# Explorations of alternative stock assessment models for Eastern Bering Sea Pacific cod 


https://afsc-assessments.github.io/EBS PCOD/2023 ASSESSMENT/SEPTEMBER MODELS/

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## 1 Introduction

For 2023 the authors wished to examine outstanding problems common to all four of the eastern Bering Sea (EBS) Pacific cod ensemble models accepted for management in 2022. The two main issues with the ensemble models were: 1) for the length composition data the Dirichlet multinomial log(theta) values approach the upper bound and therefore needed to be fixed for the models to converge, 2) failing residual runs tests for length and age composition data in all ensembles indicating autocorrelation in the residuals pointing at poor residual behavior, 3) potential confounding of aging bias, annually varying growth, and annually varying selectivity result in the models being highly unstable with considerable tuning of the annual devs. on growth and selectivity required for model convergence, and 4) the models are highly sensitive to changes in catchability and natural mortality with small changes in either resulting in substantial changes in management advice with only small changes in negative log likelihood. For Model 22.2 there was a $\sim 150,000$ ton range in 2023 ABC with a less than 10 -LL change from the MLE in likelihood profile run over catchability.

For 2023 we conducted a series of model explorations in an attempt to fix these issues.. After initial investigations to move to a more standardized way of setting input sample sizes it became apparent that a different approach was needed for model exploration as the updated model did not consistently converge on the MLE and the complex base model made investigations difficult. We chose to move to a more simplified model and then add components to the model sequentially to evaluate the impacts of different model assumptions. All models examined in this paper are built in Stock Synthesis version 3.30.21 and parameterized the same as Model 22.2 (Barbeaux et al. 2022) except for changes specified. For the simplified model we reverted to a standard multinomial for the composition data and removed all time varying components, we also fixed aging bias based on previous model results and in line with
isotope analyses validating aging methods (Kastelle et al. 2017). A full description of changes made to simplify the model is provided below. For model explorations we examined a wide range of model alternatives however for this analysis we have limited the changes to the impacts of allowing growth to be time varying, allowing survey selectivity to be time varying, reducing the maximum age from 20 to 12 , adding catch data from 1964 to 1976 while removing the regime change parameter on recruitment, and adding conditional age-at-length. Although all the models were generally well fit, the results in terms of both stock size and management implications from these models remained highly variable and individual models remained sensitive with small changes in fit resulting in large changes in management advice. The models explored displayed notable variation in survey catchability spanning a range from 0.74 to 1.10 and was highly negatively correlated ( $\mathrm{R}^{2}=-0.92$ ) with natural mortality ranging from 0.33 to 0.44 . Likelihood profiles on catchability showed that for many of the seemingly reasonable individual models $A B C$ recommendations could vary by more than 100,000 $t$ with changes in log likelihood of less than 2 points. A major finding therefore of this work is that for Bering Sea Pacific cod very disparate outcomes in terms of management advice could be generated from models with very little difference in fit to data and retrospective bias.

All model files are provided here:

## https://afsc-

assessments.github.io/EBS PCOD/2023 ASSESSMENT/SEPTEMBER MODELS/APPENDICES/September Models.zip

For 2023 the authors recommend the following:

1) Moving away from the ensemble approach currently employed. The authors believe that the current ensemble of models are too similar in nature. A better ensemble approach would be to include models with more varied structures such as the multispecies model (CEATTLE) and simplified random effects models. This effort would require a much larger team of researchers to evaluate individual model performance. As it stands the evaluation of individual model performance in the ensemble may have been hindered by the volume of work required of a single author to complete an ensemble of models given their unstable nature in the limited time between the September and November Plan Teams. In addition it is the opinion of the assessment authors that if the ensemble approach is to continue the SSC should consider setting up a working group outside the normal Plan Team process to develop a science-based and transparent model selection and weighting scheme for new ensembles.
2) Changing the current modeling approach to a simpler model with fewer and/or more constrained, annually varying parameters on growth and selectivity as these parameters are confounded. Generally we add this type of complexity to deal with poor residual behavior and retrospective bias issues, here the residuals of the complex models don't appear to differ substantially from the more complex models and retrospective patterns are within reasonable bounds even with the simplest models considered.
3) Fixing one or more key parameters (e.g. $M, Q, L_{\text {min }}, L_{\text {max }}, K$, etc.) in the model or using more constrained priors would provide improved model stability. Constraining natural mortality or growth parameters would be better candidates than survey catchability as more refined aging methods developed recently should better inform these parameters. Either fixing or providing a
strong prior on natural mortality would have the added effect of constraining catchability within a model as these parameters are highly correlated.

If a single model is to be used for management this year, of the models presented the authors would recommend Model 23.1.0.d be considered as an option for further exploration for management in 2024. Model 23.1.0.d is the simplified model with the addition of constrained annual variability in growth (Figure 1 and Figure 2 ) and survey selectivity (Figure 3 and Figure 4). The model employs bootstrapped input sample sizes, the fit to the survey index is improved over Model 22.2 (Figure 5), retrospective bias remains within acceptable limits, and residual runs analyses imply that the fit to the composition data are better with less autocorrelation in the residuals while maintaining similar effective sample sizes. The mean absolute scaled error (MASE) analyses show the predictive skill of Model 23.1.0.d for the survey index was improved over Model 22.2 ( 0.40 vs 0.69 ) and remains similar for the fishery mean length and survey mean age.

Model 23.1.0.d has improved performance while reducing the model by 86 parameters, however this reduction in complexity comes at a cost of increased uncertainty (higher standard deviations) in some key parameter estimates ( $M, Q$, and $\mathrm{R}_{0}$; Figure 6) which translates into increased uncertainty in derived quantities such as $\mathrm{B}_{0}, \mathrm{~F}_{40 \%}$, current recruitment (Figure 7), current spawning biomass (Figure 8 and Figure 9), and future catch recommendations. Another potential red-flag in Model 23.1.0.d is that natural mortality at 0.429 is higher than most methods external to the model indicate and catchability at 0.765 is lower than most other models examined previously. Profiles over catchability show little change in likelihood over a wide range of natural mortality and catchability suggesting little information in the data to inform these quantities (Figure 10).

One solution may be found in applying a maximum age-based method for deriving a prior for natural mortality (Sulliven at al. 2022; Thorson et al. 2023), which suggests a lower value with $M$ at 0.387 . When natural mortality is fixed in Model 23.1.0.d to this value, catchability increases to 0.972 and may be a reasonable alternative to allowing natural mortality to be freely fit. Fixing $M$ in this model has a minor cost of degrading the overall model performance by +1.4 negative log likelihood (-LL) with an improvement to the fit to the age composition of $-3.95-\mathrm{LL}$, but a poorer fit to the survey index at +2.56 LL and length composition at $+2.40-\mathrm{LL}$.

## 2 Model 22.2 updated changes in input sample size

Hulson et al. (2023) found that there was not a consistent approach to setting input sample sizes for composition data in assessment models at the Alaska Fisheries Science Center. They proposed a unifying bootstrap approach that would evaluate the variance and autocorrelation within the survey composition data collections to appropriately calculate annual input sample sizes. For the 2022 Pacific cod ensemble models the input sample sizes for the survey size and age composition data were set at the number of surveyed hauls for each year, and the fishery size composition data were set at the number of hauls sampled standardized to the mean sample of hauls from the survey over all years surveyed which is unique to Eastern Bering Sea Pacific cod. This method led to an average input sample size of 369 for both survey and fishery length and age compositions (Table 1). As noted the ensemble models were fit using the Dirichlet-multinomial (DM), which as coded in Stock Synthesis uses a parameter (log theta) to re-weight the data and in effect reduce the size and age composition input sample sizes, if needed, to appropriately weight data components within the model. In all of the 2022 ensemble models the DM log theta parameters approaches the upper bound for both the fishery and survey size composition data and must be fixed in order for the models to converge. Having the DM theta parameter fixed at the
upper bound is not optimal and may indicate that input sample sizes for the size composition data are too small and therefore underweighted in the model compared to other data components. Using a bootstrap approach (Hulson et al. 2023) for calculating input sample size for the survey length and age composition data resulted in an on average smaller age composition sample size of 250 (in agreement with the fitted DM theta value of -0.47 ) and a much larger on average input sample size of for the size composition data of 1661 (Table 1). A bootstrap approach is not yet available for the fishery composition data and therefore for the fishery size composition data input sample size we used the annual number of hauls sampled standardized to the mean survey size composition input sample size so that both means were equal for the two size composition data sets. As in previous years it was assumed that the raw numbers of hauls were far too high as they numbered in the tens of thousands for some year, far higher than the survey input sample size. Model 22.2 was then fit using these new sample sizes. Although not useful for judging differences in model fits the overall negative log likelihood (-LL) increased from 10,875 to 18,362, with a sharp increase in the length composition -LL from 9,990 to 17,383 which does indicate a shift in the model weighting with more weight being given to the composition data. The DM log theta values for the survey age composition data changed from -0.47 to 0.60 and the survey size composition changed from near the bound at 10 in the old model to 1.32 in the updated model. However the fishery size composition DM log theta remained at the bound suggesting the sample sizes remained too low in comparison with the other data components or potentially an additional issue with model misspecification. The change resulted in substantially more weight on the composition data than in the old Model 22.2 and a degradation in fit to the survey (Table 2 and Table 3) from -6-LL to 68 -LL. Retrospective and mean absolute scaled error (MASE) values were the same between the two model configurations (Table 4) and the fishery mean length residuals remained significantly autocorrelated as determined by the residual runs test (ss3diags; Winker et al.2023). However, the change in input samples sizes resulted in better residual behavior for the survey mean length and age (Table 5). Convergence was impacted with a large number of jitter runs failing to converge at the MLE for the updated model. The profile over catchability for the updated Model 22.2 (Figure 1) shows a highly irregular profile resulting from the models not converging to the MLE for each of the fixed catchability value runs. In conducting likelihood profiles over catchability for both the old and updated Model 22.2 survey catchability ( Q ) and natural mortality ( M ) are strongly negatively correlated $\left(R^{2}=-0.999\right)$ with a slightly higher $M$ (Table 10) in the model with updated input sample size. The updated Model 22.2 growth parameters were different from the old Model 22.2 driven entirely by the different input sample sizes (Table 7) and changes in relative weighting of the data components. However, changes in growth parameters had little impact on the overall size at age (Figure 1). Additional changes in influential parameters are shown in Table 8. Despite changes in important parameters such as catchability and natural mortality both model configurations resulted in similar estimates of recruitment over time (Figure 7), spawning stock biomass over time (Figure 8 and Figure 9), reference points (within 5\%; Table 9), and management advice (2024 ABC within 2\%; Table 9).

We explored changing from the DM to standard multinomial and implementing the Francis TA1.8 weighting method for Model 22.2 to improve model performance. However, we found that when iteratively fitting using Francis TA1.8 methods the model the suggested correction to the fishery size composition value continued to increase without settling until the model no longer converged. This suggests model misspecification in Model 22.2, potentially due to the same issue contributing to the DM log theta approaching the upper bound for the fishery length composition data despite rather large input samples sizes. We theorized that the model misspecification could be due to confounding among the freely fit aging bias, annually varying growth, and annually varying selectivity.

## 3 Model 23.1.0.a description

To allow an easier understanding of the interaction of model components on model results and model sensitivities we created a simplified version of Model 22.2 (Barbeaux et al. 2022). The new simplified Model 23.1.0.A had the following changes:

1. Removing length composition data for years with age composition data (1994-2021) which were duplicated in the age comps.
2. Reconfiguring both survey and fishery selectivity to be static instead of including annually varying parameters.
3. Reconfiguring the Richard's growth to be static instead of including annually varying $L_{\text {min }}$.
4. Reconfiguring the survey double normal selectivity (Stock Synthesis pattern 24; Methot et al. 2023) to estimate parameters 1-4 and using new asymptotic option for parameter 6.
5. Fixing the pre-2007 aging bias to Model 22.2 values.
6. For the growth model fixing CV at older ages at 0.06 and fixing CV at younger ages at 0.2 based on the previous ensemble model fits.
7. Changing from the Dirichlet-multinomial to standard multinomial for length and age composition data.
8. Using the iterative Francis TA1.8 weighting method to tune the model.

### 3.1 Duplicate composition data

For the 2022 ensemble models both survey and age composition are included for all years in which they are available, resulting in 1994-2021 having both survey age and survey size composition data included in the model. Therefore the survey composition data for these years are potentially more highly weighted in the models than other data components. In the exploratory models for the years with bottom trawl survey age composition data available (1994-2019, and 2021) the bottom trawl survey size composition data were removed.

