# Implementing the 2014 three-stock model for Southern Hemisphere Breeding Stocks E1, Oceania and G 

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#### Abstract

The three stock model approach with mixing on feeding grounds, which was implemented in 2014 to assess the BSD, BSE1 and BSO breeding stocks, is applied in a similar manner to the BSE1, BSO and BSG breeding stocks. This is intended as a first step in applying this approach consecutively around the globe to check for consistency of results in circumstances of uncertainty in the allocation of feeding ground catches. Compared to earlier assessments of the BSE1, BSO and BSG breeding stocks in partly separate analyses, there is an appreciable change in that the pre-exploitation level is estimated higher for BSE1and lower for BSO; correspondingly, BSE1 is estimated as less and BSO as more recovered towards those pre-exploitation levels. Comparable likelihoods do not differ greatly for data for these stocks used in both the 2014 BSD+BSE1+BSO and the 2015 BSE1+BSO+BSG assessments, suggesting that the data used here do not contain sufficient information to distinguish these rather different results.


## INTRODUCTION

In 2014, a three-stock approach was implemented in order to jointly assess the West Australian Breeding Stock D (BSD), East Australian Breeding Stock E1 (BSE1) and the Oceania breeding stock (BSO). The rationale behind a three-stock approach was primarily to address the uncertainty about how to allocate the historical feeding ground catches between the three stocks (IWC 2014). The three-stock model defined a western and an eastern feeding ground with a boundary at $130^{\circ} \mathrm{E}$. BSD and an estimable proportion of the BSE1 stock feed (and were caught) in the western feeding ground west of $130^{\circ} \mathrm{E}$, while BSO and the remaining proportion of BSE1 feed in the eastern feeding ground. These assumptions were used to allocate the historical catches to breeding stocks in proportion to the abundances of the populations present in each feeding ground.

Allocation of feeding ground catches remains a challenge for the assessment of the Southern Hemisphere humpback whale populations, since assumptions have to be made about where to place the boundaries for each stock. An attempt was made to develop an all-stock assessment which combined seven breeding populations into a single assessment (Müller et al. 2010), where all feeding ground catches could be allocated in proportion to population abundances present on the feeding ground, rather than fixed on input. This work, however, was not completed owing to convergence issues which arose as a result of too many estimable parameters. Another possible approach would be to utilise the less ambitious three-stock model to successively assess all the stocks, i.e. to follow the $\mathrm{D}+\mathrm{E} 1+\mathrm{O}$ assessment with a $\mathrm{E} 1+\mathrm{O}+\mathrm{G}$ assessment and so on, in order to allow flexibility in the allocation of historical feeding ground catches for all the stocks and so to assess how key parameter estimates change for each iteration. The aim would be to assess the potential of using successive three-stock models to develop a combined assessment of all the stocks.

This paper presents the results of a three-stock E1+O+G model. Similar to the $\mathrm{D}+\mathrm{E} 1+\mathrm{O}$ model, a western and eastern feeding area are defined (Figure 1). The western region runs from $120^{\circ} \mathrm{E}$ to $100^{\circ} \mathrm{W}$ and BSE 1 and a proportion $\left(\gamma^{\circ}\right)$ of the BSO population are assumed to feed there. The eastern region runs between $100^{\circ} \mathrm{W}$ and $50^{\circ} \mathrm{W}$ and BSG and the remaining proportion (1- $\gamma^{O}$ ) of the BSO population are assumed to feed there. Figure 2 provides a juxtaposition of the $\mathrm{D}+\mathrm{E} 1+\mathrm{O}$ and the $\mathrm{E} 1+\mathrm{O}+\mathrm{G}$ models in order to provide comparison of the boundaries and total catches taken in the respective western and eastern feeding areas.

[^0]

Figure 1: Diagrammatic representation of the E1+O+G three-stock model. BSE1 and a proportion ( $\gamma^{O}$ ) of the BSO population feed in the western feeding area, while BSG and the remaining proportion $\left(1-\gamma^{O}\right)$ of the BSO population feed in the eastern feeding ground. Historical feeding ground catches from each feeding ground are allocated to the different stocks in proportion to the model-predicted abundances in each area. Catches taken between $70^{\circ} \mathrm{W}$ and $50^{\circ} \mathrm{W}$ are further split between the eastern feeding ground and the Brazilian Breeding Stock A, as described later in the text.


## 2015 E1+O+G three-stock model (cumulative total of 59321 catches)

Figure 2: Juxtaposition of the 2014 D+E1+O three-stock model (dashed lines) with the 2015 E1+O+G three-stock model (solid lines), illustrating the boundaries of the respective eastern and western feeding areas, as well as the total historical catches taken in each area. In IWC (2014) $\gamma^{E l}$ was estimated to be 0.068 , i.e. $6.8 \%$ of BSE1 feed in western feeding area

## DATA

## Historical catch data

There are two sets of historical catch data, both of which are available from Allison's database (C. Allison, pers. commn):
i) Catches north of $40^{\circ} \mathrm{S}$

These catches are given by location. Additionally there are some Russian catch data available by 10 degree longitude and latitude bands. The allocations of these catches to the breeding stocks considered in this assessment are described below.

## Breeding Stock E1and Oceania

The catches for E1 and Oceania are given by landing station. Catches landed at LochTay, Tangalooma and Byron Bay have been allocated to BSE1. Catches landed at Norfolk Island, Tonga and Polynesia have been allocated to the Oceania breeding stock. Catches landed at Rakiura and Prince George, New Zealand, Kaikoura, Cook Strait and Tory Channel, Great Barrier Island, Whangamumu and Bay of Island are allocated to BSE1 and Oceania in proportion to the model-predicted population abundances.

## Breeding Stock $G$

Catches labelled "Chile", "Ch/Per/E" and "Peru" in the database have been allocated to BSG.
The resulting catch series are given in Table A. 1 of the Appendix.
ii) Catches south of $40^{\circ} \mathrm{S}$

These catches are given for 10 degree longitude bands, as shown in Table A. 2. Catches taken between $120^{\circ} \mathrm{E}$ and $100^{\circ} \mathrm{W}$ are allocated to the western feeding ground and catches taken between $100^{\circ} \mathrm{W}$ and $70^{\circ} \mathrm{W}$ are allocated to the eastern feeding ground. IWC (2010) splits the area between $70^{\circ} \mathrm{W}$ and $50^{\circ} \mathrm{W}$ into three blocks: one from $40^{\circ} \mathrm{S}$ $50^{\circ} \mathrm{S}$ (allocated to the Brazilian breeding stock BSA), one from $50^{\circ} \mathrm{S}-60^{\circ} \mathrm{S}$ (shared between BSA and BSG) and lastly one south of $60^{\circ} \mathrm{S}$ (see Figure 6). In light of this, all catches between $70^{\circ} \mathrm{W}$ and $50^{\circ} \mathrm{W}$ and south of $60^{\circ} \mathrm{S}$ were added to the eastern feeding ground catches for the three-stock model. Further, half the catches between $70-50^{\circ} \mathrm{W}$ and $50-60^{\circ} \mathrm{S}$ were added to the eastern feeding ground catches.

