## POP2012-02

New Zealand sea lion - demographic assessment of the causes of decline at the Auckland Islands

## 7a model results

## CSP Technical Working Group

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## Optimal model configuration

- Annual survival estimates for age groupings 0, 1, 2-5, 6-14, 15+
- Survival at Age 15+ is time-invariant
- All others have separate estimate for years where data informative
- Annual breeding probability for Age 4+ individuals
- Separate estimates for breeders and non-breeders
- All time-varying (1998-2011)
- Annual resighting probability of age groupings 1-2, 3, 4I-5I, 6I, 7I, B, N
- Separate estimates for breeders and non-breeders
- All time varying 1999-2011
- Decline in resighting probability estimated of breeders after mid-2000s suggests a problem as nearly all breeders should be resighted in every year since 1999. This can be fixed to 1 - all resighted.


## Fits to tagging observations - optimal model

| Model run | Survival estimates Age | Survival Yr groups | Breeding Prob estimates Age | Breeding Prob Yr groups | Resighting prob estimates Age | Resighting prob Yr groups | Maturation | LL | params | AIC |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 7 a | 0, 1, 2-5, 6-14, 15+ | 15+ time invariant | 4+(P), 4+(N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 1-2 time invariant | Time varying | -7976.2 | 178 | 16,308 |
| 6 b | 0, 1, 2-5, 6-14, 15+ | 15+ time invariant | $4+(\mathrm{P}), 4+(\mathrm{N})$ | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 1-2 time invariant |  | -8023.6 | 152 | 16,351 |
| 6d | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | functional form | a4 \& b4 time invariant | 1-2,3,4-5,6,7,N | 1-2 time invariant |  | -8022.8 | 154 | 16,354 |
| 6 a | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | $4+(\mathrm{P}), 4-14(\mathrm{~N}), 15+$ (N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 1-2 time invariant |  | -8020.5 | 159 | 16,359 |
| 5 j | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+(P), 15+(N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 1-2 time invariant |  | -8017.1 | 166 | 16,366 |
| 4 m | 0, 1, 2-5, 6-14, 15+ | 0 \& 15+ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -7999.6 | 185 | 16,369 |
| 5 m | 0, 1, 2-5, 6-14, 15+ | $6+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 1-2 time invariant |  | -8032.2 | 153 | 16,370 |
| 6c | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | functional form | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 1-2 time invariant |  | -8019.3 | 166 | 16,371 |
| 51 | 0, 1, 2-5, 6-14, 15+ | $0 \& 15+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 1-2 time invariant |  | -8036.4 | 149 | 16,371 |
| 5d | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+(P), 15+(N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | Separate estimates all yrs |  | -8008.5 | 179 | 16,375 |
| 5 b | 0, 1, 2-5, 6-14, 15+ | 15+ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4-5,6,7,N | Separate estimates all yrs |  | -7999.3 | 192 | 16,383 |
| 5 h | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 4-5 time invariant |  | -8023.8 | 169 | 16,386 |
| $4 i$ | 0, 1, 2-5, 6-14, 15+ | 15+ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -7992.4 | 202 | 16,389 |
| 4k | 0, 1, 2-5, 6-14, 15+ | 2-5 \& 15+ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -8008 | 187 | 16,390 |
| $5 f$ | 0, 1, 2-5, 6-14, 15+ | 15+ time invariant | 4-14 (P), 4-14 (N), 15+(P), 15+(N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 7 time invariant |  | -8025.2 | 170 | 16,390 |
| 5 i | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 3 time invariant |  | -8027.5 | 168 | 16,391 |
| 3 | 0, 1, 2-5, 6-14, 15+ | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -7987.6 | 208 | 16,391 |
| 4 j | 0, 1, 2-5, 6-14, 15+ | $6+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -8007.2 | 189 | 16,392 |
| 5 g | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 6 time invariant |  | -8026.4 | 170 | 16,393 |
| 4 h | 0, 1, 2-5, 6+ | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+(P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -8001.7 | 201 | 16,405 |
| 4 e | 0, 1, 2-4, 5-14, 15+ | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -7995.1 | 208 | 16,406 |
| 4d | 0, 1, 2, 3-5, 6-14, 15+ | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -7981.1 | 222 | 16,406 |
| 5 e | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+(P), 15+ (N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | N time invariant |  | -8038.7 | 166 | 16,409 |
| 4 g | 0, 1, 2-14, 15+ | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+(P), 15+(N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -8010.7 | 194 | 16,409 |
| 5 k | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1-2,3,4-5,6,7,N | 0-7 time invariant |  | -8087.6 | 127 | 16,429 |
| 4c | 0, 1, 2, 3, 4, 5, 6-14, 15+ | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -7977 | 243 | 16,440 |
| 5a | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4-7, N | Separate estimates all yrs |  | -8053.7 | 175 | 16,457 |
| 4 a | u1, u3, u4, max (u3) at age 3 | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+(P), 15+(N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -8140 | 145 | 16,570 |
| 4 b | u1, u3, u4, max (u3) at age 2 | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+(P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -8141.1 | 144 | 16,570 |
| 5 c | 0, 1, 2-5, 6-14, 15+ | $15+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4-6,7,N | Separate estimates all yrs |  | -8411.4 | 182 | 17,187 |
| 4 f | 0-1, 2-5, 6-14, 15+ | Separate estimates all yrs | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -8476.6 | 191 | 17,335 |
| 41 | 0, 1, 2-5, 6-14, 15+ | 1 \& $15+$ time invariant | 4-14 (P), 4-14 (N), 15+ (P), 15+ (N) | Separate estimates all yrs | 1,2,3,4,5,6,7,N | Separate estimates all yrs |  | -8483.1 | 186 | 17,338 |