### 3.2 Selectivity and growth

For the 2022 ensemble models both fishery and survey selectivity and $L_{\text {min }}$ in the Richard's growth model were set to be annually varying. This is likely confounded as the model would likely not be able to discern between annually varying growth and selectivity at the smaller sizes potentially leading to issues with model convergence and inability to settle on appropriate Francis weighting for the fishery length composition data. This may have also led to issues with the DM log theta approaching the upper bound for these data. Whether the annual variability is attributed to growth or selectivity has impacts on model results affecting management advice. For the simplified model, Model 23.1.0a, we set both growth (Figure 2) and selectivity (Figure 3) to be static over time reducing the number of devs by 220 (Table 2).

In addition, we implemented a selectivity feature new to stock synthesis for the survey that simplifies the double normal function (selectivity option 24 in Stock Synthesis; Method et al. 2023) where the values past a set length are meant to be static, here we set all selectivity values at lengths greater than 40 cm to be fixed. It should be noted that although this feature was meant to fix the survey selectivity to be asymptotic and not allow the shape to become dome-shaped, this was found not to be the case when conducting profiles over catchability. In cases where catchability was high and natural mortality was very low, the shape of the survey selectivity curve did unexpectedly become dome-shaped.

### 3.3 Aging bias

Aging bias was fit for all of the 2022 ensemble models as a two parameter linear vector from ages 2 to 20. The two parameters are the aging bias at age 2 and at the max age. These two parameters tended to vary considerably depending on assumptions of growth and selectivity as model configurations were explored leading to some fits that were improbable suggesting that these parameters were likely confounded with growth and selectivity. Changes in estimated aging bias had substantial impacts on model results and some fits were well outside what would be expected given isotope analysis (Kastelle et al. 2017). For the models explored this year in order to stabilize model explorations we fixed the two parameters based on the 2022 Model 22.2 accepted values.

### 3.4 Composition distribution from Dirichlet multinomial to standard multinomial

As described above when fitting Model 22.2 with the updated bootstrap input sample sizes the fishery size composition DM log theta continued to approach the upper bound, which in effect reverts the distribution to the standard multinomial. Although it has been common practice to fix the log(theta) parameter near the upper bound when this occurs, the fit may indicate that the input sample sizes continue to be inadequate or some other model misspecification.

In order to investigate this phenomenon and evaluate other options, we changed the presumed distribution of the composition data to the standard multinomial. We then iteratively adjusted the model as per the Francis reweighting scheme TA1.8 (Francis, 2011) as implemented in the R library r4ss (Taylor et al. 2021), a technique previously utilized in Pacific cod models prior to 2018.

### 3.5 Model 23.1.0.a Results

Model 23.1.0.a performed well overall with standard metrics for model fits similar and in some cases improved over the more complicated Model 22.2 with substantially fewer fit parameters ( 82 vs. 306). Iterative Francis reweighting for Model 23.1.0.a settled on consistent values and resulted in considerable down-weighting of the length and age composition data from the initial input sample sizes with a multiplier values of 0.03 and 0.06 for the fishery and survey length composition data and 0.25 for the survey age composition data. This resulted in the survey index having more influence on the model then in the 2022 ensemble models. This is evidenced with the improved likelihood and RMSE on the survey abundance index fit (Table 3 and Table 4 ) and shown as a much tighter fit of the model to the survey abundance index (Figure 5). Due to the change to the Francis weighting versus the DM between the two models, no direct comparison of likelihoods can be used for comparison, however looking at the effective sample size shows a drop in the effective sample size between the updated Model 22.2 and Model 23.1.0.a for the fishery size and survey size and age compositions (Table 4). The visual fits to the fishery length composition data reveal little difference in the fitted values (Figure 11) or residuals (Figure 12). However, Model 22.2 fits the survey size composition better, specifically Model 23.1.0.a tends to overestimate large incoming small fish even more than Model 22.2 when large recruitments are present (Figure 11). Visual inspection of the age composition fits (Figure 13) show Model 23.1.0.a fitting the age data less closely than Model 22.2 with lower effective same size, however visual inspection of the age composition Pearson residuals from both models shows similar patterns (Figure 14).

The retrospective (Table 5) and residual runs tests (Table 6) results were similar between updated Model 22.2 and the simplified Model 23.1.0.a with Mohn's rho retrospective values on spawning bias at 0.07 and passing runs test for all but the fishery size composition data component for both models. Mohn's rho tests show a small positive bias (0.08) for Model 23.1.0.a while the updated Model 22.2 had
a slight negative bias (-0.06). Examination using the mean absolute scaled error (MASE) provided in the ss3diags R Package (Winker et al. 2023) showed a marked improvement in the prediction skill of Model 23.1.0.a over the updated Model 22.2 for the survey index (Table 5), a slight improvement for the prediction skill of mean survey age, and a slight degradation for the prediction skill of the mean fishery length.

Despite being different from the updated Model 22.2 the growth parameter estimates between the old Model 22.2 and Model 23.1.0.a are similar (Table 6), however the standard deviation of the parameter estimates for $L_{\text {max }}$ and the Richard's parameter are nearly double in the simpler model. Similar fitted values are expected given the down-weighting of the composition samples through the Francis reweighting method and lower sample sizes in the old Model 22.2 compared to the updated model. The increase in the variance of these parameters in Model 23.1.0.a over Model 22.2 was due to removal of annual variability in growth and selectivity where some of the variability was attributed to the annual devs.

Model 23.1.0.a results in an increase in the estimated catchability over the update Model 22.2 to 1.097 from 0.974. This has the impact of scaling down recruitment (Figure 7) and spawning stock biomass overall (Figure 9). For Model 23.1.0.a both the jitter analysis and likelihood profile over catchability were well behaved with all jitter runs converging and the majority arriving at the MLE. Although the likelihood profile over survey catchability for this model is well behaved it shows very little change in the overall likelihood over a wide range of survey catchability values (Figure 10 and Table 11). For catchability ranging from 0.9 to 1.28 there is a change in negative log likelihood of less than 2 -LL from the maximum likelihood estimate (MLE) resulting in a 100,000 t difference in 2023 recommended ABC across that range. This is of similar concern in Model 22.2. This may be an issue with environmentally driven fluctuations in growth and recruitment contributing substantially to the overall biomass's variability, leading to limited insights into the consequences of fishery removals. The ability of a model to fit catchability is influenced by the degree to which catch impacts changes in survey abundance. Given the considerable impact of environmental drivers on cod abundance and mortality, there's a possibility of insufficient data for accurately determining survey catchability without a better understanding of the environmental drivers of this stock.

## 4 Further model explorations

Alternative models with increasing complexity were developed to further explore model sensitivity and performance with the following:

1. Allowing annual variability in both the $L_{\text {min }}$ and Richards $K$ parameters.
2. Adding constrained annually varying selectivity for the survey.
3. Integrating catch data from the period 1964-1976 into the model while eliminating the recruitment adjustment parameter linked to the 1977 environmental regime change.
4. Adjusting the upper age group for model dynamics from age 20 to age 12 to more accurately represent available data and speed up model runs.
5. Introducing survey conditional age-at-length (CAAL) data.

| Model | Npar. <br> +Ndevs | Annually <br> varying growth | Annually varying <br> survey selectivity | Max age <br> to 12 | Catch to 1964 no <br> regime | CAAL |
| :---: | ---: | :---: | :---: | :---: | :---: | :---: |
| $\underline{23.1 .0 . \mathrm{a}}$ | 82 |  |  |  |  |  |
| $\underline{23.1 .0 . \mathrm{b}}$ | 176 | X |  |  |  |  |
| $\underline{23.1 .0 . \mathrm{d}}$ | 218 | X | X |  |  |  |
| $\underline{23.1 .0 . \mathrm{g}}$ | 217 | X | XX | X | X |  |
| $\underline{23.1 .0 . \mathrm{h}}$ | 217 | X | X | X | X | X |

### 4.1 Model 23.1.0.b

### 4.1.1 Annually varying growth

For the 2022 ensemble models all growth was fit as a 4 parameter Richard's growth relationship with $\mathrm{L}_{\text {min }}$ fit as an annually varying deviation. All parameters in the 2022 models were fit with an uninformative prior. For the simplified Model 23.1.0.a although the four parameters were fit within the model with uninformative priors both growth and selectivity were set to be time-invariant. It has been long understood that environment, particularly temperature, is influential in the growth of Gadus species (Taylor 1958) and annual variability in growth should be expected. Growth in Pacific cod specifically has been found to be rather elastic and dependent on environmental conditions particularly for young fish (Laurel et al. 2008, Barbeaux et al. 2021). To evaluate this elasticity we explored including annually varying growth in Model 23.1.0.b. Here we used the growth parameters and standard deviations as posteriors from Model 23.1.0.a (Table 7) as 'priors' for all four growth parameters and added a mean tending random walk for $L_{\text {min }}$ and Richard's parameter. This is option 5 described in the Stock Synthesis manual (Methot et al 2023) as a mean reverting random walk with rho and a logit transformation to stay within the minimum and maximum parameter bound. In developing Model 23.1.0.b allowing all the growth parameters to vary annually was evaluated, however only varying $L_{\text {min }}$ and the Richards parameter provided substantial improvements to the model fit. The authors understand that using the values from Model 23.1.0a are not true priors, but given their large standard deviations it was considered only as a means of providing soft bounds on the parameters and kept the parameters from wandering to extreme values during fitting. The standard deviation of the devs were tuned following the methods of Thompson et al. (2021) where the standard deviation was tuned to set the variance of the estimates plus the sum of the estimates' variances equal to unity. This resulted in standard deviation of 0.4416 for $L_{\text {min }}$ and 0.2995 for the Richards parameter annual deviations.

### 4.1.2 Results of adding annually varying growth

Model 23.1.0.b performed well overall with standard metrics for model fits improved over Model 23.1.0.a. The results of allowing annually varying growth (Figure 2) was an overall improvement to fits to all data components over the static growth model with lower negative log likelihood across all data components, lower RMSE for the survey index, and higher effective N for all of the composition data (Table 3, Table 4, and Figure 5). As was expected with the use of the 'prior' and annual variability the standard deviations of the growth parameters were also reduced (Table 7) from Model 23.1.0.a. Retrospective bias remained similar to Model 23.1.0.a with a small positive bias for spawning stock biomass (Table 5). The MASE evaluation showed improved predictive skill on the index and size composition and a slight degradation in the predictive skill on age composition (Table 5). Residual runs tests resulted in an improvement for the fit to the survey index and both survey and fishery length composition fits, but a slight degradation in the survey age composition fit from Model 23.1.0.a.

However, unlike all the previous models described, all data components passed the residuals runs test (Table 6). Although fits to the length composition (Figure 15) and age composition (Figure 13) were discernably better than Model 23.1.0.a, patterns in the residuals for both length (Figure 15) and age (Figure 14) composition were visually similar.

While adding annually varying growth improved model fit, it also increased uncertainty in model estimates of management reference points (Table 8 and Table 9), annual estimates of recruitment (Figure 6 and Figure 7) and annual estimates of spawning biomass ( Figure 8 and Figure 9). There was an overall increase in uncertainty in model results compared to Model 23.1.0.a and Model 22.2. The estimate of bottom trawl survey catchability was lower and natural mortality higher in Model 23.1.0.b than in Model 23.1.0.a (Table 8). However, the likelihood profile on catchability for Model 23.1.0.b continues to show low certainty in this value as large changes in catchability continue to result in small changes in likelihood (Table 11 and Figure 10).

It is interesting to note that in likelihood profiles over catchability, natural mortality is always highly negatively correlated with catchability, however in the new models growth and selectivity parameters appear to have tipping points where the trajectory of the parameter values with catchability change suddenly (Figure 16). When examining the likelihood profiles by model component this manifests as a change in the trajectory of the index data likelihood (Figure 10). This tipping point becomes even more pronounced when annually varying growth is introduced with priors on the main growth parameters. In the 23.1.0.x series models, lower natural mortality, correlated with higher catchability values above 1.0, are compensated for by forcing the selectivity curve to be increasingly dome-shaped (Figure 17) making a subtle trade-off of fit between the survey age and size composition data and the survey index. Specifically, this trade-off is made for the initial year 1982, 1994, and 2001 where the survey abundance appeared to be higher than expected (Figure 5), out of line with adjacent years, and poorly fit in all models. This tipping point is due to the parameterization of the survey selectivity in which parameter 6, selectivity at last bin, is set to - 1040 which was meant set selectivity constant for bins greater than bin number 40, but appears to allow selectivity become dome-shaped in some cases.

