 307 | 238 | 203 | 302 | 98 | 78 | 63 | 19 | 13 | 4,555 |
| 1982 | 0 | 0 | 1,131 | 226 | 363 | 1,831 | 763 | 678 | 551 | 926 | 271 | 260 | 435 | 71 | 32 | 28 | 7,566 |
| 1981 - | 0 | 511 | 111 | 140 | 1,596 | 589 | 522 | 446 | 891 | 380 | 301 | 367 | 91 | 51 | 19 | 26 | 6,041 |
| 1980 | 0 | 6 | 34 | 1,214 | 412 | 651 | 678 | 1,093 | 550 | 599 | 537 | 169 | 127 | 85 | 40 | 53 | 6,248 |
| 1979 | 0 | 0 | 171 | 278 | 311 | 160 | 681 | 237 | 510 | 475 | 139 | 88 | 38 | 20 | 10 | 6 | 3,124 |
| 1978 | 0 | 68 | 71 | 419 | 403 | 1,507 | 252 | 550 | 1,449 | 444 | 303 | 228 | 114 | 63 | 27 | 15 | 5,913 |
| $1977-$ | 2 | 0 | 428 | 330 | 2,211 | 398 | 717 | 2,952 | 898 | 684 | 484 | 326 | 193 | 91 | 44 | 24 | 9,782 |
| 1976 | 0 | 7 | 90 | 673 | 234 | 101 | 939 | 508 | 419 | 407 | 314 | 196 | 140 | 68 | 31 | 11 | 4,138 |
| $1975-$ | 0 | 12 | 100 | 12 | 7 | 118 | 15 | 26 | 23 | 4 | 1 | 3 | 1 | 1 | 0 | 0 | 323 |
| 1974 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| $1973-$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| $1972-$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1971 - | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1970 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1969 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1968 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1967 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1966 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Sum - | 102 | 7,164 | 20,775 | 37,331 | 37,673 | 34,185 | 25,859 | 24,482 | 19,600 | 13,426 | 8,682 | 6,054 | 4,292 | 2,262 | 1,444 | 1,732 |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | 1 |
|  | 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | Sum |
|  | Age |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |

Figure 13. Sample sizes of empirical weight-at-age measurements used to fit the time-varying model used to estimate mean weight-at-age that is used in the base model. Colors and transparency are identical to Figure 12. Sample sizes of zero highlight years for which data are not available, i.e., pre 1975 and post 2023. The total sample sizes for each age used in the model over all data years are shown at the bottom and year-specific sample sizes are shown to the right using the same color scale with red indicating small sample sizes and blue indicating the large sample sizes.


Figure 14. Fecundity-at-age values used for the base model. Colors correspond to the values, with red being the least fecund fish (across all years and ages) and blue being the most fecund fish. For each age, the most transparent cells indicate the least fecund fish of that age. Fecundity is the product of maturity and weight-at-age. Weight-at-age data are only available from 1975-2023. Values based on assumptions for the pre-data and forecast years are shown outside the blue lines. Bold values between 1975-2023 represent year/age combinations where weight-at-age data were unavailable to fit the model such that weights were predicted rather than estimated.


Figure 15. Empirical mean weight-at-age ( kg ) values for ages $2-10$ used for the base model, as in Figure 12 but shown as time series. Purple lines are for the youngest ages and green lines are for the oldest ages shown, with age-5 having a thicker line and larger points as a visual aid.


Figure 16. Bridging models showing some of the sequential steps made towards the base model from the 2023 base model. Models include shifting to the newest version of Stock Synthesis, amending older data sources, and adding new data. Panels are spawning biomass (upper panel); relative spawning biomass (spawning biomass in each year relative to the unfished equilibrium spawning biomass, middle left); recruitment deviations (middle right); and age- $2+$ survey biomass (lower left) and age- 1 (lower right) indices, with triangles representing the observed survey indices. Note that in the top panel the red circle for 2024 is obscured by the orange circle.


Figure 17. Bridging models showing some of the sequential steps made towards the base model from the 2023 base model. Models include the last step from the previous figure of adding new data, changes to working up data inputs, and structural changes to the model. Panels are spawning biomass (upper panel); relative spawning biomass (spawning biomass in each year relative to the unfished equilibrium spawning biomass, middle left); recruitment deviations (middle right); and age- $2+$ survey biomass (lower left) and age- 1 (lower right) indices, with triangles representing the observed survey indices.


Figure 18. Fits (thin black lines) to the acoustic survey (points) with input $95 \%$ intervals around the observations. The thin black lines are the results of a random subset of individual Markov chain Monte Carlo (MCMC) samples. Thick, vertical black lines around observed survey points indicate $95 \%$ lognormal uncertainty intervals estimated by the kriging method and are used as input to the assessment model. Thin, vertical black lines indicate estimated $95 \%$ uncertainty intervals that account for the model estimate of additional uncertainty.


Figure 19. Assessment model fit to the relative age-1 index data that was produced from acoustic survey observations. Age-1 index observations (black dots) are input into the assessment model with uncertainty arising from sampling variability (thick vertical black lines). Additional uncertainty is estimated within the stock assessment model (thin vertical black lines) and added to the input sampling variability. A time series of the assessment model fit to the observations (blue line) represents the median of the posterior Markov chain Monte Carlo (MCMC) samples. A random subset of the individual MCMC time series samples are shown (thin blue lines) to provide context for the description of the median MCMC estimate.


Figure 20. Density of the Catchability (q) parameter for the acoustic survey index. The green vertical line is the median of the posterior for 2024, which is 0.838 ; the red line is the median of the posterior for the last assessment year, 2023, which was 0.833 .


Figure 21. Density of the Catchability (q) parameter for the age-1 index. The green vertical line is the median of the posterior for 2024, which is 0.490 ; the red line is the median of the posterior for the last assessment year, 2023, which was 0.398 .


## Age

Figure 22. Base model fits to the fishery age-composition data. Colored bars show observed proportions with colors following each cohort across years. Points with intervals indicate median expected proportions and $95 \%$ credibility intervals from the Markov chain Monte Carlo calculations.


Figure 23. Base model fits to the acoustic survey age-composition data. Colored bars show observed proportions with colors following each cohort across years. Points with intervals indicate median expected proportions and $95 \%$ credibility intervals from the Markov chain Monte Carlo calculations.


Figure 24. Pearson residuals for base model fits to the age-composition data for the medians of the Markov chain Monte Carlo posteriors for the fishery (top) and acoustic survey (bottom). Closed bubbles are positive residuals (observed > expected) and open bubbles are negative residuals (observed < expected). Green lines track cohorts from years of large recruitment events.


Figure 25. Prior (black lines) and posterior (blue histograms) distributions for natural mortality $(M)$, equilibrium $\log$ recruitment $\left(\ln R_{0}\right)$, steepness $(h)$, the additional process-error standard deviation (SD) for the acoustic survey and the age- 1 index, and the Dirichlet-multinomial parameters for the fishery $\left(\log \theta_{\text {fish }}\right)$ and the survey $\left(\log \theta_{\text {surv }}\right)$. Green triangles signify the initial value for each parameter. Red vertical lines represent the median of the posterior. The small downturns at the ends of the uniform priors for $\ln \left(R_{0}\right)$, and the Survey and age- 1 extra SD parameters represent the hard limits set for the priors in the Stock Synthesis control file.


Figure 26. As for Figure 25 but the x axis of each panel is truncated to the range of the posterior distribution, and thus, there is the potential for the full range of the prior and the initial value to be missing from individual panels.


Figure 27. Mountains plot of median fishery selectivity in each year for the base model. The range of selectivity is scaled to be between 0 and 1 in each year.


Figure 28. Fishery selectivity sampled from posterior probability distribution by year for the base model. Black dots and bars indicate the median and $95 \%$ credibility interval, respectively. The shaded polygon also shows the $95 \%$ credibility interval. The range of selectivity is scaled to be between 0 and 1 in each year. Selectivity for 1990 is shared for all years from 1966 to 1990.


Figure 29. Estimated selectivities for the acoustic survey age-2+ biomass index (top, with selectivity of zero for age- 1 fish) and fishery (bottom - shown for 2023 only, age- 1 and older) from a subsample of 1,000 draws from the posterior distribution for the base model.


Figure 30. Median (solid line) of the posterior distribution for beginning of the year female spawning biomass ( $B_{t}$ in year $t ; \mathrm{Mt}$ ) through 2024 (solid line) with $95 \%$ posterior credibility intervals (shaded area). The left-most circle with a $95 \%$ posterior credibility interval is the estimated unfished equilibrium biomass, $B_{0}$.


Figure 31. Median (solid line) of the posterior distribution for relative spawning biomass ( $B_{t} / B_{0}$ ) through 2024 with $95 \%$ posterior credibility intervals (shaded area). Dashed horizontal lines show $10 \%, 40 \%$, and $100 \%$ of the unfished equilibrium $\left(B_{0}\right)$.


Figure 32. Medians (solid circles) and means ( X ) of the posterior distribution for recruitment (billions of age-0 fish) with $95 \%$ posterior credibility intervals (vertical lines). The median of the posterior distribution for mean unfished equilibrium recruitment $\left(R_{0}\right)$ is shown as the horizontal dashed line with the $95 \%$ posterior credibility interval shaded between the dotted lines.


Figure 33. Medians (solid circles) of the posterior distribution for log-scale recruitment deviations with $95 \%$ posterior credibility intervals (vertical lines). Recruitment deviations for the years 1946-1965 are used to calculate the numbers at age in 1966, the initial year of the model.


Figure 34. Bubble plot of the medians of the posterior distributions of population numbers at age at the beginning of each year, where green diagonal lines follow each larger-than-usual year-class through time. The red line represents the mean age. The scale of the bubbles is represented in the key where the units are billions of fish; the largest overall bubble represents the 17.7 billion age-0 recruits in 1980. See Table 17 for values.


Figure 35. Medians (circles) of the posterior distribution of annual recruitment relative to recruitment in 2010 (recruitment divided by the 2010 recruitment for every MCMC sample), with $95 \%$ posterior credibility intervals (red lines). This procedure somewhat scales out the uncertainty due to uncertainty in mean unfished equilibrium recruitment ( $R_{0}$ ), and better elicits comparisons of relative cohort sizes; for example, recruitment in 2014 is clearly smaller than in 2010 (horizontal green dashed line). The year 2010 was chosen as the basis for comparison due to its well recognized size and the stability of cohort strength estimates over time. The median of $R_{0} / R_{2010}$ is shown as the horizontal dashed line with the $95 \%$ posterior credibility interval shaded between the dotted lines.


Figure 36. Estimated stock-recruitment relationship for the base model with median predicted recruitments and $95 \%$ posterior credibility intervals. Colors indicate time-period, with yellow colors in the early years and blue colors in the recent years. The thick solid black line indicates the central tendency (mean) and the red line indicates the central tendency after bias correcting for the lognormal distribution (median). Shading around the stock-recruitment relationship indicates uncertainty in shape associated with distribution of the steepness parameter $(h)$. The blue polygon on the right indicates the expected distribution of absolute recruitments.


Figure 37. Trend in median relative fishing intensity (relative to the $F_{\mathrm{SPR}=40 \%}$ management level) through 2023 with $95 \%$ posterior credibility intervals. The $F_{\mathrm{SPR}=40 \%}$ management level defined in the Joint U.S.-Canada Agreement for Pacific Hake is shown as a horizontal line at 1.0.


Figure 38. Trend in median exploitation fraction (catch divided by age-2+ biomass) through 2023 with $95 \%$ posterior credibility intervals.


Figure 39. Estimated historical path of median relative spawning biomass at the beginning of year $t$ and corresponding median relative fishing intensity in fishing year $t-1$ leading up to year $t$. Labels show the time series start and end years; labels correspond to year $t$ (i.e., year of the relative spawning biomass). Gray bars span the $95 \%$ credibility intervals for 2024 relative spawning biomass (horizontal) and 2023 relative fishing intensity (vertical).


Figure 40. The posterior distribution of the default 2024 catch limit calculated using the default harvest policy ( $F_{40 \%}-40: 10$ ). The median is $747,588 \mathrm{t}$ (vertical line), with the dark shaded area ranging from the $2.5 \%$ quantile to the $97.5 \%$ quantile, covering the range $298,355-2,124,832 \mathrm{t}$.


Figure 41. A comparison of female spawning stock biomass with fishing (black; as in Figure 30) and when the effects of fishing on the population are removed (red; unfished time series). Medians (solid lines) of the posterior distribution for beginning of the year female spawning biomass ( $B_{t}$ in year $t$; Mt) through 2024 (solid lines) with $95 \%$ posterior credibility intervals (shaded areas). The left-most circles with $95 \%$ posterior credibility intervals show the estimated unfished equilibrium biomass, $B_{0}$. The difference between the two lines shows the impact of removing fishing mortality from the population.


Figure 42. A comparison of relative spawning biomass when spawning biomass in year $t$ is related to unfished equilibrium biomass, $B_{0}$ (static $B_{0}$, black; as in Figure 31) and when spawning biomass in year $t$ is related to the unfished biomass time series in year $t$ (dynamic $B_{0}$, red). Median (solid lines) of the posterior distribution for each calculation of relative spawning biomass through 2024 with $95 \%$ posterior credibility intervals (shaded areas). Dashed horizontal lines show $10 \%, 40 \%$, and $100 \%$ of the unfished equilibrium $\left(B_{0}\right)$. The default $F_{40 \%}-40: 10$ harvest policy uses relative spawning biomass based on a static $B_{0}$ determination of stock status.


Figure 43. Median and $95 \%$ posterior credibility intervals of estimated relative spawning biomass to the start of 2024 from the base model and projections to the start of 2027 (vertical shaded rectangle) for several management actions, which are defined in the decision tables. The default harvest policy catches are $747,588 \mathrm{t}$ in 2024, $772,111 \mathrm{t}$ in 2025, and $717,464 \mathrm{t}$ in 2026.


Figure 44. Graphical representation of the probabilities related to spawning biomass, relative fishing intensity, and the 2025 default harvest policy catch for alternative 2024 catch options (explained in Table 27) as listed in Table 29. The symbols indicate points that were computed directly from model output and lines interpolate between the points.


Figure 45. Graphical representation of the probabilities related to spawning biomass, relative fishing intensity, and the 2026 default harvest policy catch for alternative 2025 catch options (including associated 2024 catch; catch options explained in Table 27) as listed in Table 30. The symbols indicate points that were computed directly from model output and lines interpolate between the points.


Figure 46. Graphical representation of the probabilities related to spawning biomass, relative fishing intensity, and the 2027 default harvest policy catch for alternative 2026 catch options (including associated 2024 and 2025 catches; catch options explained in Table 27) as listed in Table 31. The symbols indicate points that were computed directly from model output and lines interpolate between the points.


Figure 47. Forecast age compositions in numbers and in weight for the 2024 fishery catch (combined across all sectors in both countries). Light blue bars show median estimates. Thick black lines show $50 \%$ credibility intervals and thin black lines show $95 \%$ credibility intervals. These estimates are based on the posterior distribution for selectivity averaged across the most recent five years, weight-at-age data averaged across the most recent five years, and the distribution for expected numbers at age at the start of 2024 (see Table 17 for the Markov chain Monte Carlo medians of numbers-at-age for all years). The panel on the right is scaled based on the weight at each age averaged across the last five years.