## Abundance and trend data

Breeding Stock E1 was fit to the Noad et al. (2011) absolute abundance estimate, as well as the Noad et al. (2011) relative abundance series. Breeding Stock Oceania was fit to the Constantine et al. (2012) mark-recapture data. Breeding Stock G was fit to the Felix et al. (2011) absolute abundance estimate. The data are listed in the Appendix along with further details.

In their assessment of Breeding Stock G, Johnston et al. (2011) fit to the Felix et al. (2011) absolute abundance estimate, but also to the Branch (2011) IDCR/SOWER relative abundance series. This series was derived for the feeding ground area between $110^{\circ} \mathrm{W}$ and $70^{\circ} \mathrm{W}$, and since these boundaries correspond closely to the eastern feeding ground $\left(100^{\circ} \mathrm{W}-70^{\circ} \mathrm{W}\right.$, with only 114 catches taken between $\left.110^{\circ} \mathrm{W}-100^{\circ} \mathrm{W}\right)$, the population feeding in the eastern area of E1+O+G three-stock model $\left(\left(1-\gamma^{O}\right) N^{B S O}+N^{B S G}\right)$ was fit to the IDCR/SOWER relative abundance series as a sensitivity.

## $N_{\text {min }}$ constraint

The minimum population constraint was taken to be three times the minimum number of haplotypes (IWC 2012). The minimum number of haplotypes utilised were: 5 for BSE1 (IWC 2014), 33 for BSO (IWC 2014) and 27 for BSG (Rosenbaum et al. 2006).

## METHODS

## Population dynamics

The population dynamics are given by the following equation:

$$
\begin{equation*}
N_{y+1}^{i}=N_{y}^{i}+r^{i} N_{y}^{i}\left(1-\left(\frac{N_{y}^{i}}{K^{i}}\right)^{\mu}\right)-C_{y}^{i} \quad i \in\{\mathrm{E} 1, \text { Oceania, G }\} \tag{1}
\end{equation*}
$$

where
$N_{y}^{i} \quad$ is the number of whales in the breeding population $i$ at the start of year $y$,
$r^{i} \quad$ is the intrinsic growth rate (the maximum per capita the population can achieve when its size is very low) of breeding population $i$,
$K^{i} \quad$ is the carrying capacity or pristine population level of breeding population $i$,
$\mu \quad$ is the "degree of compensation" parameter; this is set at 2.39 , which fixes the level at which MSY is achieved at MSYL $=0.6 K$, as conventionally assumed by the IWC SC, and
$C_{y}^{i} \quad$ is the total catch (in terms of breeding population $i$ animals) in year $y$.

## Bayesian estimation framework

## Priors

Prior distributions are defined for the following parameters:
i) $\quad r^{i} \sim \mathrm{U}[0,0.106]^{2}$
ii) $\quad \ln \tilde{N}_{t \text { target }}^{i, o s} \sim U\left[\ln N_{\text {target }}^{i, o b s}-4 C V, \ln N_{t \text { arget }}^{i, \text { obs }}+4 C V\right]$

The target abundance estimate is fitted to the model-predicted number of whales for breeding population $i$.
The uninformative $r$ prior is bounded by zero (negative rates of growth are biologically implausible - at least over lengthy periods) and 0.106 (this corresponds to the maximum growth rate for the species agreed by the IWC Scientific Committee (IWC 2007)). The prior distribution from which the target abundance estimate $\tilde{N}_{t \text { target }}^{i, o b s}$ is drawn at random is uniform on a natural logarithmic scale. The upper and lower bounds, whose only purpose is to render the computations more efficient, are set by the CV of the abundance estimate multiplied by four.
Using the randomly drawn vector of values of $\tilde{N}_{t \text { arg et }}^{i, o b s}$ and $r^{i}$, a downhill simplex method of minimization is used to calculate $K^{i}$ such that the model estimate of $N_{t \text { arg } e t}^{i}$ is identical to the randomly drawn value $\tilde{N}_{t \text { target }}^{i, o b s}$.

For each simulation, using the $r^{i}$ and calculated $K^{i}$ values, the available data are used to assign a likelihood to that particular combination. Details for calculating the components of the negative log likelihood are given below.

## Likelihood function

## Absolute abundance data

Given an absolute abundance estimate, $N_{t a r g e t}^{o b s}$, this is assumed to be log-normally distributed with the $\log$ of the estimate as the mean and the CV as the standard deviation ${ }^{3}$. Thus the negative $\log$ likelihood contribution is:

$$
\begin{equation*}
\frac{1}{2 \sigma^{2}}\left(\ln N_{t \arg e t}^{o b s}-\ln N_{t \arg e t}\right)^{2} \tag{2}
\end{equation*}
$$

where

$$
\begin{array}{ll}
N_{t a r g e t}^{o b s} & \text { is the absolute abundance estimate obtained from observations, } \\
N_{\text {target }} & \text { is the model-estimated population size for the year of the abundance estimate, and } \\
\sigma^{2} & \text { is the variance of } \ln N_{t a r g e t}^{o b s}
\end{array}
$$

## Relative abundance data

These estimates are given in a series spanning several years. Each year has a relative abundance index $I_{y}$, obtained from observations. It is assumed that this index is log-normally distributed about its expected value:

$$
\begin{equation*}
I_{y}=q N_{y} e^{\varepsilon_{y}} \tag{3}
\end{equation*}
$$

where

[^1]$I_{y} \quad$ is the relative abundance estimate for year $y$,
$q$ is a constant of proportionality,
$N_{y} \quad$ is the model estimate of observed population size at the start of year $y$, and
$\varepsilon_{y}$ is from $N\left(0, \sigma^{2}\right)$ (see Equation (4) below).
The $\sigma$ parameter is the residual standard deviation, which is estimated in the fitting procedure by its maximum likelihood value:
\[

$$
\begin{equation*}
\hat{\sigma}=\sqrt{1 / \bar{n} \sum_{y}\left(\ln I_{y}-\ln q-\ln N_{y}\right)^{2}} \tag{4}
\end{equation*}
$$