An increase in catchability and decrease in natural mortality translated into a scaling up of the population with higher estimates of annual recruitment (Figure 7) and annual spawning biomass (Figure 9). Although unfished spawning biomass ( $B_{0}$ ) increased over Model 23.1.0.a (Table 2) it did not increase proportionally to the annual spawning biomass and remained well below that estimated in Model 22.2. This resulted in an overall increase in the estimated status of the stock to over $\mathrm{B}_{50 \%}$ in 2023 compared to Model 23.1.0.a and Model 22.2, where it was estimated to be below $\mathrm{B}_{40 \%}$.
4.2 Model 23.1.0.d
4.2.1 Annually varying survey selectivity

In all of the 2022 ensemble models both survey and fishery selectivity was modeled as annually varying. This variability was removed for the 2023 simplified model but reinstitution of annually varying selectivity on survey selectivity was examined (Figure 4). The general parameterization of selectivity remained the same with a six parameter double normal as described for the Model 23.1.0.a, however an annual additive deviation (Stock Synthesis option 2; Methot et al. 2023) was added to the ascending width of the curve with a standard error of the deviation restricted to 0.2 , this added 42 dev. parameters to the model, one for each survey year. Although the apparent variation in the fitted curves was minor (Figure 4), the additional flexibility improved the model fit by -10.19-LL. We had examined
models with annually varying selectivity in the fishery as well, however adding annually varying fishery selectivity made only a modest improvement in model fit over a Model 23.1.0.b with a substantial increase in the number of annual dev. parameters ( 141 additional dev parameters gaining a 4 point reduction in -LL). For brevity the model with fishery annually varying selectivity was dropped from consideration and not fully presented in this document.

### 4.2.2 Results of adding annually varying survey selectivity

The addition of restricted annually varying selectivity for the bottom trawl survey improved the overall fit compared to Model 23.1.0.b (-10.19-LL). The largest changes were an improved fit to the survey abundance index ( $-5.49-\mathrm{LL}$ ), and both length composition data sets ( $-10.17-\mathrm{LL}$ ) and a degraded fit to the survey age composition data ( $+2.21-\mathrm{LL}$ ) (Table 2, Table 3, and Table 4). Retrospective Mohn's rho and MASE values were the same as Model 23.1.0.b (Table 5). All modeled components passed the residual runs tests (Table 6) with the same p-values for the index and fishery length composition and slightly lower $p$-values for the survey length and age composition (Table 6). There was little difference in the growth parameter estimates with nearly identical standard deviations (Table 7). There was an increase in the estimated $\mathrm{R}_{0}$ and natural mortality and a decrease in the survey abundance index catchability with standard deviation remaining near those of Model 23.1.0.b (Table 8). Similar to all of the models examined the log likelihood profile over catchability again showed the model to have low certainty in this value as large changes in catchability continue to result in small changes in likelihood (Table 12 and Figure 10).

The increase in catchability and decrease in natural mortality once again translated into a scaling up of the population from Model 23.1.0.b with higher estimates of annual recruitment (Figure 7) and annual spawning biomass (Table 2, Table 9, and Figure 9). Although this model is very near in fit to Model 23.1.0.b the management advice provided would have increased the projected 2024 ABC by approximately 24 thousand tons.

### 4.3 Model 23.10.0.g

4.3.1 Catch data 1964-1976

Regarding the ensemble models for 2022, the catch series begins in 1977, and there is a presumed shift in the climate regime in the same year, positively influencing recruitment of Pacific cod from that point onward. The steady-state catch level for all four ensemble models was set at 42,500 tons, reflecting the average catch from 1964 to 1976. Examining the catch data available for the years 1964 to 1976 (Table 12), it's evident that the catch fell notably below this average prior to 1967. Despite the considerable catch recorded during the 1920s and 1930s, anecdotal evidence suggests that the catch levels from the 1940s to 1967 were lower compared to that of 1968 onward (Mackovjak, 2019). To test the influence of this initial catch on management reference points we added the 1964-1976 data and set an equilibrium catch value to a low $10,000 \mathrm{t}$, an approximation of potential catch levels prior to the development of the modern fishery in the late 1960's.

The notion of a regime change in the North Pacific, leading to altered recruitment patterns in groundfish, was initially proposed by Francis et al. in 2003. While the climatic regime shift of 1976-77 is well-documented (Hare and Mantua, 2000), the sustained and consistent positive influence of this warmer regime on Pacific cod recruitment lacks comprehensive documentation. Consequently, we undertook an investigation encompassing models that incorporate the catch data from 1964 to 1976, excluding the regime change parameter, and assuming an equilibrium catch of 10,000 tons. This
exploration aimed to assess the sensitivity of the reference points to these initial assumptions regarding the impacts of the regime change on Pacific cod recruitment and 10,000 tons was a best guess rough estimate of catches prior to 1964.

### 4.3.2 Maximum age from age 20 to age 12

In 2022, the age plus group for model dynamics was maintained at 20 across all four ensemble models. This decision was made despite the fact that only one Pacific cod was observed in the Bering Sea shelf survey since 1993 with an age greater than 14, and only 34 fish were aged above 12 out of a total of 32,050 ages recorded. Starting from 2017, only two fish aged over 10 were identified out of a pool of 5,524 total ages collected from the bottom trawl survey. It's worth noting that due to processing limitations, the age composition analysis from the VAST survey had to be confined to an age 12 plus group.

In the near future, a shift in aging techniques is anticipated, moving towards the utilization of Fourier transform near-infrared spectroscopy (FT-NIRS) as detailed by Benson et al. (2023). Early findings pertaining to Pacific cod indicate notable discrepancies in age predictions beyond age 12 using FT-NIRS (communicated by Helser). Given these circumstances, we undertook an assessment to gauge the model's responsiveness to the transition to an age 12 plus group for model dynamics.

### 4.3.3 Results of changing catch years and decreasing the maximum age to 12

Two bridging models had been completed treating these two changes separately, but for brevity we chose to only present the combined model, Model 23.1.0.g. Neither of the changes to the model made much difference in the overall fit to the data (Table 2) with only a minor degradation in overall fit (+4.9LL ). All three composition data components were only minutely impacted negatively with these two changes ( $+6.1-\mathrm{LL}$ ) offset slightly by an improvement to the survey index ( $-1.1-\mathrm{LL}$; Table 3 and Table 4).

Retrospective Mohn's rho for Model 23.1.0.g, although slightly positively increased over that of Model 23.1.0.d, remained within acceptable bounds at 0.11 (Table 5). The MASE analysis showed a slight improvement of predictive capability for the survey index and a slight degradation in predictive capability for all of the composition data sets (Table 6) consistent with the model component specific likelihoods, RMSE, and effective N results. These slight changes within the model did however impact survey catchability increasing it to 0.79 from 0.77 of Model 23.1.0.d. This increase in catchability was accompanied by an increase in natural mortality to 0.435 from 0.429 , contrary to the trend of natural mortality decreasing with increasing $Q$. The log equilibrium recruitment $\ln \left(R_{0}\right)$ also increased to 13.74 from 13.69 consistent with the increase in natural mortality. A likelihood profile over survey catchability indicated that these model alterations did not improve fitting catchability as large changes in catchability continued to result in only small changes in likelihood (Table 13 and Figure 10).

The main difference in growth was an increase in $L_{\max }$ over Model 23.1.0.d due to the switch to the lower maximum age (Table 7). There was also a sharp increase in size and weight at the $12+$ group and somewhat longer fish at ages 5 and 6 for the terminal year (Figure 1). The weight-at-age was nearly identical to Model 23.1.0.b (Figure 1) which is slightly lighter at age than Model 23.1.0.d. The trend in annually varying growth remained similar to Model 23.1.0.d despite restricting the age to 12+ (Figure 2) with a stable trend for young fish ages 0 to 6 , increasing trend for fish ages 7 to 11, and a decreasing trend in size for the age 12+ group. The annual growth pattern in Model 23.1.0.d was consistent except the recent trend in increasing size was apparent through age 15 before turning to a recent decreasing
trend for ages 16 to 20. This difference in the trend by ages between Model 23.1.0.g and Model 23.1.0.d was due to the shortening of the modeled ages. This did not substantially impact the trend in spawning stock biomass with only slightly smaller values ( $\sim-3 \%$ ) after 1982 (Figure 9 and Table 9) consistent with the slightly higher survey catchability.

There was a substantial shift in the $B_{0}$ reference point from 623 kt from Model 23.1.0.d to 543 kt in Model 23.1.0.g (Table 9) and therefore despite the small decrease in overall female spawning biomass (Figure 9) the status of the stock in 2023 was higher at $\mathrm{B}_{61 \%}$ than the Model 23.1.0.d value of $\mathrm{B}_{55 \%}$ (Table 2 and Figure 8). The change in $\mathrm{B}_{0}$ can be attributed to the extension of the catch time series and lower equilibrium catch. There was also a reduction in the uncertainty of $B_{0}$ to a CV of 0.057 from 0.087 from Model 23.1.0.d, however the uncertainty around the current spawning biomass and ABC projections remained the same (Table 9). $\mathrm{F}_{40 \%}$ also increased from 0.47 to 0.49 from Model 23.1.0.d to 23.1.0.g due to the addition of the 1964-1976 catch and removal of the regime change parameter. Despite this increase in allowable F the 2024 ABC projection was reduced from 244kt to 239 kt due to lower spawning biomass in Model 23.1.0.g (Table 2 and Table 9). Therefore both changes resulted in minor adjustment in management advice.

Model 23.1.0.g continues to have the same issue as all the other models examined, changing the catch start year and reducing maximum age did not improve the profile on catchability. The log likelihood profile over catchability still showed the model to have low certainty in this value as large changes in catchability continue to result in small changes in likelihood (Table 13 and Figure 10). Catchability can be changed between 0.6 and 1.2 and still be within $\pm 2$-LL from the MLE, resulting in a range in management advice for the 2024 ABC of $\pm 100,000 t$ from the MLE.

Changing to the lower maximum age for the data will likely need to happen as the AFSC switches to FTNIRS aging however changes within the model dynamics as explored here will not. Impact to the model fit and results looks to be minor with a slight increase in $L_{\text {max }}$ and allowable fishing mortality. Including the earlier catch series may not be as clear and further work should be conducted to investigate the influence of these earlier catches and assumed equilibrium catch level. There is work currently underway by Dr. Catherine West, a zooarcheologist from Boston University, through a National Science Foundation grant (NSF award \# 2220552) to better account for Pacific cod fishery catch from pre-1964 which may better inform the equilibrium catch level used in the model.

### 4.4 Model 23.1.0.h

### 4.4.1 Survey conditional-age-at length

Annually varying growth in the 2022 ensemble models is driven by length and age composition data. In one set of alternative models we explored the inclusion of survey conditional-age-at-length (CAAL) to determine if this improved model estimates of annually varying growth (Figure 20). We employed the same method for calculating CAAL and input sample sizes for the CAAL used in the Gulf of Alaska Pacific cod stock assessment and documented in Barbeaux et al. (2021). In theory the CAAL should provide an improved estimate of the growth parameters including annual devs. Note that because we changed the data in the model the total likelihood cannot be directly compared to previous models however the RMSE, effective N , and other individual likelihood components can be.

### 4.4.2 Results of adding survey conditional-age-at-length

The addition of the survey CAAL resulted in the fits to all be degraded overall (Table 3 and Table 4). Here we see an increase in the survey length composition of $+12-\mathrm{LL}$ and fishery length composition of $+15-\mathrm{LL}$ while the index had an increase of +9 -LL. There was a minor improvement to the age composition data with the addition of the CAAL 76.74 in Model 23.1.0.g to 75.45 in Model 23.1.0.h. Residual patterns remain similar to previous models (Figure 13, Figure 14, and Figure 21). Fits to the CAAL data were good for most years with an exception for 1992 and 1993 where the model predicted smaller fish at age than were observed (Figure 22). The Mohn's rho for spawning stock biomass showed an increase in positive retrospective bias over Model 23.1.0.g to 0.15. MASE evaluation of predictive skill were similar for the index and survey age, but somewhat worse for fishery length composition data. Model 23.1.0.h fails the residual runs test for both the fishery length composition and survey age composition suggesting significant autocorrelation in the residuals in fitting these data sets. A likelihood profile over survey catchability (Table 14 and Figure 10) shows little improvement with the introduction of CAAL continuing to generate small changes in likelihood over large changes in catchability. This is reflected in a large range in management advice as spawning biomass and ABC are scaled with catchability.