Figure 48. Markov chain Monte Carlo estimates of spawning biomass for the base model and alternative sensitivity runs representing changing the mean of the prior for steepness from 1.0 to 0.5 , fixing steepness at 1.0 , lower (1.0) and higher (1.6) levels of variation assumed about the stock-recruitment relationship $\left(\sigma_{r}\right)$, changing the standard deviation of the prior for natural mortality, and using the Hamel/Cope prior distribution for natural mortality.


Figure 49. Markov chain Monte Carlo estimates of stock status (relative spawning biomass) for the base model and alternative sensitivity runs representing changing key parameters. See Figure 48 for sensitivity descriptions.


Figure 50. Markov chain Monte Carlo estimates of spawning biomass for the base model and alternative sensitivity models that represent the following changes in data: removing the index of age- 1 fish and down-weighting fishery composition data using the McAllister-Ianelli method.


Figure 51. Markov chain Monte Carlo estimates of stock status (relative spawning biomass) for the base model and alternative sensitivity models that represent changes in data. See Figure 50 for sensitivity descriptions.


Figure 52. Markov chain Monte Carlo estimates of recruitment deviations for the base model and alternative sensitivity runs that represent changes in data. See Figure 50 for sensitivity descriptions.


Figure 53. Markov chain Monte Carlo estimates of the fit to the acoustic survey biomass time series for the base model and alternative sensitivity runs that represent changes in data. See Figure 50 for sensitivity descriptions.


Figure 54. Markov chain Monte Carlo estimates of spawning biomass for the base model and alternative sensitivity runs representing different standard deviations ( $\Phi$ ) associated with timevarying selectivity. Standard deviations examined are below ( 0.21 and 0.70 ) and above (2.10) the base model value of 1.4.


Figure 55. Markov chain Monte Carlo estimates of stock status (relative spawning biomass) for the base model and alternative sensitivity runs representing different standard deviations ( $\Phi$ ) associated with time-varying selectivity. See Figure 54 for sensitivity descriptions.


Figure 56. Markov chain Monte Carlo estimates of recruitment for the base model and alternative sensitivity runs representing different standard deviations $(\Phi)$ associated with time-varying selectivity. See Figure 54 for sensitivity descriptions.


Figure 57. Markov chain Monte Carlo estimates of recruitment deviations for the base model and alternative sensitivity runs representing different standard deviations ( $\Phi$ ) associated with time-varying selectivity. See Figure 54 for sensitivity descriptions.


Figure 58. Markov chain Monte Carlo estimates of the fit to the survey index of age-2+ biomass for the base model and alternative sensitivity runs representing different standard deviations $(\Phi)$ associated with time-varying selectivity. See Figure 54 for sensitivity descriptions.


Figure 59. Retrospective analysis of recruitment deviations from Markov chain Monte Carlo models over the last 10 years. Recruitment deviations are the median log-scale differences between recruitment estimated by the model and expected recruitment from the stock-recruitment relationship (shading represents $95 \%$ posterior credibility intervals for a select low, moderate, and high deviation). Age-0 recruitment deviations are non-zero because Markov chain Monte Carlo allows for sampling from the full lognormal distribution. Lines represent estimated recruitment deviations for cohorts born from 2013 to 2023, with cohort birth year marked at the right of each color-coded line. For example, the right-most point for the 2016 cohort shows the cohort at age-8 (i.e., at the start of 2024, which represents the base model and includes data through 2023). The next point to the left is the 2016 cohort at age-7, calculated by removing one year of data (so includes data up to 2022). Thus, models are fit to data available only up to the start of the year in which each cohort became a given age, such that the last year of data for a given point equals cohort birth year + cohort age -1 .


Figure 60. As for Figure 59 but with the credibility intervals shown for the 2020 and 2021 cohorts.


Figure 61. Retrospective recruitment estimates shown in Figure 59 scaled relative to the most recent estimate of the strength of each cohort.


Figure 62. Markov chain Monte Carlo estimates of spawning biomass at the start of each year (top) and recruitment (bottom) for the base model and 5-year retrospective runs.


Figure 63. Summary of historical Pacific Hake assessment estimates of spawning biomass. Estimates are MLEs or Markov chain Monte Carlo medians depending on the model structure. Shading represents the $95 \%$ credible interval from the 2024 base model. Line colors are shades of orange for the oldest models, yellow shades for the 2000's, green shades for the 2010's and into blue shades from 2013 to present.


Figure 64. Comparison of absolute (left panel) and relative (right panel) variability associated with terminal year estimates of spawning biomass from Pacific Hake stock assessments dating back to 2012 (note: terminal year is the same as assessment year). The interquartile range specifies the width from quartile 1 ( $Q_{1}: 25$ th percentile) to quartile 3 ( $Q_{3}: 75$ th percentile) of terminal year spawning biomass from the posterior distribution and is a measure of absolute variability (similar to credible intervals). The quartile coefficient of dispersion is a relative measure of variability that can be compared across different data sets (similar to the coefficient of variation but less susceptible to outliers) and is calculated as $\left(Q_{3}-Q_{1}\right) /\left(Q_{1}+Q_{3}\right)$.


Figure 65. For each year $t, \mathrm{P}\left(B_{t+1}<B_{t}\right)$ is the probability that the spawning biomass at the start of $t+1$ is below that at the start of $t$. It is calculated in two ways. Red circles: the probability is taken from year $t$ 's stock assessment document, from the row in the decision table corresponding to the consequent catch in year $t$ (with interpolation if necessary). Blue triangles: the probability is calculated using the current 2024 base model. The grey horizontal line is the $50 \%$ value. For each year except 2017 and 2023, both probabilities lie on the same side of the grey line, indicating that each year's assessment model has almost always 'correctly' estimated an increase or decrease the subsequent year's biomass. For the 2024 assessment the probabilities are shown for all catch alternatives for 2024, as described in Table 27, with $0 t$ being the lowest probability, shown in pink.


Figure 66. For each year $t, \mathrm{P}\left(B_{t+1}<B_{40 \%}\right)$ is the probability that the spawning biomass at the start of $t+1$ is below $B_{40 \%}$. The red circles and blue triangles represent probabilities calculated analogously to Figure 65. The grey horizontal line is the $50 \%$ value. For each year except 2012, both probabilities lie on the same side of the grey line, indicating that each year's assessment model almost always correctly estimated that the subsequent year's biomass will not fall below $B_{40 \%}$. For the 2024 assessment the probabilities are shown for all catch alternatives for 2024, as described in Table 27, with 0 t shown in pink.

## A BASE MODEL MCMC DIAGNOSTICS



Figure A.1. Summary of Markov chain Monte Carlo diagnostics for natural mortality (upper 4 panels) and the natural log of mean unfished equilibrium recruitment $\left(\ln \left(R_{0}\right)\right.$; lower 4 panels) in the base model. Top sub-panels show the trace of the sampled values across iterations (absolute values, top left; cumulative running median with $2.5^{\text {th }}$ and $97.5^{\text {th }}$ percentiles, top right). The lower left sub-panel indicate the autocorrelation present in the chain at different lag times (i.e., distance between samples in the chain), and the lower right sub-panel shows the distribution of the values in the chain (i.e., the marginal density from a smoothed histogram of values in the trace plot).


Figure A.2. Summary of Markov chain Monte Carlo diagnostics for steepness in the base model. Sub-panel descriptions as in Figure A.1.


Figure A.3. Summary of Markov chain Monte Carlo diagnostics for the additional standard deviation (SD) in the biomass index (upper 4 panels) and for the age- 1 index (lower 4 panels) in the base model. Sub-panel descriptions as in Figure A.1.


Figure A.4. Summary of Markov chain Monte Carlo diagnostics for the Dirichlet-multinomial age-composition parameters for the fishery ( $\theta_{\text {fish }}$, upper 4 panels) and the survey $\left(\theta_{\text {surv }}\right.$, lower 4 panels) in the base model. Sub-panel descriptions as in Figure A.1.


Figure A.5. Summary histograms of Markov chain Monte Carlo diagnostics for all base model parameters. The level of autocorrelation in the chain (distribution across lag times, i.e., distance between samples in the chain, shown in the top left panel) influences the effective sample size (top right panel) used to estimate posterior distributions. The Geweke statistic (lower left panel) tests for equality between means located in the first part of the chain against means in the last part of the chain. The Heidelberger and Welch statistic (lower right panel) tests if the sampled values come from a stationary distribution by comparing different sections of the chain. Values for the unfished equilibrium recruitment $\left(R_{0}\right)$ are explicitly highlighted. Values inside the bars represent the number of parameters counted in that bin.


Figure A.6. Posterior correlations among the objective function which is minimized during model fitting, key base-model parameters, and derived quantities. Numbers refer to the absolute correlation coefficients, with font size proportional to the square root of the coefficient. Straight lines on the scatterplots are linear regressions.


Figure A.7. Posterior correlations among recruitment deviations from recent years and mean unfished equilibrium recruitment. Numbers refer to the absolute correlation coefficients, with font size proportional to the square root of the coefficient. Straight lines on the scatterplots are linear regressions.

## B SCIENTIFIC REVIEW GROUP (SRG) REQUESTS

This appendix summarizes results produced in response to Scientific Review Group requests made during the meeting held from February 6-9, 2024.

## B. 1 Day 1

## Request 1

List the sample sizes for males, females and unsexed (for the weight-at-age), for the U.S. and Canada. This will help to understand the effect of averaging unsexed fish into empirical weight-at-age and provide some insight into the differences spatially of weight-at-age.

## JTC Response

The JTC provided two tables, one for each country containing the weight-at-age sample sizes for each sex code $($ Male $=$ M, Female $=$ F, Unsexed $=\mathrm{U}$; Tables B. 1 and B.2).

The weight-at-age for two years, 2018 and 2023 are shown in Figure B.1. During the meeting, this plot contained unsexed fish as well, which were all much heavier than both males and females. On investigation, the JTC discovered that the data file for Canada had been modified to have all the sex data set to $\mathbf{U}$ (unsexed). This was fixed and Figures B. 2 and B. 3 show assessment model results.

Table B.1. Canadian sample sizes for weight-at-age data. $\mathrm{F}=\mathrm{Female}, \mathrm{M}=\mathrm{Male}$, and $\mathrm{U}=$ Unsexed.

| Year | F Fishery | F Survey | M Fishery | M Survey | U Survey | U Fishery |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: |
| 1982 | 133 | 0 | 67 | 0 | 0 | 0 |
| 1989 | 102 | 0 | 97 | 0 | 0 | 0 |
| 1990 | 133 | 0 | 91 | 0 | 0 | 0 |
| 1995 | 0 | 287 | 0 | 163 | 0 | 0 |
| 1998 | 0 | 645 | 0 | 555 | 1 | 0 |
| 2001 | 0 | 220 | 0 | 176 | 0 | 0 |
| 2002 | 294 | 0 | 179 | 0 | 0 | 0 |
| 2003 | 0 | 1,619 | 0 | 1,384 | 3 | 0 |
| 2004 | 161 | 0 | 124 | 0 | 0 | 0 |
| 2005 | 181 | 0 | 170 | 0 | 0 | 0 |
| 2009 | 0 | 291 | 0 | 275 | 0 | 0 |
| 2011 | 37 | 216 | 34 | 162 | 0 | 0 |
| 2012 | 5 | 322 | 2 | 253 | 0 | 0 |
| 2013 | 0 | 97 | 0 | 15 | 0 | 0 |
| 2015 | 0 | 155 | 0 | 203 | 0 | 0 |
| 2016 | 85 | 0 | 91 | 0 | 0 | 0 |
| 2017 | 323 | 148 | 331 | 110 | 42 | 0 |
| 2018 | 422 | 0 | 297 | 0 | 0 | 0 |
| 2019 | 214 | 259 | 162 | 132 | 0 | 0 |
| 2020 | 872 | 0 | 566 | 0 | 0 | 0 |
| 2021 | 0 | 94 | 0 | 55 | 0 | 0 |
| 2022 | 727 | 0 | 289 | 0 | 0 | 0 |
| 2023 | 657 | 172 | 229 | 90 | 0 | 0 |

Table B.2. U.S. sample sizes for weight-at-age data. $\mathrm{F}=$ Female, $\mathrm{M}=\mathrm{Male}$, and $\mathrm{U}=$ Unsexed. Samples from the Poland survey are included in the fisheries data.

| Year | F Fishery | F Survey | M Fishery | M Survey | U Survey | U Fishery |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1975 | 204 | 0 | 119 | 0 | 0 | 0 |
| 1976 | 2,362 | 0 | 1,720 | 0 | 0 | 56 |
| 1977 | 4,142 | 0 | 3,524 | 0 | 0 | 2,116 |
| 1978 | 3,164 | 0 | 2,668 | 0 | 0 | 81 |
| 1979 | 1,652 | 0 | 1,472 | 0 | 0 | 0 |
| 1980 | 3,011 | 0 | 3,233 | 0 | 0 | 4 |
| 1981 | 2,748 | 0 | 3,289 | 0 | 0 | 4 |
| 1982 | 3,596 | 0 | 3,770 | 0 | 0 | 0 |
| 1983 | 2,386 | 0 | 2,169 | 0 | 0 | 0 |
| 1984 | 2,165 | 0 | 1,982 | 0 | 0 | 10 |
| 1985 | 1,491 | 0 | 1,521 | 0 | 0 | 10 |
| 1986 | 2,398 | 38 | 2,232 | 15 | 0 | 65 |
| 1987 | 2,100 | 0 | 2,119 | 0 | 0 | 0 |
| 1988 | 1,677 | 0 | 1,819 | 0 | 0 | 22 |
| 1989 | 1,742 | 728 | 1,647 | 778 | 49 | 0 |
| 1990 | 1,849 | 0 | 1,254 | 0 | 0 | 0 |
| 1991 | 1,219 | 0 | 1,009 | 0 | 0 | 0 |
| 1992 | 1,675 | 976 | 1,516 | 1,208 | 0 | 0 |
| 1993 | 929 | 0 | 983 | 0 | 0 | 0 |
| 1994 | 1,487 | 0 | 1,642 | 0 | 0 | 0 |
| 1995 | 1,021 | 1,131 | 1,106 | 1,032 | 1 | 0 |
| 1996 | 1,250 | 0 | 1,204 | 0 | 0 | 0 |
| 1997 | 1,290 | 0 | 1,338 | 0 | 0 | 0 |
| 1998 | 1,502 | 1,245 | 1,646 | 1,172 | 0 | 0 |
| 1999 | 1,314 | 0 | 1,145 | 0 | 0 | 0 |
| 2000 | 1,336 | 0 | 1,226 | 0 | 0 | 0 |
| 2001 | 2,089 | 1,134 | 1,993 | 1,006 | 0 | 0 |
| 2002 | 1,839 | 0 | 1,416 | 0 | 0 | 0 |
| 2003 | 1,473 | 0 | 1,735 | 0 | 0 | 0 |
| 2004 | 1,501 | 0 | 1,803 | 0 | 0 | 0 |
| 2005 | 1,676 | 898 | 1,928 | 956 | 0 | 0 |
| 2006 | 1,924 | 0 | 2,363 | 0 | 0 | 0 |
| 2007 | 1,851 | 1,642 | 2,021 | 1,668 | 42 | 0 |
| 2008 | 2,129 | 0 | 2,277 | 0 | 0 | 10 |
| 2009 | 1,627 | 1,067 | 2,081 | 1,427 | 6 | 1 |
| 2010 | 1,518 | 0 | 2,110 | 0 | 0 | 13 |
| 2011 | 1,770 | 854 | 2,165 | 1,200 | 1 | 1 |
| 2012 | 1,522 | 1,373 | 1,852 | 1,553 | 1 | 1 |
| 2013 | 2,101 | 1,282 | 2,424 | 1,489 | 0 | 1 |
| 2014 | 1,718 | 0 | 2,540 | 0 | 0 | 0 |
| 2015 | 1,510 | 1,385 | 1,984 | 1,459 | 27 | 30 |
| 2016 | 2,281 | 0 | 2,645 | 0 | 0 | 2 |
| 2017 | 2,503 | 1,031 | 2,807 | 1,109 | 7 | 4 |
| 2018 | 2,435 | 0 | 2,295 | 0 | 0 | 4 |
| 2019 | 2,464 | 1,354 | 2,687 | 1,507 | 5 | 2 |
| 2020 | 1,826 | 0 | 1,995 | 0 | 0 | 2 |
| 2021 | 2,248 | 1,019 | 1,972 | 1,118 | 2 | 1 |
| 2022 | 2,095 | 0 | 1,809 | 0 | 0 | 5 |
| 2023 | 1,682 | 1,076 | 1,366 | 1,219 | 0 | 3 |

## Request 2



Figure B.1. Weight-at-age by sex (shapes) for two fishing years, 2018 and 2023 (colors).