\]

where
$\bar{n} \quad$ is the number of data points in the series, and
$q$ is a constant of proportionality, estimated by its maximum likelihood value:

$$
\begin{equation*}
\ln \hat{q}=1 / \bar{n} \sum_{y}\left(\ln I_{y}-\ln N_{y}\right) \tag{5}
\end{equation*}
$$

The negative log-likelihood component for the relative abundance data is given by:

$$
\begin{equation*}
\bar{n} \ln \sigma+\frac{1}{2 \sigma^{2}} \sum_{y}\left(\ln I_{y}-\ln q-\ln N_{y}\right)^{2} \tag{6}
\end{equation*}
$$

In the Bayesian context, $q$ and $\sigma$ are "nuisance parameters, i.e. parameters that need to be estimated but are not of great interest themselves (McAllister et al., 1994). Walters and Ludwig (1994) show that the above approach is essentially a shortcut to avoid integrating over the prior distributions parameters and corresponds to the assumption that the $q$ prior is uniformly distributed in log-space, and that the $\sigma$ prior is proportional to $\sigma^{-3}$.

## Mark recapture data

These data are given in the form a matrix showing counts of animals that were seen in a specific year and re-seen in a subsequent year. The method for incorporating this information into the likelihood is given below.

The capture-recapture data give:
$n_{y}$, the number of animals captured in year $y$, and
$m_{y, y^{\prime}}$, the number of animals captured in year $y$ that were recaptured in year $y^{\prime}$.
If $p_{y}$ is the probability that an animal is seen in a region in year $y$, then the number of animals captured in year $y$ is given by:

$$
\begin{equation*}
n_{y}=p_{y} N_{y} \tag{7}
\end{equation*}
$$

where $N_{y}$ is the total (1+) population. The model predicted number of animals captured in year $y$ that were recaptured in year $y^{\prime}$ is given by:

$$
\begin{equation*}
\hat{m}_{y, y^{\prime}}=p_{y} p_{y^{\prime}} N_{y} e^{-M\left(y^{\prime}-y\right)} \tag{8}
\end{equation*}
$$

where $M$ is the natural mortality rate (set here to equal $0.03 \mathrm{yr}^{-1}$ as recommended by the IWC SC).
The probability of a model-predicted $\hat{m}_{y, y^{\prime}}$, given the observed $m_{y, y^{\prime}}$, is determined assuming a Poisson distribution ${ }^{4}$, with the associated likelihood contribution given by:

$$
\begin{equation*}
\frac{\left(\hat{m}_{y, y^{\prime}}\right)^{m_{y, y^{\prime}}}}{m_{y, y^{\prime}!}!} e^{-\hat{m}_{y, y^{\prime}}} \tag{9}
\end{equation*}
$$

[^2]Finally the component for the negative of the log-likelihood for capture-recapture data is then given by:

$$
\begin{equation*}
\sum_{y=y_{o}}^{y_{f}-1} \sum_{y^{\prime}=y+1}^{y_{f}}\left[-m_{y, y^{\prime}} \ln \hat{m}_{y, y^{\prime}}+\hat{m}_{y, y^{\prime}}\right] \tag{10}
\end{equation*}
$$

where $y_{0}$ is the first year of captures and $y_{f}$ is the last year of recaptures.
Note that when compiling the capture-recapture matrices, if an animal is re-seen a second time, the first resighting is treated as a new sighting that is first re-seen at the second resighting.

## SIR

The negative log likelihood is then converted into a likelihood value $(L)$. The integration of the prior distributions of the parameters and the likelihood function then essentially follows the Sampling-Importance-Resampling (SIR) algorithm presented by Rubin (1988). For a vector of parameter values $\theta_{i}$, the likelihood of the data associated with this vector of parameters $(L)$ as described above is calculated and stored as $\tilde{L}$. This process is repeated until an initial sample of $n_{1} \theta_{i} \mathrm{~s}$ is generated.

This sample is then resampled with replacement $n_{2}$ times with probability equal to weight $w_{\mathrm{j}}$, where:

$$
\begin{equation*}
w_{j}=\frac{\tilde{L}\left(\theta_{j} / \text { data }\right)}{\sum_{j=1}^{m_{1}} \tilde{L}\left(\theta_{j} / \text { data }\right)} \tag{11}
\end{equation*}
$$

The resample is thus a random sample of size $n_{2}$ from the joint posterior distribution of the parameters (Rubin, 1988).

## Importance functions for BSE1

The trend data for BSE1 (Noad et al., 2011) are highly informative, and as such high $r^{E l}$ values have a much higher likelihood associated with them and have a much better chance of being resampled. Since $r^{E l}$ is sampled from a uniform prior on the interval [0,0.106], small values of $r^{E l}$ will form a substantial proportion of the initial sample of $n_{l}$, even though they are not likely to be chosen in the resampling process. This leads to sampling inefficiency and a high number of duplicates (where the same high $r^{E l}$ values are sampled repeatedly). A very large initial sample has to then be drawn in order to generate enough samples with high $r^{E I}$ values to be able to resample without a high number of duplicates. In order to increase the sampling efficiency, an importance function was used. This function increases the likelihood of sampling high $r^{E l}$ values and reduces the number of essentially wasted low $r^{E l}$ values in the sample. To counter the fact that the resulting distribution of the $n_{l}$ values of $r^{E I}$ sampled is no longer uniform as required by the uniform prior distribution, the final likelihood values are weighted upwards in the same proportion as the probability of picking a particular $r^{E l}$ in the initial sample was weighted down. The importance function is shown in Figure 3 below.


Figure 3: Importance functions used when sampling for $r^{E l}$. The horizontal axis shows the step values of $r$ at which the importance function increases, and the vertical axis shows the probability of accepting an $r$ sample from a particular range. In other words, if a value between 0 and 0.090 is drawn for $r^{E l}$ from the uniform prior, it has a $95 \%$ chance of being discarded.

## $\boldsymbol{N}_{\text {min }}$ constraints

The assumption for these assessments is that given a minimum number of haplotypes, $h$, for a specific region, the minimum population size for that region is given by $3 * h$. This offers a constraint below which values the model
estimated population trajectory must not go. A penalty is added to the negative log likelihood to ensure that these constraints are not violated.

## RESULTS

Table 1(a) gives the posterior median values and $90 \%$ probability intervals for key parameter estimates for the E1+O+G three-stock model. Since Johnston et al. (2011) fit to the Branch (2011) IDCR/SOWER feeding ground relative abundance series in their assessment of BSG, Table 1 (b) reports on the results of the E1+O+G three-stock model where the number of whales feeding in the eastern feeding ground is also fit to the Branch (2011) series. Table 1(c) gives the posterior median values and $90 \%$ probability intervals for the BSE1, BSO and BSG populations from previous assessments: the BSE1 and BSO results are from the $2014 \mathrm{D}+\mathrm{E} 1+\mathrm{O}$ assessment (IWC 2014) and BSG results are from the Johnston et al. (2011) assessment.