The main impact of the addition of CAAL in the model was to fit a smaller $L_{\text {min }}$, a higher $K$, and reduce the standard deviation of all the growth parameters (Figure 23 and Table 7), as well as a small reductions in uncertainty for $\mathrm{R}_{0}$, natural mortality, and catchability (Table 8). The reduction in variance in parameters translated into lower uncertainty in the derived quantities such as fishing mortality, unfished spawning biomass, spawning biomass, and projected ABCs (Table 9). The trends in annually varying length-at-age were similar to Model 23.1.0.d and 23.1.0.g, but with slightly lower interannual variability for those years (1990-2022) with CAAL data. At ages 3 to 11 there is an increasing trend in size-at-age from 2000 forward, and a decreasing trend in size-at-age at age 12+. Random walk devs on $L_{\text {min }}$ are relatively consistent for Models 23.1.0.g and Model23.1.0.h (Figure 22) with an overall increasing trend over the time series suggesting an increase in size of juvenile fish in the Bering Sea. Richards parameter impacts the rate of fish growth and is consistent between models prior to the introduction of CAAL from 19771986, after which the series diverges with low values for Model 23.1.0.h that then increase over time.

The forecast for Model 23.1.0.h was set to average biological parameters back to 1964, this resulted in some aberrant behavior for the forecasted weight-at-age resulting in these values being substantially higher than the other models (Figure 1). As projections had not been considered in developing these models, this issue went unnoticed until very recently and the authors have not had to time to correct this issue. This would impact model projections including the 2024 ABC which was incorrect and has been removed from all tables. This issue would not impact the time series fits or model performance.

In summary the addition of CAAL reduces uncertainty throughout the model, but at a cost of degrading fits to all of the other model components (Table 3 and Table 4).

## 5 A Case for Fixing Natural Mortality

One way that could be used to alleviate model sensitivities and that has been explored in the past is to fix key model parameters or provide informative priors, e.g. natural mortality or catchability at 'reasonable' values. It's crucial to emphasize that at its base, fitting catchability within the model is influenced by the degree to which catch impacts changes in survey abundance. The fluctuations in growth and recruitment due to environmental factors can significantly contribute to the overall biomass's variability independent of catch, leading to limited insights into the consequences of fishery
removals and therefore little information within the data to inform catchability. There has been substantial debate in the past over catchability with an equal amount of work going into studies to try to better understand this value, the results of which have been equivocal. Given the considerable impact of environmental drivers on cod abundance and mortality, there's a possibility of never having sufficient data to accurately determine survey catchability. Issues with aging Pacific cod in the past have made estimating natural mortality unreliable with estimates varying from 0.20 to 0.96 across the spectrum of Pacific cod stocks (Thompson 2018). However recent improvements in methods may provide a more reliable means of estimating natural mortality outside the model.

### 5.1 Estimating a life-history-based prior for natural mortality

The parameter M representing natural mortality is difficult to estimate in many stock assessment models. When total removals are fitted and information exists to estimate the fishing mortality rate, estimates of M are typically correlated with estimates of survey catchability, $q$, such that including a Bayesian prior on $M$ can provide information about population scale and resulting catch limits.

Substantial empirical and theoretical evidence suggests that natural mortality is lower for large bodied individuals (Andersen, 2019). Asymptotic body length L_inf is negatively correlated with the von Bertalanffy growth parameter k, such that these two growth parameters are sometimes used to predict M (Hoenig, 1983). In fact, the ratio M/k has erroneously been called a "life-history invariant" (Roff, 1984), despite theory suggesting that higher $\mathrm{M} / \mathrm{k}$ is associated with lower L_mat/L_inf (Beverton \& Holt, 1959). In particular, some taxa evolve behavioral and morphological defenses against predators (e.g., spines) that likely contribute to a lower M/k than otherwise expected (Thorson et al., 2014). These antipredator defenses may in some cases be evolutionarily conserved, such that a lower-than-expected $\mathrm{M} / \mathrm{k}$ for a related taxa will be informative when predicting the value of M from k for a given species. This intuition gives rise to taxonomic-nested linear mixed models or phylogenetic trait imputation, which have been used to impute missing values for natural mortality (Thorson et al., 2017), recruitment density dependence (Thorson, 2020), or other behavioral and ecological traits (Thorson et al., 2023).

As an alternative to estimating natural mortality from growth parameters, researchers have also compiled estimates of longevity from aged specimens, and research suggests that longevity-based predictions of natural mortality rate are more precise than growth-based estimates (Hamel \& Cope, 2022; Then et al., 2015). Longevity can be recorded either as the maximum aged specimen, or the average of the five maximum ages (Sullivan et al., 2022). However, developing separate estimators using longevity and growth parameters then results in multiple estimators for a given species (Sullivan et al., 2022), which presents a challenge in either selecting a single estimator or weighting alternative estimators within an ensemble (Cope \& Hamel, 2022).

As alternative to developing separate models using growth or longevity information, recent research has developed phylogenetic structural equation models, which can explicitly represent the dependency among multivariate trait data (Thorson et al., 2023; van der Bijl, 2018; von Hardenberg \& GonzalezVoyer, 2013). In particular, a user-friendly R-package phylosem can impute missing trait values jointly with estimating complex dependencies among traits (Thorson \& van der Bijl, In review). Research confirms that phylosem exactly replicates results from simpler models including structural equation models, phylogenetic linear models, and phylogenetic trait imputation (Thorson \& van der Bijl, In review).

Here, we fit a phylogenetic structural equation model (PSEM) to a high-quality database of independent estimates of natural mortality (Then et al., 2015). We specifically use a PSEM that specifies three linear associations $\log \left(L_{\text {inf }}\right) \rightarrow \log \left(t_{\max }\right), \log (k) \rightarrow \log \left(t_{\text {max }}\right)$, and $t_{\max } \rightarrow \log (M)$. A jackknife experiment confirms that this PSEM can explain nearly $50 \%$ additional variance relative to a conventional linear model when using growth parameters to predict natural mortality rate, while also providing a simple method to include both growth and longevity information in a single natural mortality estimator (Thorson, In review). We then use either the maximum specimen age, or the average of the maximum ages to predict natural mortality rate for Pacific cod in the eastern Bering Sea since 2008. Both longevity metrics result in the same value t_max=14 years, and this results in a predicted value $\mathrm{M}=0.3866$ and $\log$ standard deviation of 0.4 .

All of the models considered above were refit with the maximum age derived value of natural mortality of 0.3866 and presented in Table 15. In all of the newly developed simpler models the change in model fit by fixing natural mortality was minimal with likelihoods changing by +2 -LL or less. The fixing of natural mortality also resulted in improved retrospective runs as would be expected (Figure 25). Survey catchability fit in the models changed in the direction one would expect as it is negatively correlated with natural mortality, when the fixed $M$ value was higher than the fit value catchability decreased, when it was lower it increased (Table 16).

## 6 Recommendation

If a single model is to be used for management this year, of the models presented the authors would recommend Model 23.1.0.d be considered as an option for further exploration for management in 2024. Model 23.1.0.d is the simplified model with the addition of constrained annual variability in growth (Figure 1 and Figure 2 ) and survey selectivity (Figure 3 and Figure 4). The model employs bootstrapped input sample sizes, the fit to the survey index is improved over Model 22.2 (Figure 5), retrospective bias remains within acceptable limits, and residual runs analyses imply that the fit to the composition data are better with less autocorrelation in the residuals while maintaining a similar effective n . The mean absolute scaled error (MASE) analyses show the predictive skill of Model 23.1.0.d for the survey index was improved over Model $22.2(0.40$ vs 0.69$)$ and remains similar for the fishery mean length and survey mean age.

Model 23.1.0.d has improved performance while reducing the model by 86 parameters, however this reduction in complexity comes at a cost of increased uncertainty (higher standard deviations) in some key parameter estimates ( $M, Q$, and $R_{0}$; Figure 6) that translates into increased uncertainty in derived quantities such as $\mathrm{B}_{0}, \mathrm{~F}_{40 \%}$, current recruitment (Figure 7), current spawning biomass (Figure 8 and Figure 9), and future catch recommendations. Another potential red-flag in Model 23.1.0.d is that natural mortality at 0.429 is higher than most methods external to the model indicate and catchability at 0.765 is lower than most other models examined previously. Profiles over catchability show little change in likelihood over a wide range of natural mortality and catchability (Figure 10).

One solution may be found in applying a maximum age-based method for deriving a prior for natural mortality that provides a value of natural morality within reasonable bounds of what has been fit in the simplified models presented above. When natural mortality is fixed in Model 23.1.0.d to 0.387, catchability increased to 0.972 and provides a reasonable alternative to allowing natural mortality to be
freely fit. Fixing M in Model 23.1.0.d has a minor cost of degrading the overall model performance by only +1.4 negative log likelihood (-LL) with an improvement to the fit to the age composition of $-3.95-\mathrm{LL}$, but a poorer fit to the survey index at +2.56 LL and length composition at $+2.40-\mathrm{LL}$. The change provides a more stable model but makes strong assumptions on the value of natural mortality and by association survey catchability (Table 16).

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## 8 Tables

Table 1. Input sample sizes for composition data, the old based on survey haul numbers and 'New' on a bootstrap approach (Hulson et al. 2023).

|  | Fishery |  | Survey <br> Year |  |  |  | Old | New | Old | New Length | New Age |
| :---: | ---: | ---: | ---: | ---: | ---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1977 | 6 | 26 |  |  |  |  |  |  |  |  |  |
| 1978 | 10 | 42 |  |  |  |  |  |  |  |  |  |
| 1979 | 12 | 52 |  |  |  |  |  |  |  |  |  |
| 1980 | 12 | 53 |  |  |  |  |  |  |  |  |  |
| 1981 | 14 | 61 |  |  |  |  |  |  |  |  |  |
| 1982 | 7 | 30 | 481 | 2432 |  |  |  |  |  |  |  |
| 1983 | 26 | 112 | 476 | 1171 |  |  |  |  |  |  |  |
| 1984 | 31 | 135 | 479 | 2424 |  |  |  |  |  |  |  |
| 1985 | 46 | 203 | 364 | 897 |  |  |  |  |  |  |  |
| 1986 | 47 | 207 | 481 | 2139 |  |  |  |  |  |  |  |
| 1987 | 87 | 380 | 412 | 2104 |  |  |  |  |  |  |  |
| 1988 | 89 | 387 | 354 | 1650 |  |  |  |  |  |  |  |
| 1989 | 41 | 179 | 373 | 1176 |  |  |  |  |  |  |  |
| 1990 | 42 | 184 | 354 | 1226 |  |  |  |  |  |  |  |
| 1991 | 345 | 1506 | 400 | 1200 |  |  |  |  |  |  |  |
| 1992 | 340 | 1485 | 368 | 807 |  |  |  |  |  |  |  |
| 1993 | 201 | 880 | 451 | 813 |  |  |  |  |  |  |  |
| 1994 | 317 | 1383 | 360 | 1265 | 183 |  |  |  |  |  |  |
| 1995 | 344 | 1503 | 381 | 1999 | 174 |  |  |  |  |  |  |
| 1996 | 445 | 1943 | 368 | 1343 | 151 |  |  |  |  |  |  |
| 1997 | 472 | 2063 | 354 | 1389 | 98 |  |  |  |  |  |  |
| 1998 | 451 | 1972 | 360 | 2196 | 180 |  |  |  |  |  |  |
| 1999 | 600 | 2622 | 422 | 2078 | 224 |  |  |  |  |  |  |
| 2000 | 652 | 2849 | 363 | 1396 | 154 |  |  |  |  |  |  |
| 2001 | 692 | 3025 | 402 | 1829 | 304 |  |  |  |  |  |  |
| 2002 | 759 | 3318 | 366 | 2159 | 329 |  |  |  |  |  |  |
| 2003 | 947 | 4138 | 355 | 1040 | 265 |  |  |  |  |  |  |
| 2004 | 794 | 3471 | 336 | 1887 | 308 |  |  |  |  |  |  |
| 2005 | 761 | 3328 | 362 | 1164 | 212 |  |  |  |  |  |  |
| 2006 | 594 | 2595 | 369 | 2487 | 492 |  |  |  |  |  |  |
| 2007 | 466 | 2035 | 359 | 270 | 55 |  |  |  |  |  |  |
| 2008 | 551 | 2409 | 347 | 1757 | 235 |  |  |  |  |  |  |
| 2009 | 488 | 2134 | 364 | 908 | 201 |  |  |  |  |  |  |
| 2010 | 435 | 1902 | 363 | 1191 | 150 |  |  |  |  |  |  |
| 2011 | 572 | 2498 | 332 | 1398 | 127 |  |  |  |  |  |  |
| 2012 | 611 | 2670 | 330 | 865 | 150 |  |  |  |  |  |  |
| 2013 | 726 | 3171 | 329 | 909 | 149 |  |  |  |  |  |  |
| 2014 | 793 | 3467 | 293 | 1057 | 124 |  |  |  |  |  |  |
| 2015 | 733 | 3202 | 370 | 2068 | 362 |  |  |  |  |  |  |
| 2016 | 621 | 2715 | 339 | 3149 | 536 |  |  |  |  |  |  |
| 2017 | 544 | 2377 | 349 | 2802 | 447 |  |  |  |  |  |  |
| 2018 | 418 | 1827 | 369 | 2996 | 367 |  |  |  |  |  |  |
| 2019 | 301 | 1316 | 264 | 1230 | 250 |  |  |  |  |  |  |
| 2020 | 231 | 1008 | NA | 189 | 827 |  |  |  |  |  |  |
| Mean | 128 | 1115 | 255 | 320 | 2388 |  |  |  |  |  |  |
|  | 369 | 1626 | 369 | 1661 | 250 |  |  |  |  |  |  |
|  |  |  |  |  | 531 |  |  |  |  |  |  |
| N |  |  |  |  |  |  |  |  |  |  |  |