On Figure G. 10 add the data for proportion mature at age-2 (if possible) and age- 3 to the model prediction of maturity at age-2 (if possible) and age-3 from the temperature-based model.

## JTC Response

The JTC provided two figures showing the proportion mature by year and temperature.

- Figure B. 4 shows empirical results for age two and Figure B. 5 shows empirical results for age three. For each year (denoted on the smooth line) the points representing the raw data from non-ASHOP samples appear above or below the labeled point from the same year.
- We do not expect the points to overlap or lie on the line because the curve represents the marginal effect of temperature (e.g., predicted maturity from non-ASHOP samples collected from an average sample location on July 1 but predicted to the 278th day of the year).
- Changing the reference date from July 1 to a later date for age- 3 fish shifted the curve up closer to the points because the non-ASHOP samples occur later in the year ( $25 \%$ before July $1 / 75 \%$ after). The points are also affected by variable sampling effort year to year.
- In conclusion, we would never really expect the empirical estimates to perfectly match the model-based ones. Sampling varies across space and time, and the empirical estimates do not account for any of this variation.


Figure B.2. Female spawning biomass for the pre-SRG base model compared to the final base model, where the latter model excluded unsexed fish in the weight-at-age data.


Figure B.3. Relative female spawning biomass for the pre-SRG base model compared to the final base model, where the latter model excluded unsexed fish in the weight-at-age data.

Age 3


Figure B.4. Proportion of age-2 fish mature by year and sea temperature (Celcius).


Figure B.5. Proportion of age-3 fish mature by year and sea temperature (Celcius).

## Request 3

Add $\mathrm{B}_{2023} / \mathrm{B}_{0}$ to Table 25 comparing the 2023 and 2024 assessments.

## JTC Response

The JTC provided an updated table, and updated the table in the assessment to match it. See Table 25. All other similar tables in the document containing parameter estimates now have this row as well.

## Request 4

The SRG requests data to better understand the presence of clean hake trawls and mixed species trawls in the 2023 acoustic survey by area (such as INPFC area). This will help understand where hake were in 2023 and the potential effects of mixed trawls on the determination of biomass.

## JTC Response

In 2023, the acoustic-trawl survey had 13 regions out of 282 that were classified as 'hake $\mathrm{mix}^{\prime}$. These regions were associated with 4 separate trawls. The hake percentages in those hake mix trawls ranged from $55 \%$ to $76 \%$ hake (all mixed with rockfish).

Of the un-Kriged biomass ( $3.1 \%$ ), 22.9 kt out of the 735 kt came from hake mixes in 2023.

## Request 5

The SRG requests maturity-at-age in a tabular format from 2008 to 2026. These numbers will help understand the variability of maturity-at-age without the effect of changes in weight-at-age used to calculate fecundity. It will also help understand how the projected maturity-at-age relates to estimated values.

## JTC Response

The actual years that are included are 2009-2023 and 4 projection years. The projections use the mean of the maturity-at-age for 2019-2023. Figure B. 6 shows the requested table in a heatmap format to match the other figures in the document depicting weight-at-age, sample sizes of weight-at-age, and fecundity.


Figure B.6. Heatmap showing maturity-at-age by year in the same format as the other heatmaps for weight-at-age, sample size, and fecundity found in this document (Figures 12-14). The Projected years (2024-2027) are set to the mean values of the previous 5 years (2019-2023). The years prior to the initial year shown (2009) are set to the mean of the entire time series and are shown in the Mean row.

## Request 6

The SRG requests that the JTC add a new run seriously downweighting the age composition sample sizes to increase the influence of the survey biomass index. For example, an effective sample size of no more than 50 , which would mean dividing the sample size by about 100. Alternatively, use a lambda of 0.05 on survey and fishery age composition data. Report the additional survey variance as well.

## JTC Response

A lambda value of 0.05 was multiplied by the age-composition likelihood component to calculate the overall negative $\log$ likelihood to be minimized. The bulleted list below provides parameter estimates for the base model and the requested model run, respectively. Figures B.7-B. 12 were provided to the SRG to fulfill the request thoroughly. While substantially downweighting age-composition data results in an improved survey fit, the model loses critical information on recruitment and thus overall population scale.

- DM fishery: 0.341 (base); 0.035 (lambda)
- DM survey: 0.942; 0.271
- DM ratio: 0.362; 0.129
- Extra SD survey: 0.313; 0.202
- Extra SD age-1 index: 0.385; 0.371
- Age Likelihood: 2,144; 131
- Approximate run time: 2 hours; 17 hours


Figure B.7. Fits to the acoustic age $2+$ survey observations (triangles) for the base model (black) and the adjusted lambda $=0.05$ model (red).


Figure B.8. Fits to the relative age 1 index observations (triangles) for the base model (black) and the adjusted lambda $=0.05$ model (red).


Figure B.9. Estimated female spawning biomass for the base model (black) and the adjusted lambda $=0.05$ model (red). The shaded areas represent the $95 \%$ credible interval.


Figure B.10. Estimated relative female spawning biomass for the base model (black) and the adjusted lambda $=0.05$ model (red). The shaded areas represent the $95 \%$ credible interval.


Figure B.11. Estimated recruitment deviations for the base model (black) and the adjusted lambda $=0.05$ model (red). The bars represent the $95 \%$ credible interval.


Figure B.12. Estimated absolute recruitment for the base model (black) and the adjusted lambda $=0.05$ model (red). The bars represent the $95 \%$ credible interval.

## Request 7

The SRG would like to see the proportion of survey biomass and catch estimated in Canada. JTC Response

The JTC provided two tables to show the catch-at-age in each country, which were calculated by multiplying the input proportions-at-age with the the annual catch in each country (Tables B. 3 and B.4).
To show the proportion of biomass in Canada as determined by the acoustic survey, a figure (Figure B.13) and two tables (Tables B. 6 and B.5) were provided.

Table B.3. Fishery catch-at-age for Canada ( t ).

| Year | Catch | $\mathbf{1}$ | $\mathbf{2}$ | $\mathbf{3}$ | $\mathbf{4}$ | $\mathbf{5}$ | $\mathbf{6}$ | $\mathbf{7}$ | $\mathbf{8}$ | $\mathbf{9}$ | $\mathbf{1 0}$ | $\mathbf{1 1}$ | $\mathbf{1 2}$ | $\mathbf{1 3}$ | $\mathbf{1 4}$ |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 2007 | 73,590 | 43 | 225 | 315 | 5,477 | 1,482 | 4,147 | 3,196 | 38,261 | 7,415 | 1,448 | 3,853 | 4,383 | 1,098 | 373 |
| 2008 | 71,010 | 0 | 501 | 2,759 | 653 | 4,874 | 885 | 2,917 | 3,385 | 41,070 | 5,284 | 3,978 | 916 | 1,432 | 153 |
| 2009 | 13,223 | 0 | 79 | 4,740 | 2,881 | 552 | 1,349 | 64 | 196 | 26 | 2,634 | 358 | 68 | 260 | 17 |
| 2010 | 48,934 | 0 | 250 | 921 | 7,957 | 23,635 | 3,540 | 2,332 | 567 | 329 | 966 | 5,408 | 1,951 | 670 | 281 |
| 2011 | 56,073 | 0 | 0 | 32,542 | 1,030 | 6,979 | 6,754 | 2,009 | 1,574 | 213 | 677 | 194 | 3,230 | 231 | 248 |
| 2012 | 47,059 | 0 | 262 | 4,200 | 21,222 | 2,376 | 8,960 | 3,760 | 1,071 | 1,014 | 397 | 576 | 241 | 2,078 | 285 |
| 2013 | 52,249 | 0 | 0 | 1,019 | 2,799 | 5,074 | 2,107 | 12,282 | 4,975 | 2,800 | 5,810 | 622 | 1,297 | 2,247 | 9,871 |
| 2014 | 35,118 | 0 | 0 | 276 | 6,103 | 4,393 | 9,220 | 2,807 | 4,282 | 2,849 | 769 | 1,026 | 195 | 492 | 343 |
| 2015 | 39,684 | 454 | 0 | 1,215 | 872 | 23,386 | 4,154 | 5,516 | 871 | 1,587 | 1,047 | 259 | 150 | 0 | 58 |
| 2016 | 69,743 | 177 | 3,210 | 551 | 3,403 | 4,551 | 44,086 | 4,433 | 4,974 | 1,228 | 997 | 57 | 668 | 464 | 95 |
| 2017 | 81,113 | 3,016 | 342 | 6,191 | 1,670 | 3,362 | 5,067 | 39,733 | 10,348 | 6,668 | 1,479 | 1,490 | 1,038 | 144 | 137 |
| 2018 | 92,689 | 295 | 4,576 | 1,271 | 18,810 | 1,657 | 3,934 | 5,172 | 37,048 | 10,282 | 4,453 | 2,323 | 1,019 | 759 | 858 |
| 2019 | 95,013 | 21 | 15,182 | 12,823 | 3,155 | 22,764 | 2,266 | 3,908 | 3,575 | 23,296 | 3,847 | 2,613 | 958 | 409 | 197 |
| 2020 | 30,085 | 0 | 12 | 2,885 | 5,957 | 412 | 9,073 | 814 | 1,049 | 771 | 7,240 | 861 | 636 | 67 | 144 |
| 2021 | 45,807 | 0 | 0 | 0 | 7,907 | 10,994 | 3,087 | 12,537 | 1,318 | 1,318 | 4,355 | 2,460 | 514 | 0 | 1,318 |
| 2022 | 31,671 | 0 | 0 | 5 | 123 | 2,257 | 7,044 | 2,485 | 5,614 | 2,283 | 1,384 | 1,866 | 5,133 | 2,298 | 892 |
| 2023 | 23,557 | 0 | 11 | 242 | 130 | 745 | 3,816 | 5,451 | 1,238 | 4,387 | 1,265 | 741 | 1,616 | 3,109 | 503 |

Table B.4. Fishery catch-at-age for U.S.A. (t).

| Year | Catch | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2008 | 248,395 | 2,349 | 26,390 | 88,573 | 6,174 | 39,451 | 2,671 | 9,195 | 7,945 | 57,070 | 4,244 | 1,601 | 1,340 | 463 | 533 | 396 |
| 2009 | 121,324 | 1,333 | 795 | 43,582 | 36,762 | 3,436 | 11,084 | 930 | 2,658 | 1,804 | 15,765 | 1,757 | 375 | 569 | 235 | 237 |
| 2010 | 170,960 | 45 | 50,951 | 6,367 | 66,392 | 32,785 | 2,365 | 3,531 | 347 | 673 | 1,260 | 5,016 | 828 | 94 | 126 | 180 |
| 2011 | 230,219 | 9,198 | 29,296 | 160,492 | 5,950 | 11,887 | 7,744 | 1,496 | 927 | 563 | 358 | 142 | 1,710 | 243 | 101 | 115 |
| 2012 | 159,696 | 463 | 87,120 | 16,632 | 41,442 | 3,103 | 4,407 | 2,536 | 1,371 | 594 | 178 | 320 | 428 | 465 | 304 | 333 |
| 2013 | 232,561 | 80 | 1,806 | 182,689 | 13,275 | 23,566 | 1,779 | 2,634 | 2,858 | 983 | 669 | 410 | 142 | 341 | 859 | 469 |
| 2014 | 263,945 | 0 | 10,532 | 10,893 | 187,047 | 15,985 | 25,963 | 2,720 | 4,844 | 2,934 | 1,357 | 343 | 77 | 125 | 353 | 772 |
| 2015 | 154,160 | 5,139 | 1,885 | 13,262 | 6,493 | 112,388 | 5,633 | 5,277 | 1,065 | 1,185 | 1,195 | 236 | 127 | 84 | 0 | 191 |
| 2016 | 261,582 | 973 | 152,082 | 4,763 | 12,658 | 5,582 | 71,932 | 4,087 | 5,685 | 1,602 | 684 | 759 | 524 | 90 | 111 | 51 |
| 2017 | 354,129 | 16,571 | 2,820 | 170,171 | 8,308 | 14,398 | 7,812 | 109,815 | 8,885 | 7,254 | 3,442 | 886 | 2,377 | 696 | 313 | 383 |
| 2018 | 318,306 | 33,394 | 94,236 | 3,995 | 91,906 | 4,574 | 7,674 | 7,332 | 58,134 | 7,687 | 3,790 | 1,922 | 1,393 | 1,089 | 976 | 202 |
| 2019 | 317,002 | 0 | 38,011 | 72,616 | 3,591 | 115,220 | 4,809 | 12,484 | 4,981 | 53,575 | 4,246 | 3,660 | 1,411 | 1,161 | 663 | 572 |
| 2020 | 287,813 | 0 | 161 | 25,494 | 115,322 | 5,563 | 85,014 | 4,438 | 5,549 | 5,841 | 34,093 | 2,671 | 2,555 | 789 | 74 | 248 |
| 2021 | 268,556 | 4,092 | 1,294 | 5,927 | 33,341 | 98,939 | 5,502 | 61,806 | 4,291 | 5,614 | 6,011 | 33,179 | 5,328 | 1,605 | 520 | 1,109 |
| 2022 | 291,702 | 2,902 | 111,429 | 4,878 | 6,513 | 25,074 | 65,665 | 3,098 | 43,032 | 5,333 | 2,325 | 1,887 | 13,145 | 4,077 | 1,310 | 1,034 |
| 2023 | 240,424 | 3,077 | 100,090 | 67,936 | 2,711 | 3,381 | 11,777 | 23,375 | 2,626 | 14,349 | 1,999 | 912 | 1,403 | 5,537 | 862 | 388 |



Figure B.13. The proportion of Pacific Hake in Canadian waters by year and age. The yellow tiles mean that all the fish at that age were found in the Canadian part of the survey only. Dark blue means that all fish of that age were found in the U.S. part of the survey only, or for ages 19 and 20, that there were no data. Figure produced by Rebecca Thomas (NOAA).