Figure 4 is a graphical representation of the $r, K, N_{2015} / K$ parameter estimates for the three stocks from Table 1(a)(c), comparing the median estimates and $90 \%$ probability intervals for the $\mathrm{E} 1+\mathrm{G}+\mathrm{O}$ model, the $\mathrm{E} 1+\mathrm{G}+\mathrm{O}$ model fit to the IDCR/SOWER data as well for the previous assessments. Figure 5 shows the posterior median trajectories and $90 \%$ probability envelopes for the E1+O+G model and for previous assessments.

Table 2 gives the likelihood components (in terms of median values) for the $\mathrm{D}+\mathrm{E} 1+\mathrm{O}$ and the $\mathrm{E} 1+\mathrm{O}+\mathrm{G}$ models. Likelihood components are reported for BSE1 and BSO, i.e. the two stocks common across the two three-stock models.

Table 3 gives the cumulative catches allocated in the models to each breeding stock.

## DISCUSSION

The main point of interest is how compatible the results of the E1+O+G model are with the $2014 \mathrm{D}+\mathrm{E} 1+\mathrm{O}$ threestock model, in order to assess whether a combined assessment of all the stocks could be undertaken by using successive three-stock models.

As can be seen from Figure 5 (a) and (b), there are substantial differences between the results of the two models. Table 3 helps elucidate the reason underlying these differences, namely that the E1+O+G model allocated substantially more catches to BSE1 and fewer catches to BSO than the D+E1+O model, thus resulting in the higher BSE1 and lower BSO carrying capacity values. This illustrates not only the impact that the catch allocations have on model outputs, but also the impact that the neighbouring populations have on any one breeding stock when catches are shared between neighbouring stocks. For example, the BSE1 population requires a fairly large number of catches to be allocated to it in order for the population to be sufficiently depleted to achieve the high growth rate observed by Noad et al. (2011). For the D+E1+O model, the BSE1 populations shares a total of 63892 catches with a larger BSD population to the west and a smaller BSO population to the east (Figure 2), and as such there is some flexibility in the model for balancing the catch allocation between BSD+BSE1 and BSE1+BSO. For the E1+O+G model, the BSE1 population shares a total of 59321 catches with a smaller BSO population to the east (Figure 2) and all the BSE1 catches are those shared with the western BSO population. While there are many intricate interacting factors that play a role in the final model outputs, it is clear that the different set-up of the two threestock models can easily lead to appreciably different results. Interestingly, the likelihoods in Table 2, while showing a small difference for the BSE1 fit to the Noad et al. (2011) relative abundance data, otherwise do not differ greatly amongst the models, suggesting that the data do not provide sufficient information to clearly prefer one model to the other.

Some thought could be given to developing a gradient-like boundary, rather than the hard boundaries from Figure 1 and Figure 2, so that the catches from outside the current boundaries can still be allocated to the western- and eastern-most stocks of the three-stock model and perhaps produce more compatible results when the three-stock model is shifted from one group of neighbouring stocks to the next.

## ACKNOWLEDGEMENTS

Computations were performed using facilities provided by the University of Cape Town's ICTS High Performance Computing team: http://hpc.uct.ac.za. Funding support for this work from the International Whaling Commission is gratefully acknowledged.

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Table 1 (a): Posterior median values of key model parameters are given with their $90 \%$ probability intervals for the E1+O+G threestock model. Results are from a re-sample of 1000 from an initial sample of 2000000 . The model was fit to the Noad et al. (2011) absolute and relative abundance estimates for BSE1, the Constantine et al. (2012) mark-recapture data for BSO and the Felix et al. (2011) absolute abundance estimate for BSG.

|  | BSE1 |  |  | BSO |  | BSG |  |
| :--- | ---: | :--- | ---: | :--- | ---: | :--- | :---: |
| $r$ | 0.105 | $[0.103,0.106]$ | 0.056 | $[0.010,0.087]$ | 0.055 | $[0.006,0.103]$ |  |
| K | 36682 | $[29092,38717]$ | 5700 | $[2861,15644]$ | 8629 | $[5620,18096]$ |  |
| $V$ | 0.359 | $[0.043,0.798]$ | (Proportion of BSO feeding in western feeding area) |  |  |  |  |
| $\mathrm{N}_{\text {min }}$ | 227 | $[204,259]$ | 529 | $[175,2033]$ | 1470 | $[233,5676]$ |  |
| $\mathrm{N}_{2015}$ | 21765 | $[19760,23727]$ | 3579 | $[2818,5285]$ | 7254 | $[5274,9591]$ |  |
| $\mathrm{N}_{\text {min }} / \mathrm{K}$ | 0.006 | $[0.005,0.008]$ | 0.098 | $[0.024,0.273]$ | 0.163 | $[0.034,0.431]$ |  |
| $\mathrm{N}_{2015} / \mathrm{K}$ | 0.602 | $[0.544,0.709]$ | 0.713 | $[0.223,0.998]$ | 0.921 | $[0.375,1.000]$ |  |
| $\mathrm{N}_{2020} / \mathrm{K}$ | 0.811 | $[0.760,0.885]$ | 0.818 | $[0.240,0.999]$ | 0.961 | $[0.379,1.000]$ |  |
| $\mathrm{N}_{2040} / \mathrm{K}$ | 0.999 | $[0.999,1.000]$ | 0.986 | $[0.304,1.000]$ | 0.998 | $[0.435,1.000]$ |  |

Table 1 (b): Repeat of the model run from Table 1, except that in addition to the input data described there, the eastern feeding ground population $\left(1-\gamma^{O}\right) * N^{B S O}+N^{B S G}$ was also fit to the Branch (2011) feeding ground relative abundance series.