Table 2. Results from 2023 model exploration. The table shows Natural mortality (M),bottom trawl survey catchability ( $Q$ ), unfished female spawning biomass ( $B_{0}$ ), female spawning biomass in 2023 ( $B_{23}$ ), projected Allowable Biological Catch in 2024 ( $\mathrm{ABC}_{24}$ ), number of non-dev parameters (Npars), number of annual devs (Ndevs), and negative log likelihood (-LL) by model.

| M | Q | $\begin{gathered} B_{0} \\ \text { (kt) } \end{gathered}$ | Fmsy | $\begin{gathered} B_{23} \\ (k t) \end{gathered}$ | $\mathrm{B}_{23} / \mathrm{B}_{0}$ | $\begin{gathered} \mathrm{ABC}_{24} \\ (\mathrm{kt}) \end{gathered}$ | Npars | Ndevs | -LL | Model |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0.347 | 0.960 | 661.5 | 0.326 | 249.8 | 0.378 | 145 | 20 | 284 | 10875 | MODEL 22.2 old |
| 0.328 | 0.974 | 694.7 | 0.290 | 263.2 | 0.379 | 141 | 22 | 284 | 18362 | MODEL 22.2 updated |
| 0.344 | 1.097 | 586.1 | 0.332 | 205.9 | 0.351 | 132 | 18 | 64 | 251 | MODEL 23.1.0.a |
| 0.414 | 0.822 | 605.4 | 0.441 | 314.1 | 0.519 | 220 | 18 | 158 | 143 | MODEL 23.1.0.b |
| 0.429 | 0.765 | 623.4 | 0.465 | 343.4 | 0.551 | 244 | 18 | 200 | 133 | MODEL 23.1.0.d |
| 0.435 | 0.792 | 542.6 | 0.488 | 331.8 | 0.612 | 239 | 17 | 200 | 141 | MODEL 23.1.0.g |
| 0.424 | 0.808 | 611.4 | 0.466 | 313.1 | 0.512 |  | 17 | 200 | 631 | MODEL 23.1.0.h |

Table 3. Negative log likelihoods by data component and fleet.

| Model | Label | All | Fishery | Survey |
| :---: | :---: | :---: | :---: | :---: |
| MODEL 22.2 old | Age_like | 817.80 |  | 817.80 |
| MODEL 22.2 updated | Age_like | 766.34 |  | 766.34 |
| MODEL 23.1.0.a | Age_like | 88.62 |  | 88.62 |
| MODEL 23.1.0.b | Age_like | 71.07 |  | 71.07 |
| MODEL 23.1.0.d | Age_like | 73.27 |  | 73.27 |
| MODEL 23.1.0.g | Age_like | 76.74 |  | 76.74 |
| MODEL23.1.0.h | Age_like | 75.45 |  | 75.45 |
| MODEL 22.2 old | Length_like | 9990.5 | 4502.5 | 5487.98 |
| MODEL 22.2 updated | Length_like | 17382.5 | 7682.9 | 9699.66 |
| MODEL 23.1.0.a | Length_like | 184.38 | 79.03 | 105.35 |
| MODEL 23.1.0.b | Length_like | 130.75 | 60.29 | 70.46 |
| MODEL 23.1.0.d | Length_like | 120.58 | 59.78 | 60.81 |
| MODEL 23.1.0.g | Length_like | 123.19 | 60.46 | 62.73 |
| MODEL 23.1.0.h | Length_like | 150.98 | 76.04 | 74.94 |
| MODEL 22.2 old | Surv_like | -5.96 |  | -5.96 |
| MODEL 22.2 updated | Surv like | 67.53 |  | 67.53 |
| MODEL 23.1.0.a | Surv_like | -30.05 |  | -30.05 |
| MODEL 23.1.0.b | Surv_like | -83.13 |  | -83.13 |
| MODEL 23.1.0.d | Surv_like | -88.62 |  | -88.62 |
| MODEL 23.1.0.g | Surv_like | -89.74 |  | -89.74 |
| MODEL 23.1.0.h | Surv_like | -80.49 |  | -80.49 |
| MODEL 23.1.0.h | Survey CAAL | 445.62 |  | 445.62 |

Table 4. Root mean squared error RMSE and effective $N$ for data components by model. For the fishery length on Models 23.1.1.x gear splits are in order of Trawl/Longline/Pot

|  |  | Effective N |  |  |  |
| ---: | ---: | ---: | ---: | ---: | ---: |
| Index RMSE | Recruitment RMSE/SigmaR | Fishery Length | Survey Length | Survey Age | Model |
| 0.13 | 1.01 | 2919 | 852 | 168 | MODEL 22.2_old |
| 0.16 | 1.22 | 3474 | 929 | 122 | MODEL 22.2_updated |
| 0.12 | 1.06 | 1700 | 561 | 87 | MODEL 23.1.- |
| 0.07 | 0.81 | 2263 | 813 | 132 | MODEL 23.1.0.b |
| 0.07 | 0.77 | 2288 | 899 | 132 | MODEL 23.1.0.d |
| 0.07 | 0.82 | 2242 | 860 | 120 | MODEL 23.1.0.g |
| 0.08 | 0.71 | 1867 | 691 | 33 | MODEL 23.1.0.h |

Table 5. Retrospective results (Mohn's Rho) for a ten-year peal on spawning stock biomass and mean absolute scaled error (MASE) analyses from ss3diags library for components of models assessed.

|  |  | MASE |  |  |
| :--- | :---: | :---: | :---: | ---: |
| Model | Mohn's Rho | Index | Fish Length (adj.) | Survey Age |
| M22.2 old | $-0.06(-0.07)$ | 0.69 | $0.93(0.21)$ | 0.35 |
| M22.2 updated | $-0.06(-0.07)$ | 0.69 | $0.93(0.21)$ | 0.35 |
| M23.1.0.a | $0.08(0.10)$ | 0.42 | $1.07(0.24)$ | 0.32 |
| M23.1.0.b | $0.09(0.07)$ | 0.40 | $0.97(0.22)$ | 0.38 |
| M23.1.0.d | $0.09(0.07)$ | 0.40 | $0.96(0.22)$ | 0.38 |
| M23.1.0.g | $0.11(0.08)$ | 0.39 | $0.97(0.22)$ | 0.41 |
| M23.1.0.h | $0.15(0.14)$ | 0.40 | $1.05(0.24)$ | 0.41 |

Table 6. Residual runs test for models evaluated with combined fishery comp data from ss3diags.

| Model | Type | Index | p-value | Test | Sigma3 10 | Sigma3 hi |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| MODEL 22.2 old | cpue | Survey | 0.280 | Passed | -0.376 | 0.376 |
| MODEL 22.2 updated | cpue | Survey | 0.261 | Passed | -0.433 | 0.433 |
| MODEL 23.1.0.a | cpue | Survey | 0.850 | Passed | -0.424 | 0.424 |
| MODEL 23.1.0.b | cpue | Survey | 0.903 | Passed | -0.260 | 0.260 |
| MODEL 23.1.0.d | cpue | Survey | 0.903 | Passed | -0.222 | 0.222 |
| MODEL 23.1.0.g | cpue | Survey | 0.974 | Passed | -0.227 | 0.227 |
| MODEL 23.1.0.h | cpue | Survey | 0.903 | Passed | -0.265 | 0.265 |
| Model22.2 old | len | Fishery | 0.002 | Failed | -0.024 | 0.024 |
| Model22.2 old | len | Survey | 0.000 | Failed | -0.077 | 0.077 |
| MODEL 22.2 updated | len | Fishery | 0.009 | Failed | -0.019 | 0.019 |
| MODEL 22.2 updated | len | Survey | 0.122 | Passed | -0.090 | 0.090 |
| MODEL 23.1.0.a | len | Fishery | 0.003 | Failed | -0.066 | 0.066 |
| MODEL 23.1.0.a | len | Survey | 0.625 | Passed | -0.100 | 0.100 |
| MODEL 23.1.0.b | len | Fishery | 0.155 | Passed | -0.060 | 0.060 |
| MODEL 23.1.0.b | len | Survey | 0.815 | Passed | -0.125 | 0.125 |
| MODEL 23.1.0.d | len | Fishery | 0.155 | Passed | -0.060 | 0.060 |
| MODEL 23.1.0.d | len | Survey | 0.462 | Passed | -0.087 | 0.087 |
| MODEL 23.1.0.g | len | Fishery | 0.155 | Passed | -0.061 | 0.061 |
| MODEL 23.1.0.g | len | Survey | 0.815 | Passed | -0.083 | 0.083 |
| MODEL 23.1.0.h | len | Fishery | 0.015 | Failed | -0.075 | 0.075 |
| MODEL 23.1.0.h | len | Survey | 0.625 | Passed | -0.083 | 0.083 |
| MODEL I22.2_old | age | Survey | 0.039 | Failed | -0.160 | 0.160 |
| MODEL 22.2 updated | age | Survey | 0.401 | Passed | -0.199 | 0.199 |
| MODEL 23.1.0.a | age | Survey | 0.177 | Passed | -0.250 | 0.250 |
| MODEL 23.1.0.b | age | Survey | 0.086 | Passed | -0.161 | 0.161 |
| MODEL 23.1.0.d | age | Survey | 0.298 | Passed | -0.160 | 0.160 |
| MODEL 23.1.0.g | age | Survey | 0.086 | Passed | -0.153 | 0.153 |
| MODEL 23.1.0.h | age | Survey | 0.016 | Failed | -0.152 | 0.152 |

Table 7. Growth parameter values and standard deviations.

| Label | Value | StDev | Model | Label | Value | StDev | Model |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $L_{\text {max }}$ | 112.387 | 3.05 | Model22.2_old | Richards | 1.474 | 0.04 | Model22.2_old |
| Lmax | 116.862 | 1.78 | Model22.2 updated | Richards | 1.541 | 0.02 | Model22.2 updated |
| $\mathrm{L}_{\text {MAX }}$ | 112.958 | 5.92 | MODEL23.1.0.a | Richards | 1.494 | 0.11 | MODEL23.1.0.a |
| $\mathrm{L}_{\text {MAX }}$ | 112.380 | 3.24 | MODEL23.1.0.b | Richards | 1.529 | 0.08 | MODEL23.1.0.b |
| $L_{\text {max }}$ | 112.355 | 3.24 | MODEL23.1.0.d | Richards | 1.528 | 0.08 | MODEL23.1.0.d |
| $L_{\text {max }}$ | 113.217 | 3.23 | MODEL23.1.0.g | Richards | 1.539 | 0.08 | MODEL23.1.0.g |
| $L_{\text {max }}$ | 110.918 | 2.28 | MODEL23.1.0.h | Richards | 1.535 | 0.07 | MODEL23.1.0.h |
| Lmin | 15.134 | 0.45 | Model22.2_old | VonBert K | 0.115 | 0.009 | Model22.2_old |
| $\mathrm{L}_{\text {MIN }}$ | 15.648 | 0.44 | Model22.2_updated | VonBert K | 0.100 | 0.004 | Model22.2_updated |
| L--- | 14.772 | 0.24 | MODEL23.1.0.a | VonBert K | 0.110 | 0.021 | MODEL23.1.0.a |
| $\mathrm{L}_{\text {MIN }}$ | 14.674 | 0.20 | MODEL23.1.0.b | VonBert K | 0.112 | 0.011 | MODEL23.1.0.b |
| Lmin | 14.713 | 0.21 | MODEL23.1.0.d | VonBert K | 0.113 | 0.011 | MODEL23.1.0.d |
| $\mathrm{L}_{\text {MIN }}$ | 14.708 | 0.21 | MODEL23.1.0.g | VonBert K | 0.109 | 0.011 | MODEL23.1.0.g |
| LMIN | 14.681 | 0.20 | MODEL23.1.0.h | VonBert K | 0.131 | 0.009 | MODEL23.1.0.h |

Table 8. Influential parameter values and standard deviations.