Table B.5. Proportion of survey biomass estimated in the U.S.A.

| Year | Biomass | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2005 | 707,378 | 0 | 76,133 | 10,331 | 49,462 | 36,282 | 401,433 | 45,855 | 15,065 | 26,367 | 19,430 | 11,565 | 9,384 | 655 | 3,242 | 0 | 470 | 584 | 1,119 | 0 | 0 |
| 2007 | 682,827 | 0 | 126,820 | 15,197 | 100,304 | 11,749 | 38,143 | 25,269 | 298,298 | 29,490 | 14,727 | 10,392 | 4,704 | 5,546 | 2,186 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2009 | 1,104,157 | 0 | 3,424 | 345,098 | 580,433 | 13,521 | 67,320 | 5,293 | 8,109 | 17,164 | 39,179 | 12,572 | 4,840 | 2,994 | 4,008 | 95 | 108 | 0 | 0 | 0 | 0 |
| 2011 | 602,165 | 0 | 138,137 | 370,905 | 24,062 | 19,958 | 24,340 | 4,652 | 4,385 | 1,456 | 2,135 | 0 | 10,548 | 1,171 | 416 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2012 | 1,140,742 | 0 | 614,258 | 139,229 | 253,797 | 33,700 | 37,162 | 23,940 | 16,112 | 2,376 | 2,980 | 2,874 | 3,200 | 7,858 | 3,254 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2013 | 1,805,243 | 0 | 44,844 | 1,323,574 | 108,111 | 170,667 | 16,315 | 40,496 | 59,942 | 13,214 | 13,058 | 82 | 133 | 5,092 | 7,072 | 1,821 | 790 | 33 | 0 | 0 | 0 |
| 2015 | 1,698,059 | 0 | 119,473 | 135,552 | 68,994 | 980,876 | 93,396 | 153,672 | 34,322 | 35,177 | 35,855 | 22,023 | 1,715 | 4,975 | 0 | 2,533 | 6,470 | 3,025 | 0 | 0 | 0 |
| 2017 | 1,028,202 | 0 | 6,443 | 556,809 | 28,241 | 36,013 | 32,431 | 242,600 | 40,633 | 27,601 | 12,706 | 9,815 | 18,289 | 9,043 | 3,447 | 2,132 | 0 | 0 | 452 | 0 | 1,547 |
| 2019 | 1,527,135 | 0 | 101,995 | 382,877 | 22,970 | 509,596 | 43,528 | 54,597 | 47,642 | 279,611 | 39,302 | 17,226 | 6,442 | 6,574 | 8,340 | 3,986 | 0 | 2,174 | 274 | 0 | 0 |
| 2021 | 1,458,573 | 0 | 103,310 | 86,591 | 201,176 | 400,084 | 50,190 | 297,866 | 49,639 | 29,699 | 26,877 | 165,189 | 24,819 | 4,760 | 3,739 | 7,775 | 1,165 | 4,333 | 1,361 | 0 | 0 |
| 2023 | 885,014 | 0 | 335,555 | 250,521 | 11,566 | 13,620 | 33,344 | 93,009 | 11,854 | 64,783 | 9,473 | 7,550 | 8,630 | 31,719 | 5,980 | 4,870 | 1,306 | 183 | 1,052 | 0 | 0 |

Table B.6. Proportion of survey biomass estimated in Canada

| Year | Biomass | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2005 | 668,721 | 0 | 56,703 | 3,392 | 18,366 | 32,614 | 347,059 | 64,653 | 27,257 | 45,003 | 34,090 | 20,770 | 8,601 | 7,364 | 2,199 | 0 | 141 | 175 | 336 | 0 | 0 |
| 2007 | 259,894 | 0 | 440 | 2,035 | 18,629 | 3,224 | 17,398 | 12,669 | 152,072 | 18,048 | 11,027 | 9,710 | 6,024 | 4,918 | 2,258 | 717 | 262 | 461 | 0 | 0 | 0 |
| 2009 | 398,117 | 0 | 967 | 4,483 | 30,399 | 22,526 | 74,367 | 15,335 | 22,546 | 22,742 | 114,687 | 52,557 | 23,060 | 3,092 | 6,660 | 1,490 | 3,208 | 0 | 0 | 0 | 0 |
| 2011 | 72,409 | 0 | 0 | 10,007 | 2,114 | 7,143 | 9,038 | 2,663 | 3,959 | 2,388 | 3,792 | 7,557 | 14,532 | 5,892 | 2,569 | 393 | 196 | 169 | 0 | 0 | 0 |
| 2012 | 138,679 | 0 | 520 | 6,816 | 21,233 | 5,640 | 28,505 | 19,490 | 10,281 | 6,776 | 3,452 | 4,116 | 8,292 | 13,778 | 7,801 | 1,222 | 0 | 757 | 0 | 0 | 0 |
| 2013 | 123,992 | 0 | 23 | 2,101 | 8,258 | 19,331 | 7,947 | 19,406 | 15,465 | 6,858 | 4,669 | 2,936 | 4,427 | 7,227 | 16,463 | 5,877 | 1,866 | 540 | 597 | 0 | 0 |
| 2015 | 457,794 | 0 | 1,738 | 19,326 | 17,273 | 282,824 | 30,457 | 52,041 | 12,237 | 10,888 | 10,401 | 6,637 | 1,381 | 2,063 | 2,027 | 2,179 | 4,820 | 975 | 527 | 0 | 0 |
| 2017 | 389,610 | 0 | 1,108 | 51,919 | 12,181 | 22,813 | 26,322 | 178,693 | 24,911 | 32,341 | 7,116 | 10,756 | 10,203 | 3,583 | 4,510 | 789 | 0 | 0 | 2,362 | 0 | 0 |
| 2019 | 190,892 | 0 | 27,387 | 7,271 | 4,064 | 33,244 | 5,224 | 10,259 | 9,043 | 62,313 | 12,260 | 8,901 | 1,247 | 1,848 | 4,782 | 2,637 | 0 | 27 | 386 | 0 | 0 |
| 2021 | 66,069 | 0 | 696 | 2,864 | 7,984 | 12,852 | 3,464 | 13,410 | 3,667 | 4,693 | 6,860 | 6,598 | 1,912 | 860 | 107 | 29 | 33 | 0 | 39 | 0 | 0 |
| 2023 | 22,079 | 0 | 829 | 1,150 | 296 | 633 | 3,099 | 7,706 | 544 | 4,138 | 457 | 363 | 703 | 1,355 | 476 | 148 | 56 | 125 | 0 | 0 | 0 |

## B. 2 Day 2

## Request 1

The SRG requests rerunning the GLM for weight-at-age without unsexed hake, and creating a weight-at-age matrix using the updated outputs by averaging Males and Females equally.

JTC Response

- The GLMM was run after filtering the data to remove unsexed fish.
- The model output was truncated to just produce males and females given unsexed fish were no longer in the data used to fit the GLMM.
- Results for males and females were averaged.
- The weight-at-age file for SS3 was updated.
- The updated weight-at-age file was used to process the fishery-composition data.


## Request 2

The SRG requests a plot of the day effect in the maturity analysis for age- 3 hake for any given year, assuming the peak of maturity is the same in all years (please confirm this assumption).

## JTC Response

The formula used for the non-spatial effects of the maturity-at-age model looks like the following:
maturity $_{\text {functional }} \sim-1+s($ doy $)+$ age + age $e^{2}+$ temp + temp $^{2}$
where there is NO interaction with the smoother on the day of the year. Thus, the shape of the relationship (G.5) does not change with time or temperature in this case.

## Request 3

The SRG requests that the maturity plot be recalculated for Day 278, peak of maturity.

## JTC Response

The updated figure was presented at the SRG meeting and Figure 11 was also updated.

## Request 4

The SRG asks for a presentation of the recalculated probability mature by age matrix.

## JTC Response

The JTC created a heatmap-type plot (Figure B.14) which is similar to the figures in the document for weight-at-age and fecundity-at-age.


Figure B.14. Heatmap showing the maturity-at-age by year. Projected years (2024-2027) are the means of the previous five years (2019-2023). Values used for years prior to 2009 are the means or the values by age for 2009-2023.

## Request 5

The SRG requests rerunning the base assessment model with updated weight-at-age and fecundity matrices based on day 278. Updated harvest scenario tables if possible.

## JTC Response

The JTC ran the modified base model and presents the following figures.


Figure B.15. Female spawning biomass across time for the terminal bridging steps to the new base model, as well as the pre-SRG base model. The bridge model is the step in which the student-t distribution was used for the relative age-1 index of numbers of fish. Weight-at-age data were then updated, followed by fecundity using the new maturity ogives to create the new base model.

## Request 6

The SRG requests a new executive summary based on the updated base model.

## JTC Response

The JTC completed the new Executive Summary and submitted it to the SRG in an unchecked, draft form. It has been since completed and is the new Executive Summary for this document.

## Notes:

- The forecast for a constant catch of 270,000 t was removed and the TAC from 2022 (545,000 t) was added. This results in the row letters being different for roughly half of the forecast catch streams in the tables when compared with the submitted pre-SRG Executive Summary.
- The One-page summary has been modified, in particular the penultimate bullet point.


Figure B.16. Female spawning biomass relative to unfished female spawning biomass across time for the terminal bridging steps to the new base model, as well as the pre-SRG base model. The bridge model is the step in which the student-t distribution was used for the relative age- 1 index of numbers of fish. Weight-at-age data were then updated, followed by fecundity using the new maturity ogives to create the new base model.

## C GLOSSARY OF TERMS USED IN THIS DOCUMENT

40:10 adjustment: A reduction in the overall total allowable catch that is triggered when the female spawning biomass falls below $40 \%$ of its unfished equilibrium level. This adjustment reduces the total allowable catch on a straight-line basis from the $40 \%$ level such that the total allowable catch would equal zero when the biomass is at $10 \%$ of its unfished equilibrium level. This is one component of the default harvest policy.

Acceptable biological catch (ABC): The acceptable biological catch is a scientific calculation of the sustainable harvest level of a fishery used historically to set the upper limit for fishery removals by the Pacific Fishery Management Council. It is calculated by applying the estimated (or proxy) harvest rate that produces the maximum sustainable yield from the estimated vulnerable biomass. For Pacific Hake, the calculation of the acceptable biological catch and application of the 40:10 adjustment is now replaced with the default harvest rate and the total allowable catch.

Adjusted: A term used to describe the total allowable catch or allocations that account for carryovers of uncaught catch from previous years.

Advisory Panel (AP): The Advisory Panel on Pacific Hake established by the Agreement.
Agreement ('Treaty'): The Agreement between the government of the United States and the government of Canada on Pacific Hake, signed in Seattle, Washington, on November 21, 2003 and entered into force June 25, 2008.

Alaska Fisheries Science Center (AFSC): One of six regional NOAA Fisheries Science Centers, primarily in Seattle, Washington but also present throughout Alaska.
$B_{0}$ : Unfished equilibrium female spawning biomass.
$B_{10 \%}$ : The level of female spawning biomass corresponding to $10 \%$ of unfished equilibrium female spawning biomass, i.e., $B_{10 \%}=0.1 B_{0}$. This is the level below which the calculated total allowable catch is set to 0 , based on the 40:10 adjustment.
$B_{40 \%}$ : The level of female spawning biomass corresponding to $40 \%$ of unfished equilibrium female spawning biomass, i.e., $B_{40 \%}=0.4 B_{0}$. This is the level below which the total allowable catch is decreased from the value associated with $F_{\mathrm{SPR}=40 \%}$, based on the 40:10 adjustment.
$B_{\mathrm{MSY}}$ : The estimated female spawning biomass which theoretically would produce the maximum sustainable yield under equilibrium fishing conditions (constant fishing and average recruitment in every year).

Backscatter: The scattering by a target back in the direction of an acoustic source. Specifically, the Nautical Area Scattering Coefficient (a measure of scattering per area) is frequently referred to as backscatter.

Benchmark spawning potential ratio ( $B_{\mathrm{SPR}=40 \%}$ ): The spawning potential ratio of $40 \%$, where $40 \%$ relates to the default harvest rate of $F_{\mathrm{SPR}=40 \%}$ specified in the Agreement. Even under equilibrium conditions, $F_{\mathrm{SPR}=40 \%}$ would not necessarily result in a female spawning biomass of $B_{40 \%}$ because $F_{\mathrm{SPR}=40 \%}$ is defined in terms of the spawning potential ratio that depends on the female spawning biomass per recruit.

California Current Ecosystem: The waters of the continental shelf and slope off the west coast of North America, commonly referring to the area from Central California to Southern British Columbia.

Carryover: If at the end of the year, there are unharvested allocations, then there are provisions for some of these unharvested allocations to be carried over into the next year's allocation process. The Agreement states that "If, in any year, a Party's catch is less than its individual total allowable catch, an amount equal to the shortfall shall be added to its individual total allowable catch in the following year, unless otherwise recommended by the Joint Management Committee. Adjustments under this sub-paragraph shall in no case exceed 15 percent of a Party's unadjusted individual total allowable catch for the year in which the shortfall occurred."

Catchability $(q)$ : The parameter defining the proportionality between a relative index of abundance (often a fishery-independent survey) and the estimated abundance available to that survey (as modified by selectivity) in the assessment model.

Catch-per-unit-effort (CPUE): A raw or, frequently, standardized and model-based metric of fishing success based on the catch and relative effort expended to generate that catch. Catch-per-unit-effort is often used as an index of abundance in the absence of fishery-independent indices and/or where the two types of indices are believed to be proportional.

Catch target: A general term used to describe the catch value used for management. Depending on the context, this may be a limit rather than a target and may be equal to a total allowable catch, an acceptable biological catch, the median result of applying the default harvest policy, or some other number. The Joint Technical Committee welcomes input from the Joint Management Committee on the best terminology to use for these quantities.

Closed-loop simulation: A subset of a management strategy evaluation that iteratively simulates a population using an operating model, generates data from that population and passes it to an estimation method, uses the estimation method and a management strategy to provide management advice, which then feeds back into the operating model to simulate an additional fixed set of time before repeating this process.

Coefficient of variation (CV): A measure of uncertainty defined as the standard deviation divided by the mean.

Cohort: A group of fish born in the same year. Also see recruitment and year-class.
Constant catch: A catch scenario used for forecasting in which the same catch is used in successive years.
Default harvest policy (rate): The application of $F_{\text {SPR }=40 \%}$ with the $40: 10$ adjustment. Having considered any advice provided by the Joint Technical Committee, Scientific Review Group, or Advisory Panel, the Joint Management Committee may recommend a different harvest rate if the scientific evidence demonstrates that a different rate is necessary to sustain the resource.