|  | BSE1 |  | BSO |  | BSG |  |
| :--- | ---: | :--- | ---: | :--- | ---: | :--- |
| $r$ | 0.105 | $[0.103,0.106]$ | 0.067 | $[0.036,0.087]$ | 0.068 | $[0.030,0.105]$ |
| $K$ | 37328 | $[31185,38724]$ | 4469.0 | $[2897,10743]$ | 8109 | $[5840,10913]$ |
| $\gamma$ | 0.357 | $[0.041,0.764]$ | (Proportion of BSO feeding in western feeding area) |  |  |  |
| $\mathrm{N}_{\text {min }}$ | 226 | $[202,257]$ | 335 | $[175,790]$ | 520 | $[225,2268]$ |
| $N_{2015}$ | 21719 | $[19909,23796]$ | 3726 | $[2844,5182]$ | 7280 | $[5534,8703]$ |
| $N_{\text {min }} / K$ | 0.006 | $[0.005,0.008]$ | 0.077 | $[0.025,0.169]$ | 0.066 | $[0.033,0.210]$ |
| $N_{2015} / K$ | 0.594 | $[0.532,0.674]$ | 0.859 | $[0.416,0.996]$ | 0.944 | $[0.644,1.000]$ |
| $N_{2020} / K$ | 0.805 | $[0.749,0.865]$ | 0.931 | $[0.510,0.999]$ | 0.975 | $[0.696,1.000]$ |
| $N_{2040} / K$ | 0.999 | $[0.998,0.999]$ | 0.998 | $[0.840,1.000]$ | 0.999 | $[0.899,1.000]$ |

Table 1 (c): Posterior median values from previous assessments are given with their $90 \%$ probability intervals. For BSE1 and BSO these previous assessment results are from the $2014 \mathrm{D}+\mathrm{E} 1+\mathrm{O}$ three-stock model (IWC 2014), which was fit to Bannister and Hedley (2001) and Hedley et al. (2011) relative abundance series for BSD, the Noad et al. (2011) absolute and relative abundance estimates for BSE1 and the Constantine et al. (2012) mark-recapture data for BSO. The BSG previous assessment results are from Johnston et al. (2011), where the model was fit to the Felix et al. (2011) absolute abundance estimate and the Branch (2011) IDCR/SOWER feeding ground relative abundance series.

|  | BSE1 |  |  | BSO |  | BSG |  |
| :--- | ---: | :--- | ---: | :--- | ---: | :--- | :---: |
| $r$ | 0.105 | $[0.103,0.106]$ | 0.091 | $[0.071,0.101]$ | 0.063 | $[0.023,0.093]$ |  |
| K | 26133 | $[21605,29033]$ | 14115 | $[10198,19651]$ | 11584 | $[10590,14878]$ |  |
| $\gamma$ | 0.068 | $[0.007,0.190]$ | (Proportion of BSE1 feeding in western feeding area) |  |  |  |  |
| $\mathrm{N}_{\text {min }}$ | 237 | $[203,272]$ | 132 | $[103,250]$ | 731 | $[238,2959]$ |  |
| $\mathrm{N}_{2015}$ | 19614 | $[17643,21454$ | 6404 | $[5491,7595]$ | 9173 | $[6829,10857]$ |  |
| $\mathrm{N}_{\text {min }} / \mathrm{K}$ | 0.009 | $[0.008,0.011]$ | 0.010 | $[0.007,0.014]$ | 0.063 | $[0.022,0.198]$ |  |
| $\mathrm{N}_{2015} / \mathrm{K}$ | 0.762 | $[0.692,0.841]$ | 0.466 | $[0.291,0.657]$ | 0.792 | $[0.512,0.956]$ |  |
| $\mathrm{N}_{2020} / \mathrm{K}$ | 0.915 | $[0.872,0.950]$ | 0.648 | $[0.409,0.846]$ | 0.885 | $[0.575,0.988]$ |  |
| $\mathrm{N}_{2040} / \mathrm{K}$ | 1.000 | $[0.999,1.000]$ | 0.993 | $[0.926,0.999]$ | 0.994 | $[0.818,1.00]$ |  |

Table 2: Likelihood components for BSE1 and BSO (the two stocks common across the two three-stock models) are given in terms of the posterior median values. Likelihood values are given for the 2014 D+E1+O three stock model, the current $\mathrm{E} 1+\mathrm{O}+\mathrm{G}$ three stock model as well as the $\mathrm{E} 1+\mathrm{O}+\mathrm{G}$ three-stock model where the eastern feeding ground population is fit to the IDCR/SOWER relative abundance trend.

| BSE1 | D+E1+O model | E1+O+G model | E1+O+G model, <br> fitting to IDCR |
| :--- | ---: | ---: | ---: |
| Absolute abundance estimate (Noad et al. 2011) | 0.2531 | 0.2046 | 0.2445 |
| Relative abundance series (Noad et al. 2011) | -39.61 | -41.37 | -41.52 |
|  |  |  | E1+O+G model, <br> fitting to IDCR |
| BSO | D+E1+O model | E1+O+G model | -59.17 |
| Mark-recapture (Constantine et al. 2012) | -58.81 | -59.04 | -104 |

Table 3: Posterior medians of the cumulative catches allocated in the model to the various breeding stocks for (i) the E1+O+G three stock model, (ii) the E1+O+G three stock model where the IDCR/SOWER feeding ground estimates (Branch 2011) have been included in the likelihood fit and (iii) the $2014 \mathrm{D}+\mathrm{E} 1+\mathrm{O}$ three stock model. The $5^{\text {th }}$ and $95^{\text {th }}$ percentiles have been given in parenthesis.

|  |  | BSD | BSE1 |  | BSO |  | BSG |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| (i) $\mathrm{E} 1+\mathrm{O}+\mathrm{G}$ |  | - | 41284 | [31271, 43875] | 7702 | [2542, 21791] | 10392 | [5729, 13340] |
| (ii) E1+O+G (with IDCR/SOWER) |  | - | 42048 | [31271, 43875] | 7278 | [2542, 21791] | 9991 | [5729, 13340] |
| (iii) $\mathrm{D}+\mathrm{E} 1+\mathrm{O}$ | 19967 | [16821, 21782] | 27657 | [20473, 33809] | 16439 | [11707, 22771] |  | - |



Figure 4: Graphical representation of the $r, K, N_{2015} / K$ parameter estimates from Table 1. Median estimates are indicated by points and the $90 \%$ probability intervals by bars. Estimates are shown for all three stocks for (a) the E1+O+G three stock model, (b) the E1+O+G three stock model where the IDCR/SOWER feeding ground estimates (Branch 2011) have been included in the likelihood fit and (c) previous assessments (the 2014 D+E1+O model for BSE1 and BSO, and the Johnston et al. 2011 assessment for BSG).


Figure 5: Posterior median population trajectories and their $90 \%$ probability envelopes (PE) for the three stocks. Trajectories for the $\mathrm{E} 1+\mathrm{O}+\mathrm{G}$ three-stock model are shown by the solid lines and their $90 \%$ PEs by the grey shaded areas. Posterior median trajectories from previous assessments are shown by dashed lines and their $90 \%$ P.E. by the areas filled with diagonals. For BSE1 and BSO these previous assessment results are from the 2014 D+E1+O three-stock model. The BSG previous assessment results are from Johnston et al. (2011). Fits to various input data are indicated in the legends.