| Label | Value | StDev | Model | Label | Value | StDev | Model |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $\mathrm{LN}\left(\mathrm{R}_{0}\right)$ | 13.156 | 0.100 | Model22.2_old | NatM | 0.347 | 0.012 | Model22.2_old |
| LN( $\mathrm{R}_{0}$ ) | 13.016 | 0.075 | Model22.2_updated | NatM | 0.328 | 0.009 | Model22.2_updated |
| LN( $\mathrm{RO}_{0}$ ) | 13.022 | 0.141 | MODEL23.1.0.a | NatM | 0.344 | 0.018 | MODEL23.1.0.a |
| LN(Ro) | 13.601 | 0.241 | MODEL23.1.0.b | NatM | 0.414 | 0.026 | MODEL23.1.0.b |
| $\mathrm{LN}\left(\mathrm{R}_{0}\right)$ | 13.740 | 0.248 | MODEL23.1.0.d | NatM | 0.429 | 0.025 | MODEL23.1.0.d |
| LN(Ro) | 13.688 | 0.240 | MODEL23.1.0.g | NatM | 0.435 | 0.025 | MODEL23.1.0.g |
| LN(Ro) | 13.669 | 0.175 | MODEL23.1.0.h | NatM | 0.424 | 0.021 | MODEL23.1.0.h |
| LnQ BT Shelf Survey | -0.041 | 0.064 | Model22.2_old |  |  |  |  |
| LnQ BT Shelf Survey | -0.026 | 0.049 | Model22.2 updated |  |  |  |  |
| LnQ BT Shelf Survey | 0.092 | 0.086 | MODEL23.1.0.a |  |  |  |  |
| LnQ BT Shelf Survey | -0.196 | 0.163 | MODEL23.1.0.b |  |  |  |  |
| LnQ BT Shelf Survey | -0.268 | 0.172 | MODEL23.1.0.d |  |  |  |  |
| LnQ BT Shelf Survey | -0.233 | 0.162 | MODEL23.1.0.g |  |  |  |  |
| LnQ BT Shelf Survey | -0.213 | 0.104 | MODEL23.1.0.h |  |  |  |  |

Table 9. Derived quantities values, standard deviations, and coefficient of variation.

| Label | Value | StdDev | CV | Model | Label | Value | StdDev | CV | Model |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $\mathrm{F}_{40}$ | 0.33 | 0.02 | 0.05 | Model22.2_old | $\mathrm{B}_{2023}$ | 249,809 | 17,360 | 0.07 | Model22.2_old |
| $\mathrm{F}_{40}$ | 0.29 | 0.01 | 0.04 | Model22.2 updated | B2023 | 263,189 | 14,151 | 0.05 | Model22.2 updated |
| $\mathrm{F}_{40}$ | 0.33 | 0.03 | 0.09 | MODEL23.1.0.a | B2023 | 205,914 | 19,749 | 0.10 | MODEL23.1.0.a |
| $\mathrm{F}_{40}$ | 0.44 | 0.05 | 0.10 | MODEL23.1.0.b | $\mathrm{B}_{2023}$ | 314,146 | 58,787 | 0.19 | MODEL23.1.0.b |
| $\mathrm{F}_{40}$ | 0.47 | 0.05 | 0.11 | MODEL23.1.0.d | $\mathrm{B}_{2023}$ | 343,431 | 66,590 | 0.19 | MODEL23.1.0.d |
| $\mathrm{F}_{40}$ | 0.49 | 0.05 | 0.10 | MODEL23.1.0.g | $\mathrm{B}_{2023}$ | 331,845 | 62,266 | 0.19 | MODEL23.1.0.g |
| $\mathrm{F}_{40}$ | 0.47 | 0.04 | 0.09 | MODEL23.1.0.h | $\mathrm{B}_{2023}$ | 313,052 | 38,688 | 0.12 | MODEL23.1.0.h |
| $\mathrm{ABC}_{2024}$ | 144,694 | 14,664 | 0.10 | Model22.2_old | B0 | 661,455 | 14,493 | 0.02 | Model22.2_old |
| $\mathrm{ABC}_{2024}$ | 141,115 | 11,792 | 0.08 | Model22.2 updated | B0 | 694,750 | 12,587 | 0.02 | Model22.2_updated |
| $\mathrm{ABC}_{2024}$ | 131,883 | 18,010 | 0.14 | MODEL23.1.0.a | $\mathrm{B}_{0}$ | 586,050 | 27,073 | 0.05 | MODEL23.1.0.a |
| $\mathrm{ABC}_{2024}$ | 219,817 | 49,257 | 0.22 | MODEL23.1.0.b | $\mathrm{B}_{0}$ | 605,435 | 50,776 | 0.08 | MODEL23.1.0.b |
| $\mathrm{ABC}_{2024}$ | 243,533 | 56,378 | 0.23 | MODEL23.1.0.d | B0 | 623,435 | 54,253 | 0.09 | MODEL23.1.0.d |
| $\mathrm{ABC}_{2024}$ | 239,088 | 53,953 | 0.23 | MODEL23.1.0.g | $\mathrm{B}_{0}$ | 542,635 | 30,880 | 0.06 | MODEL23.1.0.g |
| $\mathrm{ABC}_{2024}$ |  |  | 0.20 | MODEL23.1.0.h | B0 | 611,365 | 23,726 | 0.04 | MODEL23.1.0.h |

Table 10. Likelihood profiles over survey catchability for the old input sample size and updated input sample size Model 22.2. Light shaded rows are $\pm 10 \mathrm{LL}$ from the MLE, dark shaded row is the closest to MLE.

| M | Q | $\mathrm{B}_{0}$ | $\mathrm{F}_{\text {MSY }}$ | $\mathrm{B}_{2023}$ | $\mathrm{B}_{2023} / \mathrm{B}_{0}$ | $\mathrm{ABC}_{2024}$ | Model | -LL |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0.408 | 0.607 | 702,510 | 0.493 | 372,577 | 0.530 | 243,473 | Model 22.2 old | 10893 |
| 0.402 | 0.638 | 690,370 | 0.489 | 355,882 | 0.515 | 231,967 | Model 22.2 old | 10889 |
| 0.396 | 0.670 | 680,070 | 0.480 | 340,092 | 0.500 | 220,867 | Model 22.2 old | 10886 |
| 0.390 | 0.705 | 671,990 | 0.468 | 325,376 | 0.484 | 210,231 | Model 22.2 old | 10884 |
| 0.384 | 0.741 | 665,730 | 0.453 | 311,391 | 0.468 | 199,983 | Model 22.2 old | 10883 |
| 0.378 | 0.779 | 660,385 | 0.448 | 297,939 | 0.451 | 190,460 | Model 22.2 old | 10880 |
| 0.370 | 0.819 | 659,620 | 0.427 | 287,355 | 0.436 | 181,953 | Model 22.2 old | 10879 |
| 0.363 | 0.861 | 656,310 | 0.424 | 273,501 | 0.417 | 167,867 | Model 22.2 old | 10876 |
| 0.356 | 0.905 | 657,270 | 0.412 | 262,236 | 0.399 | 155,598 | Model 22.2 old | 10876 |
| 0.348 | 0.951 | 660,520 | 0.399 | 251,553 | 0.381 | 146,301 | Model 22.2 old | 10875 |
| 0.340 | 1.000 | 666,530 | 0.385 | 241,723 | 0.363 | 137,214 | Model 22.2 old | 10875 |
| 0.331 | 1.051 | 674,500 | 0.371 | 232,258 | 0.344 | 128,133 | Model 22.2 old | 10876 |
| 0.323 | 1.105 | 685,095 | 0.357 | 223,100 | 0.326 | 119,010 | Model 22.2 old | 10878 |
| 0.314 | 1.162 | 698,290 | 0.343 | 213,998 | 0.306 | 109,750 | Model 22.2 old | 10882 |
| 0.304 | 1.221 | 715,160 | 0.329 | 205,686 | 0.288 | 100,832 | Model 22.2 old | 10885 |
| 0.294 | 1.284 | 735,990 | 0.316 | 198,475 | 0.270 | 92,520 | Model 22.2 old | 10889 |
| 0.284 | 1.350 | 760,510 | 0.302 | 190,942 | 0.251 | 83,996 | Model 22.2 old | 10895 |
| 0.274 | 1.419 | 789,875 | 0.288 | 183,723 | 0.233 | 75,680 | Model 22.2 old | 10902 |
| 0.263 | 1.492 | 824,435 | 0.275 | 176,848 | 0.215 | 67,701 | Model 22.2 old | 10911 |
| 0.251 | 1.568 | 876,375 | 0.258 | 171,031 | 0.195 | 59,398 | Model 22.2 old | 10919 |
| 0.239 | 1.649 | 928,585 | 0.245 | 164,735 | 0.177 | 51,936 | Model 22.2 old | 10929 |
| 0.387 | 0.607 | 721,175 | 0.457 | 384,413 | 0.533 | 233,327 | Model 22.2 update | 18423 |
| 0.382 | 0.638 | 710,870 | 0.447 | 367,428 | 0.517 | 222,109 | Model 22.2 update | 18416 |
| 0.375 | 0.670 | 705,800 | 0.422 | 355,832 | 0.504 | 213,090 | Model 22.2 update | 18437 |
| 0.369 | 0.705 | 695,605 | 0.413 | 339,071 | 0.487 | 202,348 | Model 22.2 update | 18428 |
| 0.363 | 0.741 | 690,235 | 0.402 | 325,001 | 0.471 | 192,820 | Model 22.2 update | 18423 |
| 0.358 | 0.779 | 687,200 | 0.393 | 312,083 | 0.454 | 184,339 | Model 22.2 update | 18421 |
| 0.355 | 0.819 | 686,660 | 0.384 | 306,398 | 0.446 | 179,781 | Model 22.2 update | 18396 |
| 0.343 | 0.861 | 686,650 | 0.379 | 285,626 | 0.416 | 161,963 | Model 22.2 update | 18386 |
| 0.339 | 0.905 | 686,880 | 0.370 | 279,310 | 0.407 | 155,643 | Model 22.2 update | 18363 |
| 0.329 | 0.951 | 689,040 | 0.365 | 259,098 | 0.376 | 139,004 | Model 22.2 update | 18370 |
| 0.323 | 1.000 | 695,900 | 0.354 | 249,146 | 0.358 | 130,620 | Model 22.2 update | 18372 |
| 0.315 | 1.051 | 703,190 | 0.343 | 238,344 | 0.339 | 121,292 | Model 22.2 update | 18369 |
| 0.304 | 1.105 | 716,200 | 0.325 | 228,459 | 0.319 | 111,728 | Model 22.2 update | 18379 |
| 0.293 | 1.162 | 734,860 | 0.310 | 219,578 | 0.299 | 102,230 | Model 22.2 update | 18376 |
| 0.283 | 1.221 | 760,165 | 0.296 | 212,295 | 0.279 | 93,747 | Model 22.2 update | 18378 |
| 0.279 | 1.284 | 781,845 | 0.289 | 204,729 | 0.262 | 86,500 | Model 22.2 update | 18381 |
| 0.264 | 1.350 | 823,400 | 0.270 | 196,966 | 0.239 | 76,732 | Model 22.2 update | 18392 |
| 0.255 | 1.419 | 844,445 | 0.260 | 188,304 | 0.223 | 69,103 | Model 22.2 update | 18397 |
| 0.245 | 1.492 | 888,745 | 0.247 | 181,505 | 0.204 | 61,267 | Model 22.2 update | 18406 |
| 0.233 | 1.568 | 946,240 | 0.233 | 175,511 | 0.185 | 53,735 | Model 22.2 update | 18416 |
| 0.222 | 1.649 | 1,009,605 | 0.220 | 169,199 | 0.168 | 46,607 | Model 22.2 update | 18429 |

Table 11. Model 23.1.0.a and Model 23.1.0.b likelihood profiles over catchability. Light shaded rows are $\pm 2 L L$ from the MLE, dark shaded row is the closest to MLE. *Note hit the lower bound for natural mortality at 0.3 .