Department of Fisheries and Oceans (DFO) Canada: See Fisheries and Oceans Canada.
Depletion: Prior to the 2015 assessment, depletion was used instead of relative spawning biomass. 'Relative depletion' was also used.

El Niño: Abnormally warm ocean climate conditions in the California Current Ecosystem as a result of broad changes in the Eastern Pacific Ocean across the eastern coast of Latin America (centered on Peru) often around the end of the calendar year.

Exploitation fraction: A metric of fishing intensity that represents the total annual catch divided by the estimated population biomass over a range of ages assumed to be vulnerable to the fishery (set to ages $2+$ in this assessment; note that
in some previous assessments it was $3+$ ). This value is not equivalent to the instantaneous rate of fishing mortality or the spawning potential ratio.
$F_{\mathrm{SPR}=40 \%}$ : The rate of fishing mortality estimated to give a spawning potential ratio of $40 \%$. Therefore, by definition this satisfies

$$
\begin{equation*}
0.4=\frac{\text { spawning biomass per recruit with } F_{40 \%}}{\text { spawning biomass per recruit with no fishing }} \tag{C.1}
\end{equation*}
$$

and SPR $\left(F_{40 \%}\right)=40 \%$. The $40 \%$ value is specified in the Agreement.
$F_{\mathrm{SPR}=40 \%}-40: 10$ harvest policy: The default harvest policy.
Female spawning biomass: The biomass of mature female fish at the beginning of the year. Sometimes abbreviated to spawning biomass.

Fisheries and Oceans Canada (DFO): Federal organization that delivers programs and services to support sustainable use and development of Canada's waterways and aquatic resources. Was previously called Department of Fisheries and Oceans.
Fishing intensity: A measure of the magnitude of fishing, defined for a fishing rate $(F)$ as:

$$
\begin{equation*}
\text { fishing intensity for } F=1-\operatorname{SPR}(F) \tag{C.2}
\end{equation*}
$$

where $\operatorname{SPR}(F)$ is the spawning potential ratio for the value of $F$ accumulated over the entire year. It is often given as a percentage. Relative fishing intensity (Figures C. 1 and C.2) is the fishing intensity relative to that at the SPR fishing rate $F_{\mathrm{SPR}=40 \%}$, where $F_{\mathrm{SPR}=40 \%}$ is the $F$ that gives an SPR of $40 \%$ such that, by definition, $\operatorname{SPR}\left(F_{40 \%}\right)=40 \%$ (the benchmark spawning ratio). Therefore

$$
\text { relative fishing intensity for } \begin{align*}
F & =\frac{1-\operatorname{SPR}(F)}{1-\operatorname{SPR}\left(F_{40 \%}\right)}  \tag{C.3}\\
& =\frac{1-\operatorname{SPR}(F)}{1-0.4}  \tag{C.4}\\
& =\frac{1-\operatorname{SPR}(F)}{0.6} \tag{C.5}
\end{align*}
$$

For brevity we use $F_{\mathrm{SPR}=40 \%}=\operatorname{SPR}\left(F_{40 \%}\right)$ in the text. Although this simply equals $40 \%$, it can be helpful to explicitly write:

$$
\begin{equation*}
\text { relative fishing intensity for } F=\frac{1-\operatorname{SPR}(F)}{1-\operatorname{SPR}_{40 \%}} \tag{C.6}
\end{equation*}
$$

Fishing mortality rate or instantaneous rate of fishing mortality $(F)$ : A metric of fishing intensity that is usually reported in relation to the most highly selected ages(s) or length(s), or occasionally as an average over an age range that is vulnerable to the fishery. Because it is an instantaneous rate operating simultaneously with natural mortality, it is not equivalent to exploitation fraction, percent annual removal, or the spawning potential ratio.
$F_{\text {MSY }}$ : The rate of fishing mortality estimated to produce the maximum sustainable yield from the population.
Harvest strategy: A formal system for managing a fishery that includes the elements shown in Figure A. 1 of Taylor et al. (2015).

Harvest control rule: A process for determining an acceptable biological catch from a stock assessment. Also see default harvest policy.
Joint Management Committee (JMC): The Joint Management Committee is established by the Agreement.

Joint Technical Committee (JTC): The Joint Technical Committee established by the Agreement. The formal name is 'Joint Technical Committee of the Pacific Hake/Whiting Agreement Between the Governments of the United States and Canada'.

Kilotonne (kt). Metric abbreviation for 1,000 metric tonnes.
Logistic transformation: A mathematical transformation used to translate between numbers bounded within some range to numbers on the real line $(-\infty$ to $+\infty)$.
Magnuson-Stevens Fishery Conservation and Management Act (MSFCMA): The MSFCMA, sometimes known as the 'Magnuson-Stevens Act', established the 200mile fishery conservation zone, the regional fishery management council system, and other provisions of U.S. marine fishery law.
Management Strategy Evaluation (MSE): A formal process for evaluating harvest strategies.

Markov chain Monte Carlo (MCMC): A numerical method used to sample from the posterior distribution of parameters and derived quantities in a Bayesian analysis. It is more computationally intensive than computing the maximum likelihood estimate but provides a more accurate depiction of parameter uncertainty. See Stewart et al. (2013) for a discussion of issues related to differences between Markov chain Monte Carlo and maximum likelihood estimation.

Maximum sustainable yield (MSY): An estimate of the largest sustainable annual catch that can be continuously taken over a long period of time from a population under equilibrium ecological and environmental conditions.

Megatonne (Mt): Metric abbreviation for 1,000,000 metric tonnes.
Metric tonne ( t ): A unit of mass (often referred to as weight) equal to 1,000 kilograms or $2,204.62$ pounds. Some previous stock assessments used the abbreviation ' $\mathrm{mt}^{\prime}$.

National Marine Fisheries Service (NMFS): See NOAA Fisheries.
No-U-Turn Sampler (NUTS): An advanced Hamiltonian Bayesian Markov chain Monte Carlo sampling algorithm used to efficiently create posterior distributions and used in Pacific Hake Bayesian assessments beginning in 2021.

NOAA Fisheries: The division of the United States National Oceanic and Atmospheric Administration (NOAA) responsible for conservation and management of off-
shore fisheries (and inland salmon). This is also known as the National Marine Fisheries Service (NMFS), and both names are commonly used at this time.

North Pacific Groundfish and Halibut Observer Program Database (NORPAC): A database that stores data collected at sea by U.S. fishery observers.

Northwest Fisheries Science Center (NWFSC): One of six regional NOAA Fisheries Science Centers, primarily in Seattle, Washington but also in various locations in Oregon and Washington.

Not available (NA): Something that is not available, e.g., an entry in a table.
Operating model (OM): A model used to simulate data for use in the management strategy evaluation. The operating model includes components for the population and fishery dynamics, as well as the simulation of the data sampling process, potentially including observation error. Cases in the management strategy evaluation represent alternative configurations of the operating model.
Pacific Biological Station (PBS): The Pacific Biological Station of Fisheries and Oceans Canada located in Nanaimo, British Columbia.

Pacific Coast Fisheries Information Network (PacFIN): A database that provides a central repository for commercial fishery information from Washington, Oregon, and California.

Pacific Fishery Management Council (PFMC): The U.S. organization under which historical stock assessments for Pacific Hake were conducted.

Pacific Hake: Common name for Merluccius productus, the species whose offshore population in the waters of the United States and Canada is subject of this assessment.
Pacific whiting: An alternative name for Pacific Hake commonly used in the United States.
Posterior distribution: The probability distribution for parameters or derived quantities from a Bayesian model representing the result of the prior probability distributions being updated by the observed data via the likelihood equation. For stock assessments, posterior distributions are approximated via numerical methods; one frequently employed method is Markov chain Monte Carlo sampling.

Prior distribution: A probability distribution for a parameter in a Bayesian analysis that represents the information available before evaluating the observed data via the likelihood equation. For some parameters, uninformative priors can be constructed which allow the data to dominate the posterior distribution. For other parameters, informative priors can be constructed based on auxiliary information and/or expert knowledge or opinions.
$R_{0}$ : Estimated annual recruitment at unfished equilibrium.
Random walk Metropolis Hastings (rwMH): Bayesian Markov chain Monte Carlo sampling algorithm used to create posterior distributions used in Pacific Hake Bayesian stock assessment models prior to 2021.

Recruits/recruitment: the estimated number of new members in a fish population born in the same age. In this assessment, recruitment is reported at age 0 . See also cohort and year-class.

Recruitment deviation: The offset of the recruitment in a given year relative to the stock-recruitment relationship; values occur on a logarithmic scale and are relative to the expected recruitment at a given female spawning biomass.

Relative fishing intensity: See fishing intensity.
Relative spawning biomass: The ratio of the beginning-of-the-year female spawning biomass to the unfished equilibrium female spawning biomass $\left(B_{0}\right)$. Thus, lower values are associated with fewer mature female fish. This term was introduced in the 2015 stock assessment as a replacement for 'depletion'.

Scientific Review Group (SRG): The Scientific Review Group established by the Agreement.
Scientific and Statistical Committee (SSC): The scientific advisory committee to the Pacific Fishery Management Council. The Magnuson-Stevens Fishery Conservation and Management Act requires that each council maintain a Scientific and Statistical Committee to assist in gathering and analyzing statistical, biological, ecological, economic, social, and other scientific information that is relevant to the management of the Council.

Simulation: A model evaluation under a particular state of nature, including combinations of parameters controlling stock productivity, stock status, and the time series of recruitment deviations. In this assessment, there are 8,000 simulations used to characterize alternative states of nature, each of which are based on a sample from the posterior distribution of the parameters, as calculated using Markov chain Monte Carlo, for a particular model (e.g., the base model).

Spawning biomass: Abbreviated term for female spawning biomass.
Spawning biomass per recruit: The expected lifetime contribution of an age-0 recruit, calculated as the sum across all ages of the product of spawning biomass at each age and the probability of surviving to that age. See Figure C. 2 for a graphical demonstration of the calculation of this value, which is found in both numerator and denominator of the spawning potential ratio.

Spawning potential ratio (SPR): The ratio of the spawning biomass per recruit under a given level of fishing to the estimated spawning biomass per recruit in the absence of fishing; i.e., for fishing mortality rate F. Often expressed as a percentage, it achieves a value of $100 \%$ in the absence of fishing and declines toward zero as fishing intensity increases (Figure C.2).

Standard deviation (sd): A measure of variability within a sample.
Steepness (h): A parameter of the stock-recruitment relationship representing the proportion of $R_{0}$ expected (on average) when the female spawning biomass is reduced to $20 \%$ of $B_{0}$ (i.e., when relative spawning biomass is equal to $20 \%$ ).

Stock Synthesis (SS): The age-structured stock assessment model applied in this stock assessment.

Target strength (TS): The amount of backscatter from an individual acoustic target.
Total allowable catch (TAC): The maximum fishery removal under the terms of the Agreement.
U.S./Canadian allocation: The division of the total allowable catch of $73.88 \%$ as for the U.S. share and $26.12 \%$ for the Canadian share.

Vulnerable biomass: The demographic portion of the population available for harvest by the fishery.

Year-class: A group of fish born in the same year. See also 'cohort' and 'recruitment'.


Figure C.1. Fishing intensity as a function of the spawning potential ratio (SPR; top axis) and 1 -SPR (bottom axis); given the benchmark SPR of $40 \%$, the solid blue line is simply $1 / 0.6$, as shown in equation (C.3).

Numbers per recruit


Biomass per recruit


Female spawning biomass per recruit


Figure C.2. Illustration of the spawning potential ratio (SPR) calculation based on the combination of maturity and fecundity used in the model, using the Markov chain Monte Carlo (MCMC) estimates of natural mortality, selectivity, and fishing mortality in the final year of the base model used in this year's assessment. The light blue bars represent unfished values, the dark blue bars represent fished values.

## D REPORT OF THE 2023 PACIFIC HAKE FISHERY IN CANADA

Prepared by the Canadian Advisory Panel and submitted on February 13, 2024 for inclusion in this assessment document.

The Canadian Offshore Pacific Hake fishery was very poor in 2023. The 2023 year recorded the lowest catch since the fishery began in the late 1970s, with slightly more than 22,000 tonnes caught, or $21 \%$ of the 105,000 tonne TAC for the fishery. This is down from the 31,000 tonnes caught in 2022 when $29 \%$ of the 105,000 tonne TAC was harvested. Generally, the fishermen found the fishing very poor and similar or worse than 2022. Catches were inconsistent and usually very scratchy. Most of the fishing occurred around Father Charles and the Finger Bank and south of there. The fish encountered were large older fish and small fish were rarely encountered. The bycatch was primarily Yellowtail rockfish, pollock, and herring. The market for HGT was also poor.

## E REPORT OF THE 2023 PACIFIC HAKE FISHERY IN THE UNITED STATES

## Prepared by the United States Advisory Panel on 17 January 2024 for inclusion in this assessment document.

Based on data from the Pacific Fisheries Information Network (PacFIN), total U.S. harvest in the 2023 whiting fishery was $240,189 \mathrm{t}$, which is $52 \%$ of the U.S. allocation of $461,750 \mathrm{t}$. Total U.S. harvest from these same sectors in the 2022 whiting fishery was 289,726 t. Compared to 2022, total U.S. catch was down 17\% in 2023 (see Tables E. 1 and E.2).

Shoreside (SS) Fishery: Spring fishing in the SS sector started strong shortly after May 1 but slowed considerably by the end of May, and didn't pick back up until later in the year. Spring fishing was described as very spotty for both the fleet fishing around the Columbia River and north, as well as the fleet fishing out of Newport, Oregon. By June, almost all of the fish were observed to be off Newport and well to the south. The most consistent volumes in the SS fishery were observed from late June to early September, and the majority of the SS late summer and fall fishery occurred off the central OR coast and closer to the California border. Fishing in the SS sector slowed by October, which is normal.

Larger fish were reported further north during the times when fish were up north. The fleet fishing north of the Columbia River and in WA landed fish as large as 900 grams. Fish caught off the south coast of OR and CA were reported to be sub- 450 gram at best and got smaller later in the season.

Bycatch encounters in the SS sector consisted primarily of Chinook salmon, sablefish, darkblotched rockfish, yellowtail rockfish, and some mackerel off Newport late in the season. The SS whiting cooperative implemented several hot spot closures/advisory areas during the season to avoid/minimize bycatch.

Overall, the total 2023 SS whiting catch (100,392 t) represented $56 \%$ of the 2023 SS allocation and was slightly lower than total 2022 SS catch (104,323 t).

At-Sea Fishery: The U.S. at-sea fishery is comprised of the Mothership (MS) and Catcher Processor (CP) sectors.

Mothership (MS) Sector - In the spring fishery, one MS fleet was on the grounds on May 2, and another joined on May 10. Two other platforms joined on May 15. Spring fishing was strong until late May, and all four MS fleets concluded operations by May 30.

The size of whiting in the MS spring fishery generally ranged from 400-600 grams. However, location of fish further south and in shallower waters during the spring increased bycatch. By the end of May, slower fishing rates combined with high bycatch encounters and restrictive bycatch movement rules led to an early end to the spring fishery for the MS sector.