Figure 6: Nucleus and Margin regions associated with each of the seven breeding stocks according to Hypothesis 1 (map adapted from IWC, 2010).

## APPENDIX A: CATCH, ABDUNANCE AND TREND DATA

Table A. 1: Historical catches taken north of $40^{\circ} \mathrm{S}$ from Allison's database (C.Allison, pers. commn). The catches for E1 and Oceania are available by landing station. Catches landed at LochTay, Tangalooma and Byron Bay have been allocated to BSE1. Catches landed at Norfolk Island, Tonga and Polynesia have been allocated to the Oceania breeding stock. Catches landed at Rakiura and Prince George, New Zealand, Kaikoura, Cook Strait and Tory Channel, Great Barrier Island, Whangamumu and Bay of Island are allocated to BSE1 and Oceania in proportion to the model-predicted population abundances (labelled BSE1/BSO in the table). Catches labelled "Chile", "Ch/Per/E" and "Peru" in the database have been allocated to BSG.

| Year | BSE1 | BSO | $\begin{gathered} \text { BSE1/ } \\ \text { BSO } \end{gathered}$ | BSG | Year | BSE1 | BSO | $\begin{gathered} \text { BSE1/ } \\ \text { BSO } \end{gathered}$ | BSG |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1890 | 0 | 0 | 8 | 0 | 1935 | 0 | 0 | 57 | 31 |
| 1891 | 0 | 0 | 8 | 0 | 1936 | 0 | 0 | 69 | 18 |
| 1892 | 0 | 0 | 8 | 0 | 1937 | 0 | 0 | 55 | 28 |
| 1893 | 0 | 0 | 8 | 0 | 1938 | 0 | 0 | 75 | 6 |
| 1894 | 0 | 0 | 8 | 0 | 1939 | 0 | 0 | 80 | 7 |
| 1895 | 0 | 0 | 8 | 0 | 1940 | 0 | 0 | 107 | 0 |
| 1896 | 0 | 0 | 8 | 0 | 1941 | 0 | 0 | 86 | 0 |
| 1897 | 0 | 0 | 8 | 0 | 1942 | 0 | 0 | 71 | 0 |
| 1898 | 0 | 0 | 8 | 0 | 1943 | 0 | 0 | 90 | 0 |
| 1899 | 0 | 0 | 8 | 0 | 1944 | 0 | 0 | 88 | 0 |
| 1900 | 0 | 0 | 8 | 0 | 1945 | 0 | 0 | 107 | 0 |
| 1901 | 0 | 0 | 8 | 0 | 1946 | 0 | 0 | 110 | 15 |
| 1902 | 0 | 0 | 8 | 0 | 1947 | 0 | 0 | 101 | 19 |
| 1903 | 0 | 0 | 8 | 0 | 1948 | 0 | 0 | 92 | 5 |
| 1904 | 0 | 0 | 8 | 0 | 1949 | 0 | 3 | 141 | 6 |
| 1905 | 0 | 0 | 8 | 0 | 1950 | 0 | 0 | 79 | 5 |
| 1906 | 0 | 0 | 8 | 0 | 1951 | 0 | 0 | 111 | 26 |
| 1907 | 0 | 0 | 8 | 0 | 1952 | 600 | 0 | 121 | 27 |
| 1908 | 0 | 0 | 8 | 16 | 1953 | 700 | 0 | 109 | 29 |
| 1909 | 0 | 0 | 16 | 44 | 1954 | 718 | 0 | 180 | 106 |
| 1910 | 0 | 0 | 77 | 62 | 1955 | 720 | 0 | 112 | 7 |
| 1911 | 0 | 0 | 77 | 92 | 1956 | 720 | 150 | 143 | 10 |
| 1912 | 0 | 0 | 93 | 86 | 1957 | 721 | 136 | 184 | 5 |
| 1913 | 348 | 0 | 92 | 45 | 1958 | 720 | 136 | 183 | 0 |
| 1914 | 0 | 0 | 93 | 195 | 1959 | 810 | 166 | 318 | 3 |
| 1915 | 0 | 0 | 106 | 30 | 1960 | 810 | 186 | 361 | 2 |
| 1916 | 0 | 0 | 82 | 15 | 1961 | 731 | 186 | 80 | 3 |
| 1917 | 0 | 0 | 94 | 15 | 1962 | 173 | 4 | 32 | 4 |
| 1918 | 0 | 0 | 90 | 23 | 1963 | 0 | 0 | 9 | 1 |
| 1919 | 0 | 0 | 119 | 24 | 1964 | 0 | 0 | 0 | 35 |
| 1920 | 0 | 0 | 107 | 21 | 1965 | 0 | 0 | 0 | 143 |
| 1921 | 0 | 0 | 89 | 21 | 1966 | 0 | 0 | 0 | 58 |
| 1922 | 0 | 0 | 57 | 19 | 1967 | 0 | 0 | 0 | 0 |
| 1923 | 0 | 0 | 79 | 16 | 1968 | 0 | 0 | 0 | 3 |
| 1924 | 0 | 0 | 107 | 34 | 1969 | 0 | 0 | 0 | 1 |
| 1925 | 0 | 0 | 96 | 248 | 1970 | 0 | 0 | 0 | 0 |
| 1926 | 0 | 0 | 78 | 277 | 1971 | 0 | 0 | 0 | 0 |
| 1927 | 0 | 0 | 127 | 40 | 1972 | 0 | 0 | 0 | 0 |
| 1928 | 0 | 0 | 105 | 36 | 1973 | 0 | 3 | 0 | 0 |
| 1929 | 0 | 0 | 102 | 26 | 1974 | 0 | 4 | 0 | 0 |
| 1930 | 0 | 0 | 78 | 33 | 1975 | 0 | 8 | 0 | 0 |
| 1931 | 0 | 0 | 109 | 53 | 1976 | 0 | 4 | 0 | 0 |
| 1932 | 0 | 0 | 18 | 21 | 1977 | 0 | 4 | 0 | 0 |
| 1933 | 0 | 0 | 44 | 11 | 1978 | 0 | 11 | 0 | 0 |
| 1934 | 0 | 0 | 52 | 13 | Total | 7771 | 1001 | 5690 | 2119 |

 western feeding ground and catches taken between $100^{\circ} \mathrm{W}$ and $70^{\circ} \mathrm{W}$ were allocated to the eastern feeding ground. Catches in the column 'S. Shet/Falk' are $100 \%$ of catches taken between $70^{\circ} \mathrm{W}$ and $50^{\circ} \mathrm{W}$ and south of $60^{\circ} \mathrm{S}$ as well as $50 \%$ of catches taken between $70-50^{\circ} \mathrm{W}$ and $50-60^{\circ} \mathrm{S}$. These catches were allocated to the eastern feeding ground.