| M | Q | $\mathrm{B}_{0}$ | Fmsr | $\mathrm{B}_{2023}$ | $\mathrm{B}_{2023} / \mathrm{B}_{0}$ | $\mathrm{ABC}_{2024}$ | Model | -LL |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0.428 | 0.607 | 628,560 | 0.448 | 358,028 | 0.570 | 266,750 | Model23.1.0.a | 263.27 |
| 0.422 | 0.638 | 615,975 | 0.440 | 340,813 | 0.553 | 253,241 | Model23.1.0.a | 261.84 |
| 0.416 | 0.670 | 604,905 | 0.432 | 324,522 | 0.536 | 240,370 | Model23.1.0.a | 260.43 |
| 0.410 | 0.705 | 595,370 | 0.424 | 309,117 | 0.519 | 228,105 | Model23.1.0.a | 259.04 |
| 0.404 | 0.741 | 587,395 | 0.415 | 294,561 | 0.501 | 216,417 | Model23.1.0.a | 257.69 |
| 0.397 | 0.779 | 581,020 | 0.406 | 280,816 | 0.483 | 205,276 | Model23.1.0.a | 256.39 |
| 0.390 | 0.819 | 576,295 | 0.396 | 267,847 | 0.465 | 194,658 | Model23.1.0.a | 255.16 |
| 0.383 | 0.861 | 573,285 | 0.386 | 255,623 | 0.446 | 184,534 | Model23.1.0.a | 254.04 |
| 0.375 | 0.905 | 572,080 | 0.375 | 244,109 | 0.427 | 174,603 | Model23.1.0.a | 253.04 |
| 0.368 | 0.951 | 572,765 | 0.365 | 233,276 | 0.407 | 160,938 | Model23.1.0.a | 252.2 |
| 0.359 | 1.000 | 575,445 | 0.353 | 223,096 | 0.388 | 149,870 | Model23.1.0.a | 251.55 |
| 0.351 | 1.051 | 580,240 | 0.342 | 213,531 | 0.368 | 140,094 | Model23.1.0.a | 251.15 |
| 0.342 | 1.105 | 587,305 | 0.330 | 204,553 | 0.348 | 130,375 | Model23.1.0.a | 251.04 |
| 0.333 | 1.162 | 596,790 | 0.318 | 196,137 | 0.329 | 120,757 | Model23.1.0.a | 251.28 |
| 0.324 | 1.221 | 608,855 | 0.306 | 188,270 | 0.309 | 111,313 | Model23.1.0.a | 251.93 |
| 0.315 | 1.284 | 623,360 | 0.294 | 182,351 | 0.293 | 103,208 | Model23.1.0.a | 253.05 |
| 0.307 | 1.350 | 639,265 | 0.283 | 178,992 | 0.280 | 97,047 | Model23.1.0.a | 254.52 |
| 0.300* | 1.419 | 652,660 | 0.276 | 176,069 | 0.270 | 92,037 | Model23.1.0.a | 256.36 |
| 0.300* | 1.492 | 654,240 | 0.275 | 173,938 | 0.266 | 90,100 | Model23.1.0.a | 259.23 |
| 0.300* | 1.568 | 655,845 | 0.275 | 172,085 | 0.262 | 88,328 | Model23.1.0.a | 263.39 |
| 0.300* | 1.649 | 657,355 | 0.275 | 170,532 | 0.259 | 86,749 | Model23.1.0.a | 268.84 |
| 0.452 | 0.607 | 673,405 | 0.515 | 416,043 | 0.618 | 298,565 | Model23.1.0b | 144.63 |
| 0.446 | 0.638 | 658,845 | 0.504 | 396,918 | 0.602 | 284,082 | Model23.1.0b | 144.26 |
| 0.440 | 0.670 | 642,510 | 0.483 | 377,148 | 0.587 | 270,404 | Model23.1.0b | 144.25 |
| 0.434 | 0.705 | 634,510 | 0.479 | 361,847 | 0.570 | 256,961 | Model23.1.0b | 143.64 |
| 0.428 | 0.741 | 624,665 | 0.466 | 345,823 | 0.554 | 244,284 | Model23.1.0b | 143.40 |
| 0.421 | 0.779 | 615,195 | 0.454 | 330,336 | 0.537 | 232,235 | Model23.1.0b | 143.26 |
| 0.414 | 0.819 | 606,740 | 0.441 | 315,592 | 0.520 | 220,737 | Model23.1.0b | 143.21 |
| 0.407 | 0.861 | 599,275 | 0.429 | 301,553 | 0.503 | 209,764 | Model23.1.0b | 143.25 |
| 0.400 | 0.905 | 592,795 | 0.417 | 288,186 | 0.486 | 199,290 | Model23.1.0b | 143.40 |
| 0.392 | 0.951 | 587,295 | 0.405 | 275,456 | 0.469 | 189,292 | Model23.1.0b | 143.65 |
| 0.384 | 1.000 | 582,790 | 0.394 | 263,335 | 0.452 | 179,746 | Model23.1.0b | 144.03 |
| 0.376 | 1.051 | 579,350 | 0.382 | 251,806 | 0.435 | 168,994 | Model23.1.0b | 144.55 |
| 0.367 | 1.105 | 578,495 | 0.371 | 243,038 | 0.420 | 158,466 | Model23.1.0b | 145.19 |
| 0.367 | 1.162 | 594,405 | 0.372 | 259,770 | 0.437 | 170,202 | Model23.1.0b | 144.71 |
| 0.360 | 1.221 | 595,900 | 0.364 | 254,814 | 0.428 | 163,101 | Model23.1.0b | 145.08 |
| 0.354 | 1.284 | 597,815 | 0.356 | 250,051 | 0.418 | 156,223 | Model23.1.0b | 145.48 |
| 0.347 | 1.350 | 600,160 | 0.348 | 245,405 | 0.409 | 149,506 | Model23.1.0b | 145.91 |
| 0.340 | 1.419 | 602,040 | 0.341 | 240,473 | 0.399 | 142,939 | Model23.1.0b | 146.41 |
| 0.333 | 1.492 | 606,265 | 0.333 | 236,288 | 0.390 | 137,854 | Model23.1.0b | 146.91 |
| 0.326 | 1.568 | 609,185 | 0.326 | 231,425 | 0.380 | 132,598 | Model23.1.0b | 147.50 |
| 0.319 | 1.649 | 614,605 | 0.317 | 227,219 | 0.370 | 127,351 | Model23.1.0b | 148.09 |

Table 12. Eastern Bering Sea Pacific cod catch for 1964-2022.

| Year | Catch (t) | Year | Catch (t) | Year | Catch (t) |
| :--- | ---: | :--- | ---: | :--- | ---: |
| 1964 | 13,408 | 1984 | 125,103 | 2004 | 183,748 |
| 1965 | 14,719 | 1985 | 143,447 | 2005 | 182,940 |
| 1966 | 18,200 | 1986 | 135,605 | 2006 | 168,818 |
| 1967 | 32,064 | 1987 | 149,903 | 2007 | 140,129 |
| 1968 | 57,902 | 1988 | 203,071 | 2008 | 139,802 |
| 1969 | 50,351 | 1989 | 178,323 | 2009 | 147,174 |
| 1970 | 70,094 | 1990 | 172,067 | 2010 | 142,844 |
| 1971 | 43,054 | 1991 | 210,241 | 2011 | 209,201 |
| 1972 | 42,905 | 1992 | 164,210 | 2012 | 232,623 |
| 1973 | 53,386 | 1993 | 133,186 | 2013 | 236,691 |
| 1974 | 62,462 | 1994 | 172,263 | 2014 | 238,718 |
| 1975 | 51,551 | 1995 | 228,498 | 2015 | 232,829 |
| 1976 | 50,481 | 1996 | 209,067 | 2016 | 247,620 |
| 1977 | 33,335 | 1997 | 232,601 | 2017 | 237,851 |
| 1978 | 42,543 | 1998 | 158,529 | 2018 | 199,867 |
| 1979 | 33,761 | 1999 | 145,867 | 2019 | 178,904 |
| 1980 | 35,058 | 2000 | 151,376 | 2020 | 155,665 |
| 1981 | 56,507 | 2001 | 142,542 | 2021 | 121,749 |
| 1982 | 61,104 | 2002 | 166,555 | 2022 | 152,146 |
| 1983 | 94,801 | 2003 | 175,443 |  |  |

Table 13. Model 23.1.0.d and 23.1.0.g likelihood profiles over catchability. Light shaded rows are $\pm 2 \mathrm{LL}$ from the MLE, dark shaded row is the closest to MLE.

| M | Q | B0 | Fmsy | $\mathrm{B}_{2023}$ | $\mathrm{B}_{2023} / \mathrm{B}_{0}$ | $\mathrm{ABC}_{2024}$ | Model | -LL |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0.457 | 0.607 | 679,640 | 0.522 | 425,216 | 0.626 | 307,231 | Model23.1.0d | 133.82 |
| 0.452 | 0.638 | 665,475 | 0.510 | 405,972 | 0.610 | 292,310 | Model23.1.0d | 133.54 |
| 0.446 | 0.670 | 644,305 | 0.490 | 383,401 | 0.595 | 278,523 | Model23.1.0d | 133.75 |
| 0.440 | 0.705 | 632,835 | 0.478 | 366,254 | 0.579 | 264,915 | Model23.1.0d | 133.56 |
| 0.434 | 0.741 | 622,430 | 0.466 | 349,934 | 0.562 | 251,929 | Model23.1.0d | 133.45 |
| 0.427 | 0.779 | 613,055 | 0.454 | 334,398 | 0.545 | 239,537 | Model23.1.0d | 133.42 |
| 0.420 | 0.819 | 604,685 | 0.442 | 319,606 | 0.529 | 227,711 | Model23.1.0d | 133.47 |
| 0.413 | 0.861 | 597,300 | 0.430 | 305,521 | 0.512 | 216,424 | Model23.1.0d | 133.63 |
| 0.406 | 0.905 | 590,875 | 0.418 | 292,103 | 0.494 | 205,655 | Model23.1.0d | 133.89 |
| 0.398 | 0.951 | 585,460 | 0.406 | 279,330 | 0.477 | 195,369 | Model23.1.0d | 134.28 |
| 0.391 | 1.000 | 581,080 | 0.395 | 267,246 | 0.460 | 185,600 | Model23.1.0d | 134.79 |
| 0.382 | 1.051 | 580,115 | 0.384 | 259,479 | 0.447 | 178,331 | Model23.1.0d | 135.41 |
| 0.383 | 1.105 | 603,225 | 0.385 | 286,700 | 0.475 | 192,596 | Model23.1.0d | 134.26 |
| 0.377 | 1.162 | 603,805 | 0.377 | 281,452 | 0.466 | 187,133 | Model23.1.0d | 134.49 |
| 0.370 | 1.221 | 604,715 | 0.368 | 276,276 | 0.457 | 181,787 | Model23.1.0d | 134.75 |
| 0.364 | 1.284 | 605,975 | 0.360 | 271,122 | 0.447 | 176,529 | Model23.1.0d | 135.05 |
| 0.357 | 1.350 | 607,595 | 0.352 | 265,972 | 0.438 | 169,713 | Model23.1.0d | 135.38 |
| 0.350 | 1.419 | 609,605 | 0.344 | 260,820 | 0.428 | 162,337 | Model23.1.0d | 135.75 |
| 0.344 | 1.492 | 612,035 | 0.336 | 255,666 | 0.418 | 155,045 | Model23.1.0d | 136.17 |
| 0.337 | 1.568 | 614,925 | 0.329 | 250,521 | 0.407 | 147,842 | Model23.1.0d | 136.64 |
| 0.330 | 1.649 | 618,320 | 0.321 | 245,392 | 0.397 | 141,201 | Model23.1.0d | 137.17 |
| 0.469 | 0.607 | 593,295 | 0.699 | 425,484 | 0.717 | 313,571 | Model23.1.0g | 142.10 |
| 0.463 | 0.638 | 578,390 | 0.680 | 404,851 | 0.700 | 298,556 | Model23.1.0g | 141.95 |
| 0.457 | 0.670 | 570,505 | 0.664 | 387,512 | 0.679 | 283,536 | Model23.1.0g | 141.46 |
| 0.450 | 0.705 | 561,410 | 0.645 | 370,082 | 0.659 | 269,474 | Model23.1.0g | 141.22 |
| 0.444 | 0.741 | 552,870 | 0.627 | 353,318 | 0.639 | 256,106 | Model23.1.0g | 141.06 |
| 0.437 | 0.779 | 545,455 | 0.610 | 337,364 | 0.619 | 243,336 | Model23.1.0g | 140.99 |
| 0.430 | 0.819 | 539,170 | 0.592 | 322,179 | 0.598 | 231,135 | Model23.1.0g | 141.00 |
| 0.423 | 0.861 | 529,465 | 0.578 | 306,090 | 0.578 | 219,879 | Model23.1.0g | 141.27 |
| 0.416 | 0.905 | 523,905 | 0.558 | 291,697 | 0.557 | 208,847 | Model23.1.0g | 141.56 |
| 0.408 | 0.951 | 523,590 | 0.548 | 279,700 | 0.534 | 197,917 | Model23.1.0g | 141.73 |
| 0.400 | 1.000 | 525,145 | 0.521 | 268,281 | 0.511 | 187,520 | Model23.1.0g | 142.25 |
| 0.392 | 1.051 | 523,205 | 0.513 | 258,255 | 0.494 | 179,501 | Model23.1.0g | 142.89 |
| 0.392 | 1.105 | 543,845 | 0.510 | 284,780 | 0.524 | 193,289 | Model23.1.0g | 142.05 |
| 0.386 | 1.162 | 545,660 | 0.499 | 279,366 | 0.512 | 187,809 | Model23.1.0g | 142.38 |
| 0.379 | 1.221 | 548,460 | 0.487 | 274,431 | 0.500 | 182,550 | Model23.1.0g | 142.74 |
| 0.373 | 1.284 | 548,815 | 0.477 | 268,000 | 0.488 | 177,088 | Model23.1.0g | 143.24 |
| 0.367 | 1.350 | 555,755 | 0.463 | 265,142 | 0.477 | 172,450 | Model23.1.0g | 143.61 |
| 0.360 | 1.419 | 559,670 | 0.452 | 260,441 | 0.465 | 167,525 | Model23.1.0g | 144.14 |
| 0.354 | 1.492 | 563,400 | 0.441 | 255,520 | 0.454 | 162,646 | Model23.1.0g | 144.74 |
| 0.347 | 1.568 | 568,140 | 0.430 | 250,993 | 0.442 | 157,951 | Model23.1.0g | 145.40 |
| 0.341 | 1.649 | 573,885 | 0.419 | 246,846 | 0.430 | 151,632 | Model23.1.0g | 146.12 |