In the "fall" fishery (which began mid-summer this year), one MS fleet was on the grounds on August 11, with another joining on September 27. Fall catch occurred primarily off of

Table E.1. Final 2022 allocations (after reallocation of tribal quota) and catch totals (tonnes, t). Note that $402,646 \mathrm{t}$ U.S. TAC is reduced by 750 t for research and incidental catch set. Minor differences in catch in this table compared to that used in the stock assessment can occur due to the timing of data extractions. Source: 2023 Whiting Stock Assessment Report and PacFIN Whiting Report.

|  | U.S. TAC | Shoreside | Catcher Processor | Mothership | Tribal |
| :--- | ---: | ---: | ---: | ---: | ---: |
| Allocation (t) | 401,896 | 156,002 | 126,287 | 89,144 | 30,463 |
| Catch (t) | 289,726 | 104,323 | 126,247 | 59,157 | 1,174 |
| Utilization (\%) | $72.1 \%$ | $66.9 \%$ | $100.0 \%$ | $66.4 \%$ | $3.9 \%$ |

Oregon, with clean fishing and low bycatch encounters, but smaller fish size, averaging around 300 grams. The 2023 MS fishery concluded on November 1.

Overall, the total 2023 MS sector catch ( $32,744 \mathrm{t}$ ) represented $32 \%$ of the 2023 MS allocation and was significantly lower than the total 2022 MS catch ( $59,157 \mathrm{t}$ ). Four out of six MS platforms participated in the 2023 fishery, with one MS platform changing ownership mid-year.

Catcher Processor (CP) Sector - In the spring fishery, the first CPs were on the grounds on May 5. The CP sector ended their spring fishery the first week of June, as vessels transition to the Alaska pollock fishery which begins June 10. Overall, spring whiting harvest rates for the CPs were lower as fish were mostly absent from the WA and northern OR coastlines. Whiting were mainly schooled in southern OR and shallower than previous years. The shoreward and more shallow distribution of whiting contributed to much higher levels of incidental catch. The CPs observed higher encounters with Chinook salmon compared to previous years and record high encounters with some rockfish species during the spring fishery. The average whiting size in the spring CP fishery was approximately 450 grams.
The fall fishery began with the first CPs arriving in OR late August and continued through the beginning of November. Unlike spring fishing efforts, the fall was a turnaround for whiting harvest levels. A greater abundance of whiting were seen further north into OR and more normal depths. Southern OR exhibited steady fishing with good catch per unit effort (CPUE), and far lower bycatch than the spring fishery. Overall, there was very little CP catch and effort off Washington. The observed fish size in the fall fishery was smaller compared to the spring fishery, averaging approximately 350 grams.

Overall, the total 2023 CP sector catch $(107,053$ t) represented $74 \%$ of the 2023 CP allocation and was lower than total 2022 CP catch $(126,247 \mathrm{t})$, due to two vessels not participating during the 2023 whiting fishery. One CP vessel underwent repairs during the fall. Another vessel experienced a fire and was not operational in the whiting fishery this year.

Table E.2. Final 2023 allocations (after reallocation of tribal quota) and catch totals (tonnes, t). Note that $461,750 \mathrm{t}$ U.S. TAC is reduced by 750 t for research and incidental catch set. Minor differences in catch in this table compared to that used in the stock assessment can occur due to the timing of data extractions. Source: PacFIN Whiting Report.

|  | U.S. TAC | Shoreside | Catcher Processor | Mothership | Tribal |
| :--- | ---: | ---: | ---: | ---: | ---: |
| Allocation (t) | 461,000 | 178,581 | 144,566 | 102,047 | 35,806 |
| Catch (t) | 240,190 | 100,392 | 107,053 | 32,744 | 0 |
| Utilization (\%) | $52.1 \%$ | $56.2 \%$ | $74.1 \%$ | $32.1 \%$ | $0.0 \%$ |

## F ESTIMATED PARAMETERS IN THE BASE ASSESSMENT MODEL

Table F.1. Medians of estimated parameters for the base model.

| Parameter | Posterior median |
| :---: | :---: |
| NatM_uniform_Fem_GP_1 | 0.2348 |
| SR_LN(R0) | 14.7710 |
| SR_BH_steep | 0.8118 |
| Q_extraSD_Acoustic_Survey(2) | 0.3225 |
| Q_extraSD_Age1_Survey(3) | 0.3810 |
| $\ln$ (DM_theta)_Age_P1 | -0.6631 |
| $\ln ($ DM_theta)_Age_P2 | 2.7705 |
| Early_InitAge_20 | -0.3065 |
| Early_InitAge_19 | -0.0891 |
| Early_InitAge_18 | -0.1063 |
| Early_InitAge_17 | -0.1203 |
| Early_InitAge_16 | -0.1496 |
| Early_InitAge_15 | -0.1937 |
| Early_InitAge_14 | -0.2506 |
| Early_InitAge_13 | -0.2520 |
| Early_InitAge_12 | -0.2990 |
| Early_InitAge_11 | -0.3331 |
| Early_InitAge_10 | -0.4485 |
| Early_InitAge_9 | -0.4587 |
| Early_InitAge_8 | -0.4887 |
| Early_InitAge_7 | -0.5836 |
| Early_InitAge_6 | -0.5343 |
| Early_InitAge_5 | -0.4851 |
| Early_InitAge_4 | -0.2706 |
| Early_InitAge_3 | 0.0148 |
| Early_InitAge_2 | 0.3822 |
| Early_InitAge_1 | 0.6536 |
| Early_RecrDev_1966 | 0.5827 |
| Early_RecrDev_1967 | 1.6581 |
| Early_RecrDev_1968 | 1.2563 |
| Early_RecrDev_1969 | -0.2794 |
| Main_RecrDev_1970 | 2.3070 |
| Main_RecrDev_1971 | -0.0674 |
| Main_RecrDev_1972 | -0.5308 |
| Main_RecrDev_1973 | 1.8778 |
| Main_RecrDev_1974 | -0.9790 |
| Main_RecrDev_1975 | 0.6857 |
| Main_RecrDev_1976 | -1.5334 |
| Main_RecrDev_1977 | 1.9687 |
| Main_RecrDev_1978 | -1.9460 |
| Main_RecrDev_1979 | 0.3948 |

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| Parameter | Posterior median |
| :--- | ---: |
| Main_RecrDev_1980 | 2.9165 |
| Main_RecrDev_1981 | -1.2655 |
| Main_RecrDev_1982 | -1.0860 |
| Main_RecrDev_1983 | -0.5714 |
| Main_RecrDev_1984 | 2.6899 |
| Main_RecrDev_1985 | -1.9600 |
| Main_RecrDev_1986 | -1.6559 |
| Main_RecrDev_1987 | 1.9252 |
| Main_RecrDev_1988 | 0.7740 |
| Main_RecrDev_1989 | -2.1642 |
| Main_RecrDev_1990 | 1.5005 |
| Main_RecrDev_1991 | 0.2631 |
| Main_RecrDev_1992 | -2.0430 |
| Main_RecrDev_1993 | 1.2331 |
| Main_RecrDev_1994 | 1.2647 |
| Main_RecrDev_1995 | 0.3080 |
| Main_RecrDev_1996 | 0.6996 |
| Main_RecrDev_1997 | 0.1019 |
| Main_RecrDev_1998 | 0.8010 |
| Main_RecrDev_1999 | 2.6847 |
| Main_RecrDev_2000 | -1.0589 |
| Main_RecrDev_2001 | 0.2921 |
| Main_RecrDev_2002 | -3.1098 |
| Main_RecrDev_2003 | 0.5723 |
| Main_RecrDev_2004 | -3.1280 |
| Main_RecrDev_2005 | 1.1299 |
| Main_RecrDev_2006 | 0.8335 |
| Main_RecrDev_2007 | -3.5752 |
| Main_RecrDev_2008 | 1.8693 |
| Main_RecrDev_2009 | 0.4833 |
| Main_RecrDev_2010 | 2.9079 |
| Main_RecrDev_2011 | -0.8300 |
| Main_RecrDev_2012 | 0.5710 |
| Main_RecrDev_2013 | -0.9457 |
| Main_RecrDev_2014 | 2.1502 |
| Main_RecrDev_2015 | -3.3347 |
| Main_RecrDev_2016 | 1.8102 |
| Main_RecrDev_2017 | 0.4993 |
| Main_RecrDev_2018 | -0.8865 |
| Main_RecrDev_2019 | -1.2643 |
| Main_RecrDev_2020 | 1.6148 |
| Main_RecrDev_2021 | 2.3944 |
| Main_RecrDev_2022 | 0.6988 |
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| Parameter | Posterior median |
| :--- | ---: |
| Late_RecrDev_2023 | 0.0142 |
| ForeRecr_2024 | -0.0152 |
| ForeRecr_2025 | 0.0616 |
| ForeRecr_2026 | 0.0164 |
| ForeRecr_2027 | 0.0017 |
| AgeSel_P3_Fishery(1) | 3.0812 |
| AgeSel_P4_Fishery(1) | 0.8957 |
| AgeSel_P5_Fishery(1) | 0.4069 |
| AgeSel_P6_Fishery(1) | 0.1832 |
| AgeSel_P7_Fishery(1) | 0.4995 |
| AgeSel_P4_Acoustic_Survey(2) | 0.5250 |
| AgeSel_P5_Acoustic_Survey(2) | -0.1734 |
| AgeSel_P6_Acoustic_Survey(2) | 0.3007 |
| AgeSel_P7_Acoustic_Survey(2) | 0.3055 |
| AgeSel_P3_Fishery(1)_DEVadd_1991 | 0.5162 |
| AgeSel_P3_Fishery(1)_DEVadd_1992 | 0.0062 |
| AgeSel_P3_Fishery(1)_DEVadd_1993 | -0.0147 |
| AgeSel_P3_Fishery(1)_DEVadd_1994 | 0.0846 |
| AgeSel_P3_Fishery(1)_DEVadd_1995 | -0.1565 |
| AgeSel_P3_Fishery(1)_DEVadd_1996 | 0.3852 |
| AgeSel_P3_Fishery(1)_DEVadd_1997 | 0.0602 |
| AgeSel_P3_Fishery(1)_DEVadd_1998 | 0.1898 |
| AgeSel_P3_Fishery(1)_DEVadd_1999 | 0.9011 |
| AgeSel_P3_Fishery(1)_DEVadd_2000 | 0.4173 |
| AgeSel_P3_Fishery(1)_DEVadd_2001 | 0.0514 |
| AgeSel_P3_Fishery(1)_DEVadd_2002 | 0.0801 |
| AgeSel_P3_Fishery(1)_DEVadd_2003 | 0.0060 |
| AgeSel_P3_Fishery(1)_DEVadd_2004 | 0.2656 |
| AgeSel_P3_Fishery(1)_DEVadd_2005 | 0.0085 |
| AgeSel_P3_Fishery(1)_DEVadd_2006 | 0.5499 |
| AgeSel_P3_Fishery(1)_DEVadd_2007 | 0.5194 |
| AgeSel_P3_Fishery(1)_DEVadd_2008 | -0.0130 |
| AgeSel_P3_Fishery(1)_DEVadd_2009 | 0.3617 |
| AgeSel_P3_Fishery(1)_DEVadd_2010 | 0.8387 |
| AgeSel_P3_Fishery(1)_DEVadd_2011 | -0.2112 |
| AgeSel_P3_Fishery(1)_DEVadd_2012 | 0.0761 |
| AgeSel_P3_Fishery(1)_DEVadd_2013 | 0.2066 |
| AgeSel_P3_Fishery(1)_DEVadd_2014 | 0.2674 |
| AgeSel_P3_Fishery(1)_DEVadd_2015 | -0.8431 |
| AgeSel_P3_Fishery(1)_DEVadd_2016 | -0.0241 |
| AgeSel_P3_Fishery(1)_DEVadd_2017 | -0.4552 |
| AgeSel_P3_Fishery(1)_DEVadd_2018 | 0.629 |
| AgeSel_P3_Fishery(1)_DEVadd_2019 |  |
| Ca |  |