| Year | $\begin{aligned} & \hline 120- \\ & 129 \mathrm{E} \end{aligned}$ | $\begin{aligned} & \hline 130- \\ & 139 \mathrm{E} \end{aligned}$ | $\begin{aligned} & \hline 140- \\ & 149 \mathrm{E} \end{aligned}$ | $\begin{aligned} & \hline 150- \\ & 159 \mathrm{E} \end{aligned}$ | $\begin{aligned} & \hline 160- \\ & 169 \mathrm{E} \end{aligned}$ | $\begin{aligned} & \hline 170- \\ & 180 \mathrm{E} \end{aligned}$ | $\begin{aligned} & \hline 180- \\ & 170 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 169- \\ & 160 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 159- \\ & 150 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 149- \\ & 140 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 139- \\ & 130 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 129- \\ & 120 \mathrm{~W} \\ & \hline \end{aligned}$ | $\begin{aligned} & \hline 119- \\ & 110 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & 109- \\ & 100 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 99- \\ & 90 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 89- \\ & 80 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 79- \\ & 70 \mathrm{~W} \end{aligned}$ | S. Shet/ Falk |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1903 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 |
| 1904 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1905 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 23 |
| 1906 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 498 |
| 1907 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 366 |
| 1908 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1250.5 |
| 1909 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1528 |
| 1910 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 2562 |
| 1911 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 2047.5 |
| 1912 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 980 |
| 1913 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1041.5 |
| 1914 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 662 |
| 1915 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 219 |
| 1916 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 21 |
| 1917 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 69 |
| 1918 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 81 |
| 1919 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 181 |
| 1920 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 149 |
| 1921 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1922 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 189 |
| 1923 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 96 |
| 1924 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 102 |
| 1925 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 163 |
| 1926 | 0 | 0 | 0 | 0 | 0 | 82 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 101 |
| 1927 | 0 | 0 | 0 | 0 | 0 | 16 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 3 |
| 1928 | 0 | 0 | 0 | 0 | 0 | 17 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 16 |
| 1929 | 0 | 0 | 0 | 0 | 0 | 775 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1930 | 1 | 0 | 32 | 49 | 3 | 55 | 96 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 |
| 1931 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1932 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1933 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1934 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1935 | 0 | 0 | 4 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1936 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 |


| Year | $\begin{aligned} & 120- \\ & 129 \mathrm{E} \\ & \hline \end{aligned}$ | $\begin{aligned} & 130- \\ & 139 \mathrm{E} \end{aligned}$ | $\begin{aligned} & 140- \\ & 149 \mathrm{E} \end{aligned}$ | $\begin{aligned} & 150- \\ & 159 \mathrm{E} \end{aligned}$ | $\begin{aligned} & \hline 160- \\ & 169 \mathrm{E} \end{aligned}$ | $\begin{aligned} & 170- \\ & 180 \mathrm{E} \end{aligned}$ | $\begin{aligned} & \hline 180- \\ & 170 \mathrm{~W} \\ & \hline \end{aligned}$ | $\begin{aligned} & 169- \\ & 160 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & 159- \\ & 150 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 149- \\ & 140 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & 139- \\ & 130 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 129- \\ & 120 \mathrm{~W} \\ & \hline \end{aligned}$ | $\begin{aligned} & 119- \\ & 110 \mathrm{~W} \end{aligned}$ | $\begin{aligned} & \hline 109- \\ & 100 \mathrm{~W} \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { 99- } \\ & 90 \mathrm{~W} \\ & \hline \end{aligned}$ | $\begin{aligned} & \hline 89- \\ & 80 \mathrm{~W} \\ & \hline \end{aligned}$ | $\begin{aligned} & \hline 79- \\ & 70 \mathrm{~W} \\ & \hline \end{aligned}$ | S. Shet/ Falk |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1937 | 32 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1938 | 24 | 24 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1939 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1940 | 342 | 342 | 342 | 342 | 342 | 342 | 342 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 |
| 1941 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1942 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1943 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1944 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1945 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1946 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1947 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1948 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1949 | 109 | 30 | 760 | 118 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1950 | 0 | 0 | 0 | 0 | 0 | 85 | 86 | 316 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 271.8 |
| 1951 | 232 | 0 | 1 | 0 | 66 | 103 | 189 | 37 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1952 | 0 | 0 | 0 | 0 | 166 | 216 | 135 | 13 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1953 | 0 | 0 | 0 | 0 | 0 | 0 | 14 | 136 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1954 | 0 | 2 | 0 | 749 | 5 | 17 | 167 | 269 | 69 | 2 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1955 | 411 | 769 | 416 | 777 | 0 | 0 | 0 | 278 | 56 | 0 | 0 | 0 | 0 | 0 | 0 | 14 | 0 | 0 |
| 1956 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 7 | 3 | 27 | 39 | 571 | 11 | 8 | 37.6 |
| 1957 | 0 | 30 | 19 | 38 | 133 | 0 | 0 | 0 | 35 | 27 | 29 | 76 | 31 | 0 | 37 | 18 | 4 | 0 |
| 1958 | 882.1 | 104.7 | 157.1 | 185.7 | 525.8 | 209.4 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 26.2 |
| 1959 | 44.8 | 1043.5 | 4057.1 | 3673 | 2228.5 | 998.7 | 317.8 | 112.8 | 73.2 | 106.8 | 73.2 | 73.2 | 74.1 | 7 | 14 | 5 | 0 | 182 |
| 1960 | 71 | 163.7 | 742.3 | 1184.3 | 3703.8 | 2630.2 | 740 | 962.5 | 565.27 | 508.3 | 428.57 | 292.87 | 0 | 0 | 0 | 0 | 81 | 10.9 |
| 1961 | 14 | 14 | 61 | 436 | 581 | 342 | 123 | 226 | 1010 | 401 | 452 | 189 | 54 | 44 | 806 | 334 | 24 | 2.5 |
| 1962 | 58.2 | 18.2 | 35.4 | 39.7 | 302.2 | 9.2 | 10 | 49.5 | 87.7 | 66.1 | 63.5 | 18.1 | 18.1 | 24.4 | 36.2 | 70 | 88.1 | 85.5 |
| 1963 | 0.2 | 1.2 | 23.6 | 20.9 | 225 | 13 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1964 | 0.9 | 2.8 | 11.3 | 26.2 | 45 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1965 | 8.8 | 12.6 | 43.6 | 26.6 | 80 | 97.1 | 85.3 | 474.6 | 1.3 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1966 | 7 | 4 | 3 | 1 | 11 | 14 | 16 | 93 | 118 | 26 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1967 | 7 | 0 | 1 | 11 | 12 | 2 | 1 | 6 | 47 | 57 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1968 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1969 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1970 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1971 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 3 | 0 | 0 | 0 | 0 | 0 |
| 1972 | 0 | 0 | 0 | 0 | 2 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Total | 2245 | 2561.7 | 6709.4 | 7677.4 | 8431.3 | 6023.6 | 2322.1 | 2973.4 | 2062.47 | 1195.2 | 1055.27 | 652.17 | 207.2 | 114.4 | 1464.2 | 452 | 205.1 | 12969 |