Table 14. Model 23.1.0.h likelihood profile over catchability. Light shaded rows are $\pm 2$ LL from the MLE, dark shaded row is the closest to MLE. *Note the model hit the lower bound for natural mortality at 0.3.

| $\mathbf{M}$ | $\mathbf{Q}$ | $\mathbf{B}_{\mathbf{0}}$ | $\mathbf{F}_{\text {MSY }}$ | $\mathbf{B}_{2023}$ | $\mathbf{B}_{2023} / \mathbf{B}_{\mathbf{0}}$ | $\mathbf{A B C}_{2024}$ | Model | -LL |
| ---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0.470 | 0.607 | 666,785 | 0.542 | 403,687 | 0.605 | 337,328 | Model23.1.0.h | 633.52 |
| 0.462 | 0.638 | 654,040 | 0.527 | 386,153 | 0.590 | 320,677 | Model23.1.0.h | 632.70 |
| 0.455 | 0.670 | 642,440 | 0.514 | 369,347 | 0.575 | 304,868 | Model23.1.0.h | 632.01 |
| 0.447 | 0.705 | 630,285 | 0.505 | 352,520 | 0.559 | 290,158 | Model23.1.0.h | 631.44 |
| 0.439 | 0.741 | 621,895 | 0.491 | 337,432 | 0.543 | 275,664 | Model23.1.0.h | 631.00 |
| 0.430 | 0.779 | 615,225 | 0.477 | 323,154 | 0.525 | 261,761 | Model23.1.0.h | 630.74 |
| 0.421 | 0.819 | 610,240 | 0.462 | 309,616 | 0.507 | 248,426 | Model23.1.0.h | 630.69 |
| 0.412 | 0.861 | 606,980 | 0.446 | 296,768 | 0.489 | 235,623 | Model23.1.0.h | 630.87 |
| 0.402 | 0.905 | 605,490 | 0.430 | 284,563 | 0.470 | 223,327 | Model23.1.0.h | 631.32 |
| 0.392 | 0.951 | 605,800 | 0.414 | 272,945 | 0.451 | 211,513 | Model23.1.0.h | 632.08 |
| 0.381 | 1.000 | 607,970 | 0.397 | 261,850 | 0.431 | 200,149 | Model23.1.0.h | 633.18 |
| 0.371 | 1.051 | 610,170 | 0.381 | 250,663 | 0.411 | 189,431 | Model23.1.0.h | 634.66 |
| 0.360 | 1.105 | 616,560 | 0.365 | 241,103 | 0.391 | 178,228 | Model23.1.0.h | 636.54 |
| 0.356 | 1.162 | 624,225 | 0.360 | 245,238 | 0.393 | 179,138 | Model23.1.0.h | 638.05 |
| 0.347 | 1.221 | 632,205 | 0.349 | 241,256 | 0.382 | 172,501 | Model23.1.0.h | 640.02 |
| 0.339 | 1.284 | 641,035 | 0.338 | 237,399 | 0.370 | 165,824 | Model23.1.0.h | 642.18 |
| 0.331 | 1.350 | 650,730 | 0.328 | 233,657 | 0.359 | 159,122 | Model23.1.0.h | 644.55 |
| 0.322 | 1.419 | 661,315 | 0.318 | 230,023 | 0.348 | 152,413 | Model23.1.0.h | 647.12 |
| 0.314 | 1.492 | 672,810 | 0.308 | 226,494 | 0.337 | 145,719 | Model23.1.0.h | 649.92 |
| 0.306 | 1.568 | 685,235 | 0.298 | 223,063 | 0.326 | 139,061 | Model23.1.0.h | 652.95 |
| $0.300 *$ | 1.649 | 694,810 | 0.292 | 220,578 | 0.317 | 134,275 | Model23.1.0.h | 656.26 |

Table 15. Negative log likelihood and derived quantities for assessed models for models with (Free M) M fit with a uninformative prior, (Fixed M) fixed at 0.386623 , and the AIC weighted (Burnham and Anderson 2002) values from the likelihood profiles on survey index catchability between -0.5 and 0.5 for the free $M$ models. Negative log likelihood (-LL), unfished female spawning biomass ( $\mathrm{B}_{0}$ ), female spawning biomass in 2023 ( $\mathrm{B}_{2023}$ ), and projected 2024 allowable biological catch ( $\mathrm{ABC}_{2024}$ ).

| Model | Free M |  |  |  |  | Fixed M |  |  |  |  | Change in -LL | AIC Weighted |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | -LL | $\mathrm{B}_{0}$ | $\mathrm{B}_{2023}$ | $\mathrm{B}_{2023} / \mathrm{B}_{0}$ | $\mathrm{ABC}_{2024}$ | -LL | $\mathrm{B}_{0}$ | $\mathrm{B}_{2023}$ | $\mathrm{B}_{2023} / \mathrm{B}_{0}$ | $\mathrm{ABC}_{2024}$ |  | $\mathrm{B}_{0}$ | B2023 | $\mathrm{B}_{2023} / \mathrm{B}_{0}$ | $\mathrm{ABC}_{2024}$ |
| M22.2 old | 10875 | 661,455 | 249,809 | 0.378 | 144,694 | 10,881 | 653,795 | 295,111 | 0.451 | 192,152 | 6 | 663,556 | 249,862 | 0.377 | 144,781 |
| M22.2 up | 18362 | 694,750 | 263,189 | 0.379 | 141,115 | 18,405 | 683,985 | 332,473 | 0.486 | 204,657 | 43 | 686,923 | 279,187 | 0.406 | 155,540 |
| M23.1.0.a | 251 | 586,050 | 205,914 | 0.351 | 131,883 | 253 | 568,340 | 246,505 | 0.434 | 178,060 | 2 | 587,786 | 209,209 | 0.356 | 134,930 |
| M23.1.0.b | 143 | 605,435 | 314,146 | 0.519 | 219,817 | 144 | 590,270 | 274,837 | 0.466 | 187,374 | 1 | 610,263 | 314,592 | 0.516 | 218,929 |
| M23.1.0.d | 133 | 623,435 | 343,431 | 0.551 | 243,533 | 134 | 594,955 | 276,042 | 0.464 | 188,263 | 1 | 620,013 | 333,797 | 0.538 | 236,289 |
| M23.1.0.g | 141 | 542,635 | 331,845 | 0.612 | 239,088 | 143 | 531,915 | 264,534 | 0.497 | 181,473 | 2 | 547,403 | 330,233 | 0.603 | 236,739 |
| M23.1.0.h | 631 | 611,365 | 313,052 | 0.512 |  | 632 | 613,550 | 276,694 | 0.451 |  | 1 | 616,616 | 318,377 | 0.516 |  |

Table 16. Survey catchability (Q) estimates for models fit with (Free M) a non-informative prior on natural mortality and (Fixed M) models fit with a fixed natural mortality of 0.386623 .

|  | Survey catchability |  |
| :--- | ---: | ---: |
| with |  |  |
| Model | Free M | Fixed $\mathbf{M}$ |
| M22.2 old | 0.960 | 0.772 |
| M22.2 up | 0.974 | 0.683 |
| M23.1.0.a | 1.097 | 0.902 |
| M23.1.0.b | 0.822 | 0.953 |
| M23.1.0.d | 0.765 | 0.972 |
| M23.1.0.g | 0.792 | 1.017 |
| M23.1.0.h | 0.808 | 0.949 |

9 Figures


Figure 1. End year (top) length (cm) at age (bottom left) difference in length by age from Model 22.2 old, and (bottom right) difference in weight by age from Model 22.2 old for all models. Please note that the weight difference for Model 23.1.0.h (pink) is in error and should not be considered.






Figure 2. Length at age over time in each of the models examined.


Figure 3. Fishery (left) and survey (right) selectivity for 2022.



Figure 4. Annually varying selectivity for (top) fishery and (bottom) survey.


Figure 5. Model fits to the log of the bottom trawl survey index for all models examined.


Figure 6. The distribution of the log of virgin recruitment $\left(R_{0}\right)$ for all models.


Figure 7. Recruitment in numbers of Age-0 Pacific cod with $95 \%$ confidence bounds for all models.


Figure 8. Spawning stock biomass/unfished spawning biomass with $\mathrm{B}_{35 \%}$ management target and (shaded) $95 \%$ confidence bounds for all models.


Figure 9. Total spawning biomass (males and females) with (shaded) $95 \%$ confidence bounds for all models.


Figure 10. Likelihood profiles scaling the log survey catchability index from -0.5 to 0.5 for the main model components and in total.


Figure 11. Length comps, aggregated across time by fleet for (left) Model 22.2 and (right) Model 23.1.0a


Figure 12. Pearson residuals for length composition, comparing across fleets for (left) Model 22.2 and (right) Model 23.1.0.a. Closed bubbles are positive residuals (observed > expected) and open bubbles are negative residuals (observed < expected).


Figure 13. Bottom trawl survey age composition distributions and model fits (green line). Note that models 23.1.0.b, .d, and .g are nearly indistinguishable visually.


Figure 14. Pearson residuals for survey age composition. Closed bubbles are positive residuals (observed > expected) and open bubbles are negative residuals (observed < expected).


Figure 15. Model 23.1.0.b (Left) length comps, aggregated across time by fleet and (right) Pearson residuals, comparing across fleets for (left) Model 22.2 and (right) Model 23.1.0.a. Closed bubbles are positive residuals (observed > expected) and open bubbles are negative residuals (observed < expected).


Model 23.1.0.b


Figure 16. Key parameters fit for the likelihood profile scaling the log survey index catchability from -0.5 to 0.5 for Model 22.2 and Model 23.1.0.b.


Figure 17. Bottom trawl survey selectivity for Model 23.1.0.b with log catchability fixed at between (far left) -0.25 and (far right) 0.5 showing change to dome-shaped selectivity appearing in the likelihood profile over catchability.


Figure 18. Model 23.1.0.d (Left) length comps, aggregated across time by fleet and (right) Pearson residuals, comparing across fleets for (left) Model 22.2 and (right) Model 23.1.0.a. Closed bubbles are positive residuals (observed > expected) and open bubbles are negative residuals (observed < expected).


Figure 19. Model 23.1.0.g (Left) length comps, aggregated across time by fleet and (right) Pearson residuals, comparing across fleets for (left) Model 22.2 and (right) Model 23.1.0.a. Closed bubbles are positive residuals (observed > expected) and open bubbles are negative residuals (observed < expected).


Figure 20. Conditional age-at-length data used in Model 23.1.0.h.


Figure 21. Model 23.1.0.h (Left) length comps, aggregated across time by fleet and (right) Pearson residuals, comparing across fleets for (left) Model 22.2 and (right) Model 23.1.0.a. Closed bubbles are positive residuals (observed > expected) and open bubbles are negative residuals (observed < expected).


Figure 22. Conditional age-at-length (CAAL) data distributions and Model 23.1.0.h fits to the data including (bottom right) mean age from the CAAL data.



Figure 23. Mean length at age and 95\% confidence intervals for (left) Model 23.1.0.g and right (Model 23.1.0.h) showing reduction in uncertainty in growth estimates.


Figure 24. Annual devs on (left) Richards parameter and (right) $L_{\text {min }}$ for Model23.1.0.g and Model 23.1.0.h.


Figure 25 Total spawning biomass from 10-year retrospective peels of Model 23.1.0d with (left) natural mortality fit with an uninformative prior and (right) with natural mortality fixed at 0.387.