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| Parameter | Posterior median |
| :---: | :---: |
| AgeSel_P3_Fishery(1)_DEVadd_2020 | 0.0127 |
| AgeSel_P3_Fishery(1)_DEVadd_2021 | -0.5725 |
| AgeSel_P3_Fishery(1)_DEVadd_2022 | 1.7423 |
| AgeSel_P3_Fishery(1)_DEVadd_2023 | 0.6769 |
| AgeSel_P4_Fishery(1)_DEVadd_1991 | 0.3254 |
| AgeSel_P4_Fishery(1)_DEVadd_1992 | 0.5886 |
| AgeSel_P4_Fishery(1)_DEVadd_1993 | 0.8088 |
| AgeSel_P4_Fishery(1)_DEVadd_1994 | 0.1478 |
| AgeSel_P4_Fishery(1)_DEVadd_1995 | 0.2448 |
| AgeSel_P4_Fishery(1)_DEVadd_1996 | -0.3570 |
| AgeSel_P4_Fishery(1)_DEVadd_1997 | 1.2656 |
| AgeSel_P4_Fishery(1)_DEVadd_1998 | 0.9667 |
| AgeSel_P4_Fishery(1)_DEVadd_1999 | -0.0935 |
| AgeSel_P4_Fishery(1)_DEVadd_2000 | 0.8008 |
| AgeSel_P4_Fishery(1)_DEVadd_2001 | 0.9377 |
| AgeSel_P4_Fishery(1)_DEVadd_2002 | 0.6972 |
| AgeSel_P4_Fishery(1)_DEVadd_2003 | 0.6910 |
| AgeSel_P4_Fishery(1)_DEVadd_2004 | 0.4503 |
| AgeSel_P4_Fishery(1)_DEVadd_2005 | 0.6481 |
| AgeSel_P4_Fishery(1)_DEVadd_2006 | -0.1227 |
| AgeSel_P4_Fishery(1)_DEVadd_2007 | 0.2426 |
| AgeSel_P4_Fishery(1)_DEVadd_2008 | 0.4494 |
| AgeSel_P4_Fishery(1)_DEVadd_2009 | 0.7286 |
| AgeSel_P4_Fishery(1)_DEVadd_2010 | 0.1541 |
| AgeSel_P4_Fishery(1)_DEVadd_2011 | 1.0462 |
| AgeSel_P4_Fishery(1)_DEVadd_2012 | 0.2448 |
| AgeSel_P4_Fishery(1)_DEVadd_2013 | 0.8045 |
| AgeSel_P4_Fishery(1)_DEVadd_2014 | 0.4961 |
| AgeSel_P4_Fishery(1)_DEVadd_2015 | 0.1647 |
| AgeSel_P4_Fishery(1)_DEVadd_2016 | -0.9255 |
| AgeSel_P4_Fishery(1)_DEVadd_2017 | -0.5714 |
| AgeSel_P4_Fishery(1)_DEVadd_2018 | -0.5257 |
| AgeSel_P4_Fishery(1)_DEVadd_2019 | -0.7931 |
| AgeSel_P4_Fishery(1)_DEVadd_2020 | 0.8083 |
| AgeSel_P4_Fishery(1)_DEVadd_2021 | -0.0104 |
| AgeSel_P4_Fishery(1)_DEVadd_2022 | -1.3424 |
| AgeSel_P4_Fishery(1)_DEVadd_2023 | -0.1153 |
| AgeSel_P5_Fishery(1)_DEVadd_1991 | -0.8527 |
| AgeSel_P5_Fishery(1)_DEVadd_1992 | 0.0980 |
| AgeSel_P5_Fishery(1)_DEVadd_1993 | -0.0093 |
| AgeSel_P5_Fishery(1)_DEVadd_1994 | 0.8687 |
| AgeSel_P5_Fishery(1)_DEVadd_1995 | 0.2673 |
| AgeSel_P5_Fishery(1)_DEVadd_1996 | -0.3229 |
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| Parameter | Posterior median |
| :---: | :---: |
| AgeSel_P5_Fishery(1)_DEVadd_1997 | -0.1132 |
| AgeSel_P5_Fishery(1)_DEVadd_1998 | -0.6136 |
| AgeSel_P5_Fishery(1)_DEVadd_1999 | 0.0951 |
| AgeSel_P5_Fishery(1)_DEVadd_2000 | -0.1591 |
| AgeSel_P5_Fishery(1)_DEVadd_2001 | 0.3224 |
| AgeSel_P5_Fishery(1)_DEVadd_2002 | 0.5311 |
| AgeSel_P5_Fishery(1)_DEVadd_2003 | 0.7475 |
| AgeSel_P5_Fishery(1)_DEVadd_2004 | 0.6686 |
| AgeSel_P5_Fishery(1)_DEVadd_2005 | 0.7351 |
| AgeSel_P5_Fishery(1)_DEVadd_2006 | -0.0276 |
| AgeSel_P5_Fishery(1)_DEVadd_2007 | -0.0707 |
| AgeSel_P5_Fishery(1)_DEVadd_2008 | -0.3018 |
| AgeSel_P5_Fishery(1)_DEVadd_2009 | -0.3118 |
| AgeSel_P5_Fishery(1)_DEVadd_2010 | 0.5130 |
| AgeSel_P5_Fishery(1)_DEVadd_2011 | -0.7007 |
| AgeSel_P5_Fishery(1)_DEVadd_2012 | 0.1580 |
| AgeSel_P5_Fishery(1)_DEVadd_2013 | -0.1910 |
| AgeSel_P5_Fishery(1)_DEVadd_2014 | -0.4967 |
| AgeSel_P5_Fishery(1)_DEVadd_2015 | -0.0246 |
| AgeSel_P5_Fishery(1)_DEVadd_2016 | -0.0070 |
| AgeSel_P5_Fishery(1)_DEVadd_2017 | -0.1511 |
| AgeSel_P5_Fishery(1)_DEVadd_2018 | -0.2096 |
| AgeSel_P5_Fishery(1)_DEVadd_2019 | -0.0752 |
| AgeSel_P5_Fishery(1)_DEVadd_2020 | 0.6741 |
| AgeSel_P5_Fishery(1)_DEVadd_2021 | 0.6345 |
| AgeSel_P5_Fishery(1)_DEVadd_2022 | 0.1278 |
| AgeSel_P5_Fishery(1)_DEVadd_2023 | -0.8338 |
| AgeSel_P6_Fishery(1)_DEVadd_1991 | -0.0546 |
| AgeSel_P6_Fishery(1)_DEVadd_1992 | -0.4801 |
| AgeSel_P6_Fishery(1)_DEVadd_1993 | -0.0557 |
| AgeSel_P6_Fishery(1)_DEVadd_1994 | -0.0871 |
| AgeSel_P6_Fishery(1)_DEVadd_1995 | 0.7385 |
| AgeSel_P6_Fishery(1)_DEVadd_1996 | -0.1070 |
| AgeSel_P6_Fishery(1)_DEVadd_1997 | -0.3419 |
| AgeSel_P6_Fishery(1)_DEVadd_1998 | 0.3681 |
| AgeSel_P6_Fishery(1)_DEVadd_1999 | -0.3918 |
| AgeSel_P6_Fishery(1)_DEVadd_2000 | 0.1502 |
| AgeSel_P6_Fishery(1)_DEVadd_2001 | -0.1392 |
| AgeSel_P6_Fishery(1)_DEVadd_2002 | 0.1388 |
| AgeSel_P6_Fishery(1)_DEVadd_2003 | 0.2638 |
| AgeSel_P6_Fishery(1)_DEVadd_2004 | -0.5599 |
| AgeSel_P6_Fishery(1)_DEVadd_2005 | 0.2660 |
| AgeSel_P6_Fishery(1)_DEVadd_2006 | 0.2131 |

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| Parameter | Posterior median |
| :--- | ---: |
| AgeSel_P6_Fishery(1)_DEVadd_2007 | -0.2244 |
| AgeSel_P6_Fishery(1)_DEVadd_2008 | 0.2367 |
| AgeSel_P6_Fishery(1)_DEVadd_2009 | -0.3308 |
| AgeSel_P6_Fishery(1)_DEVadd_2010 | -0.3864 |
| AgeSel_P6_Fishery(1)_DEVadd_2011 | -0.2272 |
| AgeSel_P6_Fishery(1)_DEVadd_2012 | -0.4511 |
| AgeSel_P6_Fishery(1)_DEVadd_2013 | -0.0421 |
| AgeSel_P6_Fishery(1)_DEVadd_2014 | 0.0535 |
| AgeSel_P6_Fishery(1)_DEVadd_2015 | -0.0197 |
| AgeSel_P6_Fishery(1)_DEVadd_2016 | -0.0199 |
| AgeSel_P6_Fishery(1)_DEVadd_2017 | -0.1621 |
| AgeSel_P6_Fishery(1)_DEVadd_2018 | -0.3403 |
| AgeSel_P6_Fishery(1)_DEVadd_2019 | 0.1470 |
| AgeSel_P6_Fishery(1)_DEVadd_2020 | -0.3350 |
| AgeSel_P6_Fishery(1)_DEVadd_2021 | 0.1884 |
| AgeSel_P6_Fishery(1)_DEVadd_2022 | 0.5586 |
| AgeSel_P6_Fishery(1)_DEVadd_2023 | 0.1653 |
| AgeSel_P7_Fishery(1)_DEVadd_1991 | -0.0949 |
| AgeSel_P7_Fishery(1)_DEVadd_1992 | 0.0732 |
| AgeSel_P7_Fishery(1)_DEVadd_1993 | -0.3617 |
| AgeSel_P7_Fishery(1)_DEVadd_1994 | 0.0914 |
| AgeSel_P7_Fishery(1)_DEVadd_1995 | -0.1230 |
| AgeSel_P7_Fishery(1)_DEVadd_1996 | 0.4078 |
| AgeSel_P7_Fishery(1)_DEVadd_1997 | 0.1310 |
| AgeSel_P7_Fishery(1)_DEVadd_1998 | -0.5020 |
| AgeSel_P7_Fishery(1)_DEVadd_1999 | -0.2628 |
| AgeSel_P7_Fishery(1)_DEVadd_2000 | -0.0763 |
| AgeSel_P7_Fishery(1)_DEVadd_2001 | -0.2944 |
| AgeSel_P7_Fishery(1)_DEVadd_2002 | -0.4180 |
| AgeSel_P7_Fishery(1)_DEVadd_2003 | -0.2696 |
| AgeSel_P7_Fishery(1)_DEVadd_2004 | -0.1766 |
| AgeSel_P7_Fishery(1)_DEVadd_2005 | -0.3799 |
| AgeSel_P7_Fishery(1)_DEVadd_2006 | -0.3073 |
| AgeSel_P7_Fishery(1)_DEVadd_2007 | 0.0703 |
| AgeSel_P7_Fishery(1)_DEVadd_2008 | -0.1160 |
| AgeSel_P7_Fishery(1)_DEVadd_2009 | -0.0300 |
| AgeSel_P7_Fishery(1)_DEVadd_2010 | -0.7561 |
| AgeSel_P7_Fishery(1)_DEVadd_2011 | -0.4840 |
| AgeSel_P7_Fishery(1)_DEVadd_2012 | -0.2938 |
| AgeSel_P7_Fishery(1)_DEVadd_2013 | 0.1362 |
| AgeSel_P7_Fishery(1)_DEVadd_2014 | -0.0977 |
| AgeSel_P7_Fishery(1)_DEVadd_2015 | -0.4649 |
| AgeSel_P7_Fishery(1)_DEVadd_2016 | -0.3844 |
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| Parameter | Posterior median |
| :--- | ---: |
| AgeSel_P7_Fishery(1)_DEVadd_2017 | -0.0375 |
| AgeSel_P7_Fishery(1)_DEVadd_2018 | 0.2466 |
| AgeSel_P7_Fishery(1)_DEVadd_2019 | -0.1914 |
| AgeSel_P7_Fishery(1)_DEVadd_2020 | -0.0499 |
| AgeSel_P7_Fishery(1)_DEVadd_2021 | -0.2999 |
| AgeSel_P7_Fishery(1)_DEVadd_2022 | 0.0538 |
| AgeSel_P7_Fishery(1)_DEVadd_2023 | 0.5701 |

## G MODELING TEMPORAL AND SPATIAL TRENDS IN HAKE MATURITY-AT-AGE DATA

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## G. 1 Methods

## G.1.1 Data filtering

We assembled a dataset representing hake functional maturity collected from three sources, the West Coast Groundfish Bottom Trawl Survey (WCGBTS; Keller et al. 2017), the Northwest Fisheries Science Center Integrated Pacific Hake Ecosystem and Acoustic-Trawl Survey, and gonad tissue samples from the At-Sea Hake Observer program (ASHOP). The first two sources represent fishery-independent samples, while ASHOP samples are fishery dependent. After filtering, this dataset consisted of 2836 samples, from years 2009 to 2021 (Table G.1; Figure G. 1 ). An additional 180 samples exist with unknown ages and 394 samples exist with uncertain functional maturity; but, these were omitted from our analysis. Similarly, there are 73 samples that were collected in British Columbia ( 72 of 73 mature; 70 of 73 age 5 or older) but they have limited temporal coverage (primarily 2013, 2015) and were omitted from our analysis. Note that these samples from British Columbia were included in the 2018 analysis of maturity-at-age that was previously included in the fecundity relationship.

## G.1.2 Statistical modeling

As each sample in our dataset is geo-referenced with a unique latitude and longitude, we constructed a series of spatiotemporal models to model variation in Pacific Hake maturity. Our statistical modeling framework can be seen as a version of generalized linear mixed model (GLMM), with a series of fixed and random components. We constructed three models to test hypotheses and help improve understanding of spatiotemporal variability in Pacific Hake maturity; these models can generally be described as

1. a null model representing the status quo, similar to models used in previous Pacific Hake assessments but with additional data and an effect for calendar day;
2. a spatiotemporal model incorporating temporal and spatial trends but no covariates;
3. a spatiotemporal model incorporating temperature as an environmental predictor.

To maintain consistency with the status quo, our null model contained no spatial or temporal variation. We included quadratic effects of age (to account for potential skip spawning of older individuals), a smooth effect of calendar day (modeled with a penalized regression or P-spline; Eilers and Marx 1996), and a linear offset corresponding to the ASHOP samples (allowing for differences between fishery independent and dependent samples). Previous Pacific Hake assessments have not allowed maturity to vary over time or space and have not estimated coefficients corresponding to ASHOP samples. Earlier versions of our first model also included offsets allowing for differences between the WCGBTS and Acoustic-Trawl samples, however these estimates were small in magnitude
and had large standard errors, which suggests there were not meaningful differences between these sampling platforms. The null model can be written as

$$
\begin{equation*}
E[\mathbf{y}]=g^{-1}(\mathbf{X} \boldsymbol{\beta}+\mathbf{Z} \mathbf{b}) \tag{G.1}
\end{equation*}
$$

where $g^{-1}()$ represents the inverse logit function, $\mathbf{X}$ represents the design matrix of fixed effects with estimated parameters $\boldsymbol{\beta}$, and the P -spline is represented with random effect design matrix $\mathbf{Z}$ and corresponding coefficients $\mathbf{b}$.

As a second model, we extended our null model to include spatial and temporal variation. We modeled year effects (intercepts) with a time-varying AR(1) random walk. Spatial effects were modeled using the stochastic partial differential equation (SPDE) approximation to Gaussian Random Fields (Lindgren et al. 2011). This approach can be seen as a predictive process method, where a spatial surface is approximated by a series of estimated random effects (locations referred to knots or vertices) which are then projected to locations of sample collections. Our mesh representing the spatial field used a cutoff distance of 50 km , resulting in 97 mesh vertices. Due to the sparsity of data in some years, we only considered models with spatial, and not spatiotemporal, fields. However, to account for interactions between space and age, we also included spatially varying coefficients in age effects, as a quadratic relationship (spatial fields for the intercept, age, age ${ }^{2}$ ). This model can be written as

$$
\begin{equation*}
E[\mathbf{y}]=g^{-1}\left(\mathbf{X} \boldsymbol{\beta}+\mathbf{Z} \mathbf{b}+\boldsymbol{\omega}_{s}+\mathbf{x}_{a g e} * \boldsymbol{\zeta}_{s, a g e}+\mathbf{x}_{a g e^{2}} * \boldsymbol{\zeta}_{s, a g e^{2}}+\mathbf{X}_{t}^{t v c} \boldsymbol{\gamma}_{\mathbf{t}}\right) \tag{G.2}
\end{equation*}
$$

where new components $\boldsymbol{\omega}_{s}$ represent a constant spatial field shared across years (analogous to a spatial intercept), $\boldsymbol{\zeta}_{s, a g e}$ and $\boldsymbol{\zeta}_{s, a g e^{2}}$ represent spatially varying coefficient effects of age, and $\mathbf{X}_{t}^{t v c}$ represents the design matrix of year effects with estimated coefficients $\gamma_{\mathbf{t}}$. The $\gamma_{\mathbf{t}}$ terms were modeled as $\operatorname{AR}(1)$ terms such as in the example $\gamma_{2010} \sim N\left(\rho \gamma_{2009}, \sqrt{1-\rho^{2}} \sigma\right)$, where $\rho$ represents the $\operatorname{AR}(1)$ parameter.
Our third model also extended the null model to include an estimated spatial field and spatially varying coefficients for the quadratic function of age; but, instead of modeling year effects as a random walk, year effects were modeled as a function of subsurface (at 130.67 m ; see below for details) temperature indices in the domain of the WCGBTS survey. Replacing random year effects with a temperature covariate has the potential to add mechanistic relationships to the modeling and also reduce uncertainty in years where no or few samples are collected. We explored several forms of the temperature relationship, including linear, quadratic, or P-splines and chose to use quadratic relationships because these introduced fewer parameters than P-splines, while also allowing for parabolic relationships. This model can be written in the same form as Model 2, without the time varying intercept,

$$
\begin{equation*}
E[\mathbf{y}]=g^{-1}\left(\mathbf{X} \boldsymbol{\beta}+\mathbf{Z} \mathbf{b}+\boldsymbol{\omega}_{s}+\mathbf{x}_{a g e} * \boldsymbol{\zeta}_{s, a g e}+\mathbf{x}_{a g e^{2}} * \boldsymbol{\zeta}_{s, a g e^{2}}\right) \tag{G.3}
\end{equation*}
$$

where the added quadratic effects of temperature are in the fixed effects components $\mathbf{X}$ and $\beta$.