## Breeding Stock E1

## Absolute abundance estimate

A land-based survey was conducted at Point Lookout on the east coast of Australia over eight weeks in June and July 2010 (Noad et al. 2011). The average number of whales passing per 10h over the peak four weeks of the northward migration was $84.7 \pm 3.2$ whales. A correction for whales available but missed was applied using double blind counts, as well as other corrections for sighting heterogeneity ( $1.212+/-0.049$, Dunlop et al., 2010). Using this correction the abundance estimate for 2010 was 14,522 whales ( $95 \% \mathrm{Cl}^{5} 12,777-16,504$ ) (Noad et al., 2011).

## Relative abundance estimates

Table A. 3: BSE1 Relative Abundance Index I (Noad et al., 2011): A count of northward migrating whales from land-based surveys conducted at Point Lookout and two other locations. The values give the number of whales passing per 10 h during four weeks of the peak migration. (M. Noad, pers. commn) and are as used for estimates of abundance provided by Noad et al., (2008), Noad et al., (2011). These data were used to in estimated annual rate of increase of $10.9 \% /$ year $(95 \% \mathrm{CI}=10.5-11.3 \% /$ year $)$ for a 24 year period ( 1984 to 2010) (Noad et al., 2011).

| Year | Estimate |
| ---: | ---: |
| 1984 | 6.12 |
| 1985 | 5.92 |
| 1986 | 8.25 |
| 1987 | 8.53 |
| 1988 | 9.15 |
| 1989 | 10.22 |
| 1990 | 11.58 |
| 1991 | 12.93 |
| 1992 | 14.36 |
| 1994 | 17.75 |
| 1996 | 20.91 |
| 1998 | 28.45 |
| 1999 | 27.45 |
| 2001 | 34.67 |
| 2002 | 37.34 |
| 2004 | 47.11 |
| 2007 | 70.73 |
| 2010 | 84.7 |

## Minimum number of haplotypes

The minimum number of haplotypes for BSE1 5 (IWC 2014).

[^3]
## Oceania breeding stock

## Absolute abundance estimate

The estimate in 2005 of 4,329 individuals ( $\mathrm{CV}=0.12$ ) arises from a sighting-resighting analysis of microsatellite genotypes collected from 1999 to 2005 across four survey areas in Oceania: New Caledonia (E2), Tonga (E3), the Cook Islands and French Polynesia (F2) (Constantine et al., 2012). It is a doubled male-specific estimate assuming equal numbers of males and females in the region.

## Mark recapture data

Table A. 4: Synoptic genotypic mark recapture data underlying male specific Oceania-wide abundance estimate. This is the males-only subset of the sexes combined dataset from Constantine et al. (2012), as provided by Jackson (pers. commn, 2012).

| Year initial capture (males) | 1999 | 2000 | 2001 | 2002 | 2003 | 2004 | 2005 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| Total individuals captured | 25 | 70 | 112 | 78 | 114 | 24 | 82 |
| 1999 | - | 3 | 4 | 0 | 3 | 0 | 1 |
| 2000 |  | - | 5 | 3 | 8 | 2 | 6 |
| 2001 |  |  | - | 7 | 12 | 3 | 7 |
| 2002 |  |  |  | - | 4 | 0 | 6 |
| 2003 |  |  |  |  | - | 1 | 11 |
| 2004 |  |  |  |  |  | - | 3 |
| 2005 |  |  |  |  |  |  | - |

## Minimum number of haplotypes

The minimum number of haplotypes for Oceania is 33 (IWC 2014).

## BREEDING STOCK G

## Absolute abundance estimate

A breeding ground estimate of $6504(\mathrm{CV}=0.21)$ for the year 2006 is from a photographic capture-recapture study in Ecuador, and is based on Chapman modified-Peterson estimator (Felix et al., 2011).

## Relative abundance estimates

Table A. 5: IDCR/SOWER estimates for the feeding grounds (Branch 2011). The area for G is $110^{\circ} \mathrm{W}-50^{\circ} \mathrm{W}$, south of $60^{\circ} \mathrm{S}$.

| Year | N | CV |
| :---: | :---: | :---: |
| 1982 | 1452 | 0.65 |
| 1989 | 2817 | 0.38 |
| 1996 | 3310 | 0.21 |

## Minimum number of haplotypes

The minimum number of haplotypes for BSG is 27 (Rosenbaum et al. 2006).


[^0]:    ${ }^{1}$ MARAM (Marine Resource Assessment and Management Group), Department of Mathematics and Applied Mathematics, University of Cape Town, Rondebosch, 7701, South Africa

[^1]:    ${ }^{2}$ Note that an importance function was used for $r^{E l}$ to improve sampling efficiency. Details are given later.
    ${ }^{3}$ If $N$ is assumed to be log-normally distributed, then $\ln N$ is normally distributed with some mean $\mu$ and standard deviation $\sigma$. The median value of $N$ is then $e^{\mu}$ while the CV of $N$ is given by $\sqrt{e^{\sigma^{2}-1}}$. Since the CV of $N$ is relatively small, $\sigma$ has been approximated here by the value of the CV of $N$.

[^2]:    ${ }^{4}$ The equations given here imply a multinomial distribution. However, because the annual capture probabilities are so small, the Poisson distribution is an adequate and convenient approximation.

[^3]:    ${ }^{5}$ This $95 \%$ CI was converted into a rough CV by assuming that the estimate was log-normally distributed. An approximation of the standard error of the $\log$ of the estimate was obtained by computing $0.5 *(\ln (16504)-\ln (12777)) / 1.96$. The resulting value of 0.065 was then taken to be the CV of the estimate.