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## Fits to tagging obs - optimal model



## Pups dead at tagging

－Some pups recorded as dead at the time of tagging，e．g．disease mortalities in 2002 \＆ 2003 －we are overestimating Surv0 if these are not accounted for
－We included additional＂phantom tag＂observations in SeaBird input files－ animals that are tagged and then not observed again
－SeaBird decreases Surv0（and not resighting probability－prob．resight ages $1 \& 2=0.104$ with phantom tags $\& 0.105$ without）．Minor effect on Surv1 in のヘロの



## MCMC runs 400,000 length; 400 samples - survival






## MCMC runs - resighting probability



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## MCMC runs - maturation







## MCMC runs - breeding probability



- Probable cohort effects on breeding probability (1998 \& 2000 cohort)
- Also year effects e.g. 2008 (low survivorship too)


## Retrospective analysis



## Retrospective analysis



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## Tag loss model (tag observations only)

- Partitions for 2 tags, 1 tag and 0 tag (PresightOtag set to zero)
- Retagged animals - assume same tag frequency in all subsequent years
- Two parameters for probability of losing 1 tag in a year (time-invariant):
- $\mathrm{T}_{\text {age } 0}=0.085 ; \mathrm{T}_{\text {age1+ }}=0.049$
- Compares with $0.15(2>1) \& 0.09(1>0)$ MacKenzie \& Chilvers (2012)




## Fitting to pup count observations with tag loss



- C.V. of 0.03
- Good fits (including low pup counts in 2009 - low prob puppers pupping)
- Minimal conflict with estimates from tag only though increased survival of groupings Age2+
- $\mathrm{T}_{\text {ageo }}=0.103 ; \mathrm{T}_{\text {age }}=0.063$



## Fitting to age distribution observations

- Good fits to lactating female age distribution observations 1998 to 2001 (Childerhouse et al 2010)



- Only really informative for survival at ages 0-5
- Confirms strong cohorts (1990-1993) evident from mark-resighting analysis
- Cohort effect on survival of animals - not just tags!


## Summary model development

- Goals of demographic modelling
- Conflict between observations
- More pups born than calculated from tag only survival and pupping rates
- Relocation effects?
- Underestimating tag shedding?
- Biases and uncertainty around estimates
- Final model development


## Proximate causes of "population decline"

Survival
-Decline in survival at ages 0 and 1 relative to strong cohorts in 1990-93
-Some very weak cohorts e.g. 1998 (disease epidemic year) with reduced survival at different ages
-2008 may be another very weak cohort
-Cohort (e.g. 1998) \& year (e.g. 2007) effects on adult survival
-A need to better understand biology of juvenile animals - low resighting probability at breeding rookeries ( $\sim 0.1$ per annum)

## Pupping

-Year effects on pupping rates, e.g. 2009, cause single-year declines in pup production. Also potential cohorts effects.

- Increased age at first pupping during period of decline \& low proportion pupping each year for an otariid species (0.6-0.7)
-Do non-pupping animals breed then abort? Do they skip breeding/suckle yearlings?


## References

Childerhouse, S. J., Dawson, S. M., Slooten, E., Fletcher, D. J., Wilkinson, I. S. (2010). Age distribution of lactating New Zealand sea lions: Interannual and intersite variation. Marine Mammal Science, 26: 123-139.

Gilbert, D.J., Chilvers B.L. (2008). Final report on New Zealand sea lion pupping rate. POP2006-01. Objective 3. Analysis from sea lion database to estimate pupping rate and associated parameters.

MacKenzie, D.I. (2012). Estimation of Demographic Parameters for New Zealand Sea Lions Breeding on the Auckland Islands - Final Report: 1997/982010/11. Objective 3: POP2010/1