Parameter estimation was done using the sdmTMB package (Anderson et al. 2022) in R ( R Core Development Team 2024). sdmTMB provides a convenient interface between R and

Template Model Builder (TMB; Kristensen et al. 2016), which allows for fast marginal maximum likelihood estimation. Model convergence was assessed by examining the Hessian and standard errors of parameter estimates and the maximum gradient at convergence. Area under the curve (AUC) estimates were also calculated using the pROC package (Robin et al. 2011) in R.

## G.1.3 Deriving temperature indices

Subsurface ocean temperature has previously been linked to Pacific Hake distribution (Malick et al. 2020) and co-occurrence of Pacific Hake with prey (Phillips et al. 2003); and, more recently, marine heat waves have been found to delay maturity in other groundfish species (Rosemond 2023). Given that temperature sampling is not done from all sampling platforms, we relied on modeled temperature products. Specifically, we used sea water potential temperature (referred to as 'temperature' throughout) from the GLORYS12v1 product ( $1 / 12^{\circ}$ resolution, 50 vertical levels, Global Ocean Physics Reanalysis for 1993-September 2023 and Global Ocean Physics Analysis and Forecast for October 2023-January 2024; Global Ocean Physics Reanalysis 2024, Global Ocean Physics Analysis and Forecast 2024; Lellouche et al. 2021). We processed monthly averages of temperature and used data from a subset of depths corresponding to the vertical distribution of Pacific Hake ( 25.21 m , $40.34 \mathrm{~m}, 55.76 \mathrm{~m}, 77.85 \mathrm{~m}, 92.33 \mathrm{~m}, 109.73 \mathrm{~m}, 130.67 \mathrm{~m})$. These data show that, as expected, temperature generally decreases with depth (Figure G.2) and unlike surface temperature, which is warmest in summer months, the warmest temperatures at depth often occur in winter (Figure G.3). Data from each month-depth combination was alternately used to generate annual indices of temperature to relate to maturity. As a final stratification, we also considered indices generated using (1) coastwide GLORYS12v1 temperatures, (2) temperature north of Point Conception, and (3) temperature south of Point Conception.
Rather than take a spatial average of temperature by year for each month-depth combination, we generated biomass weighted averages of temperature (weighting temperature by the spatial distribution of Pacific Hake biomass, rather than weighting each spatial location equally). To generate biomass weights, we constructed a spatial model of Pacific Hake catch per unit effort (CPUE, $\mathrm{kg} / \mathrm{km}^{2}$ ), using 2003-2023 data from the WCGBTS. We adopted the same SPDE approach used in modeling maturity. Our biomass model included a smooth effect of calendar day (as a P-spline, dates range from May-October), a time varying intercept modeled with a random walk, a spatial field (representing spatial variation shared across years), and spatiotemporal variation (representing year to year variation in spatial patterning) modeled as an $\operatorname{AR}(1)$ process. Given the skewed distribution of catches, we modeled CPUE with a Tweedie distribution (Shono 2008). We identified spatial cells from the GLORYS12v1 re-analysis that were in the spatial domain of the WCGBTS (Keller et al. 2017) and made predictions of biomass to those cells on July 1 of each year (July 1 was arbitrarily chosen as a date in the middle of the WCGBTS survey). Because of the random walk model for year effects, our predictions of Pacific Hake biomass in future years 2024-2025 are identical to 2023. Temperature indices were then generated by taking a weighted average of temperature across space, with estimated Pacific Hake biomass used as weights. As the WCGBTS survey only provides an annual snapshot of biomass, the same biomass weights were used for all depths and months
with temperature data. In general, weighted indices remained highly correlated with unweighted temperature indices ( $\rho=0.98$ across all depths and months; Figure G.4).

## G. 2 Model selection and sensitivity analysis

We performed two main sensitivity analyses across the three models used. First, we evaluated the sensitivity of our model selection results to include (or not) the fishery dependent ASHOP samples. Second, we evaluated a sensitivity to including samples south of Point Conception (previous maturity modeling has imposed a cutoff and did not include samples south of Point Conception). After ensuring that models converged (using the sanity () function in sdmTMB), we evaluated the relative support of models using Akaike information criterion (AIC; Akaike 1973). Similarly, alternative temperature indices were also compared using AIC, and the model(s) with the lowest AIC values were deemed to have the most support.

## G.2.1 Generation of annual maturity at age

To create a single index of maturity for each age-year combination and generate maturity ogives, we used results from the best model selected with temperature as a driver alongside model results for Models 1-2. For each of these three models, we generated model predictions onto the WCGBTS design grid for non-ASHOP samples (with a larger intercept, ASHOP samples have earlier maturity at age) on the 278th day of the year or October 5th (Table G.3). Rather than take a simple average of these estimates (which weights each spatial cell equally), we calculated a weighted average, using biomass weights. Biomass was predicted using the same spatiotemporal model fit to the WCGBTS CPUE data, used in generating weighted indices of temperature. Weights were applied to estimated probabilities of maturity in logit space, and the total weighted average for each age-year combination was converted to normal space with an inverse logit transformation.

## G. 3 Results

Comparing the spatiotemporal model of Pacific Hake maturity to the baseline model representing the status quo provides an evaluation of support for maturation varying over time and space. In our comparison, most models showed greater support for models that included spatiotemporal processes than the null model (Table G.2). Similarly, we found that the model with the temperature index to predict year effects received the most support when ASHOP samples were included and data south of Pt. Conception were excluded (Table G.2; AIC > 10). For other analysis with different subsets of data, the time varying random walk model without temperature as a driver received more support. For consistency with previous Pacific Hake stock assessment models, we focus the remainder of our results on models that were only fit to data north of Point Conception and included the ASHOP samples.

All three models estimated a similar quadratic effect of age and higher maturity at age in ASHOP samples (Table G.3). All three models had similar high AUC values (Model $1=0.951$, Model $2=0.963$, Model $3=0.962$ ) indicating high abilities to discriminate
between whether or not Pacific Hake are mature. Similarly, all three models estimated similar effects of calendar day, with a rapid increase in maturity estimated in summer months (Figure G.5).

For models that include spatiotemporal variation, we found similar patterns of year effects and spatial variation. Year effects from the time varying model without temperature indicated lower than average maturity in a number of years, including 2012, 2016, and 2019 (Figure G.6). Including spatially varying coefficients of age allowed each age to differ slightly; but, in general, spatial patterning was similar across ages (Figure G.7). Biomass weighted predictions of maturity also appeared similar between the spatiotemporal models with and without temperature effects (Figure G.8). The largest differences between these models appeared to be in years like 2016, when the temperature driven model predicts maturation rates lower than the random walk model (the classification ability of these models in 2016 was nearly identical, AUC $=0.957$ for both; Figure G.9).
The estimated marginal effect of temperature in our best model (Tables G. 2 and G.3) indicated a concave relationship between temperature and functional maturity (Figure G.10). Temperatures in 2018 and 2020 were near the peak of this relationship, while most years were cooler (maturity increasing with temperature over this region). Temperatures in 2016 and 2019 were high (Figures G.2-G.4), above the threshold corresponding to the peak of the temperature-maturity relationship, and maturity at age was estimated to decline in these years. Contrasts between years can also be seen in the estimated ogives (Figure G.11) with a delay in maturity occurring in years that are both cooler and warmer than 2020 (age at $50 \%$ maturity $=2.81$ in 2020, versus 3.58 in 2012 or 3.82 in 2016).

The estimate of the spatial variance for the model used to weight spatial estimates based on estimated biomass of Pacific Hake was higher than the estimate of the spatiotemporal variance suggesting that differences in locations are more prominent than differences in locations between years (Table G.4). That is, Pacific Hake have a patchy distribution but those patches appear in largely the same locations year after year. The center of gravity of the distribution was furthest to the north in years that correspond to high temperatures and subsequent decreases in maturity (Figure G.12).

## G. 4 Citations

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## G. 5 Tables

Table G.1. Samples included in our analysis (after filtering), by sampling platform and year.

| Platform | $\mathbf{2 0 0 9}$ | $\mathbf{2 0 1 2}$ | $\mathbf{2 0 1 3}$ | $\mathbf{2 0 1 4}$ | $\mathbf{2 0 1 5}$ | $\mathbf{2 0 1 6}$ | $\mathbf{2 0 1 7}$ | $\mathbf{2 0 1 8}$ | $\mathbf{2 0 1 9}$ | $\mathbf{2 0 2 1}$ |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| ASHOP | 0 | 0 | 135 | 196 | 131 | 194 | 177 | 0 | 0 | 0 |
| Acoustic | 0 | 181 | 186 | 0 | 160 | 131 | 57 | 54 | 59 | 68 |
| WCGBT | 244 | 64 | 63 | 197 | 216 | 66 | 102 | 109 | 46 | 0 |

Table G.2. Comparison of Akaike information criterion (AIC) values from the three models in our analysis. Model 1 represents the baseline model, Model 2 represents the model with spatiotemporal variation but no covariates, and Model 3 represents the model with spatial variation and a temperature effect. For the temperature effect, this table represents estimates from models using temperature in February, at a depth of 130.67 m , and only north of Point Conception. The best models (lowest AIC) for each combination are in bold.

| ASHOP | Include South | Model 1 | Model 2 | Model 3 |
| :---: | :---: | ---: | ---: | ---: |
| Y | Y | 1440.12 | $\mathbf{1 3 3 4 . 6 8}$ | 1366.41 |
| Y | N | 1277.58 | 1209.28 | $\mathbf{1 1 9 5 . 9 4}$ |
| N | N | 1688.75 | 893.45 | 902.61 |
| N | Y | 1912.61 | $\mathbf{1 0 1 4 . 1 3}$ | 1025.47 |

Table G.3. Parameter estimates and standard errors estimated for the three models of maturity in our analyses (including samples from the At-Sea Hake Observer Program (ASHOP) and only samples north of Pt. Conception). Estimates of fixed effects are in logit space. The spatial range and spatial variance $\left(\sigma_{O}\right)$ are also included for models with those terms.

| Model | Term | Estimate | Std Error |
| :---: | :---: | :---: | :---: |
| 1 | age | 4.460 | 0.2010 |
| 1 | age ${ }^{2}$ | -1.060 | 0.0557 |
| 1 | ashopFALSE | 3.510 | 0.1900 |
| 1 | ashopTRUE | 4.290 | 0.2200 |
| 2 | age | 4.520 | 0.2840 |
| 2 | age ${ }^{2}$ | -1.350 | 0.1050 |
| 2 | ashopFALSE | 3.740 | 0.3700 |
| 2 | ashopTRUE | 5.200 | 0.6910 |
| 2 | range | 191.000 | 85.4000 |
| 2 | $\sigma_{O}$ | 0.653 | 0.3660 |
| 3 | age | 4.510 | 0.2880 |
| 3 | age ${ }^{2}$ | -1.350 | 0.1020 |
| 3 | temp | 0.421 | 0.0994 |
| 3 | temp ${ }^{2}$ | -0.535 | 0.0754 |
| 3 | ashopFALSE | 4.360 | 0.3050 |
| 3 | ashopTRUE | 6.050 | 0.6770 |
| 3 | range | 202.000 | 95.2000 |
| 3 | $\sigma_{O}$ | 0.544 | 0.4480 |

Table G.4. Parameter estimates and standard errors estimated for the model of hake biomass from the West Coast Groundfish Bottom Trawl Survey (WCGBTS). The spatial range is included, with spatial variance $\left(\sigma_{O}\right)$, spatiotemporal variance $\left(\sigma_{E}\right)$, and Tweedie parameters $(\phi, p)$.

| Term | Estimate | Std. Error |
| :---: | ---: | ---: |
| range | 33.50 | 2.20000 |
| $\phi$ | 21.80 | 0.36700 |
| $\sigma_{O}$ | 3.78 | 0.19700 |
| $\sigma_{E}$ | 1.68 | 0.05410 |
| $p$ | 1.64 | 0.00368 |

## G. 6 Figures



Figure G.1. Breakdown of the number of observations by age, year, and sampling platform (i.e., West Coast Groundfish Bottom Trawl Survey (WCGBTS), At-Sea Hake Observer Program (ASHOP), and the Northwest Fisheries Science Center Integrated Pacific Hake Ecosystem and Acoustic-Trawl Survey).


Figure G.2. Average GLORYS12v1 potential temperature (degrees Celsius) within the domain of the West Coast Groundfish Bottom Trawl Survey (WCGBTS) north of Point Conception, in February by depth (m; colors).


Figure G.3. Average GLORYS12v1 potential temperature (degrees Celsius) within the domain of the West Coast Groundfish Bottom Trawl Survey (WCGBTS) north of Point Conception, at a depth of 130.67 m by month (colors).


Figure G.4. Comparison of temperature indices (Celsius) using raw averages of temperature (red) and biomass weighted averages of temperature (blue). Shown is temperature at a depth of 130.67 $m$ in January but only for north of Point Conception.


Figure G.5. The estimated effect of day of the year on maturation from Model 3 (see Table G.2) using a penalized spline.


Figure G.6. Estimated and predicted year effects from the spatiotemporal hake maturity model without covariates (Model 2 in Tables G. 2 and G.3). Year effects are included as an autocorrelated random walk (variance is larger in years without data, such as 2010-2011 or 2022-2025), and spatiotemporal effects are modeled with an $\operatorname{AR}(1)$ process.


Figure G.7. Spatial anomalies in predictions of maturity for ages 1-5. Predictions are centered such that blue represents lower than average maturation and red represents higher than average. Predictions are from Model 3 (see Table G.2) for year 2020. Slight variations between ages are driven by the spatially varying coefficient effect of age.


Figure G.8. Sensitivity results for the effect of temperature, with data coming from different months and depths, summarized by different regions ( $\mathrm{N}=$ north of Point Conception, $\mathrm{S}=$ south of Point Conception) and modeled using different functional relationships (linear, quadratic, and penalized spline, i.e., 'smooth'). In all cases, the raw Akaike information criterion (AIC) values are shown where a lower AIC corresponds to support for the model from the data.


Figure G.9. Biomass weighted estimates of maturity for each of the three models in our analysis (using samples from the At-Sea Hake Observer Program (ASHOP) and data north of Point Conception). Predictions of functional maturity are for non-ASHOP samples.


Figure G.10. Estimated marginal effect of February temperature (Celsius) at a depth of 130.67 $m$ when fit to data that included samples from the At-Sea Hake Observer Program (ASHOP) and only data north of Point Conception. Predictions are made for age-3 fish, using coefficients corresponding to the non-ASHOP surveys, and for the 278th day of the year.


Figure G.11. Estimated ogives in three years representing low temperatures (2012), average temperatures (2020), and high temperatures (2016). The model of maturity used February temperature at a depth of 130.67 m and was fit to data that included samples from the At-Sea Hake Observer Program (ASHOP) and only data north of Point Conception. Ogives represent Pacific Hake not sampled by ASHOP. The estimated age at $50 \%$ maturity is 2.81 in 2020, versus 3.58 in 2012, and 3.82 in 2016.


Figure G.12. Estimated center of gravity of Pacific Hake biomass from the West Coast Groundfish Bottom Trawl Survey (WCGBTS), using 2003-2023 data.


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