Energetic Consequences of Sonar Exposure for Cetaceans

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Abstract

The sub-lethal effects of sonar exposure for cetaceans is not well enough understood to predict population consequences. To better inform conservation planning, we developed a model to predict the energetic costs of sonar avoidance behavior. When avoiding sonar sources, cetaceans may cease foraging and/or flee the area. We estimate the potential energy intake lost to foraging cessation and the additional locomotor costs from increased swim speeds.

Although medium-sized cetaceans, especially beaked whales, make up the majority of mass strandings associated with naval sonar, our results indicate sublethal effects cause the most harm to large baleen whales. A severe response by a Cuvier's beaked whale (6.5 hour foraging cessation and 30 minutes of elevated swim speeds at $4.5~\mathrm{m/s}$) cost 49% of its daily basal metabolic requirements. Conversely, a moderate response by a blue whale (1 hour foraging cessation and 5 minutes of elevated swim speeds at $2.5~\mathrm{m/s}$) cost more than nine times its daily basal metabolic requirements. These results may be used to reduce harm to endangered cetacean species.

1 Introduction

- Naval exercises involving sonar have been linked to mass strandings of
- cetaceans worldwide since at least the 1980's (England et al., 2001; Frantzis, 1998;
- 4 Jepson et al., 2003; Simmonds and Lopez-Jurado, 1991). However, the population
- 5 consequences of sub-lethal effects are not as well understood. Controlled exposure
- 6 experiments (CEEs) show that behavioral responses may include cessation of
- ⁷ feeding and/or fleeing the sonar source at elevated speed (DeRuiter et al., 2017,

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2013; Friedlaender et al., 2016; Goldbogen et al., 2013; Kvadsheim et al., 2017;
Southall et al., 2019; Tyack et al., 2011; Wensveen et al., 2019). Quantitatively
linking these behaviors to demographics requires an understanding of the impacts
on individuals' health (Pirotta et al., 2018). The mechanism addressed here is
reduced energy stores due to lost foraging opportunities and increased locomotor
costs.

The two extant clades of cetaceans differ in their feeding styles. Toothed whales (odontocetes) are raptorial feeders and locate prey using echolocation [ref?]. Baleen whales (mysticetes) are filter feeders and capture prey by lunging (Goldbogen et al., 2012) or continuous ram filter feeding (Werth and Potvin, 2016). These feeding styles have profound effects on feeding rate, energy per feeding event, dive depth, and body size. Odontocetes feed at higher rates on smaller prey and most larger odontocetes must dive to extreme depths to find sufficient prey [ref?]. Lunge-feeding mysticetes (rorquals) engulf enormous quantities of prey-laden water, increasing the energy intake per feeding event but limiting dive depth and duration (Goldbogen et al., 2012).

Feeding rates and the energy obtained per feeding event can be empirically measured with multi-sensor tags, active acoustics, and stomach contents [jeremy's paper?]. The rapid echolocation clicking (buzzes) preceding odontocete prey capture events have a kinematic and acoustic signature that register on tag sensors [ref?]. Similarly, rorqual lunges can be identified from an increase in speed followed by a rapid deceleration, usually associated with high pitch and roll angles (Cade et al., 2016). Simultaneous prey mapping with tagging efforts using active acoustics have measured the biomass density of fish and krill schools targeted by rorquals. The sizes of squid beaks and fish otoliths (?) in odontocete stomach samples are a record of prey sizes and energy per feeding event.

Empirical estimates of cetacean metabolic rates are logistically challenging for

smaller species and infeasible for larger species. Oxygen consumption has been 35 measured for captive odontocetes trained to swim under a metabolic hood using 36 open-flow respirometry, showing that mass-specific stroke costs are largely size 37 invariant (Williams et al., 1993, 2017). Whether these metabolic estimates scale 38 to larger odontocetes or mysticetes is unknown, so other methods of estimating 39 energy expediture include breathing rates and hydrodynamic models (Potvin et al., 2012; Sumich, 1983). Animals swim efficiently by cruising at 1-2 m/s and maintaining a Strouhal 42 number of 0.25 - 0.3 (Katsufumi et al., 2007; Rohr and Fish, 2004). The Strouhal number is a dimensionless quantity $St = \frac{Af}{U}$ where A is stroke amplitude, f 44 is stroke frequency, and U is swimming speed. Cetacean stroke amplitudes are approximately one fifth body length [ref?] so there is a linear relationship between swimming speed and stroke frequency for animals of a given body size [gough?].

49 Methods

We considered the potential energy intake of lost feeding opportunities and additional energy expenditure from elevated swimming speeds in modeling the energetic consequences of sonar exposure. The model takes the form:

$$E_{sonar} = P_{in} \times t_d + P_{out}(U_f) \times t_f \tag{1}$$

Where E_{sonar} is the energy cost of sonar exposure, P_{in} is consumption power (i.e. rate of energy intake) during undisturbed foraging, t_d is the time displaced from foraging, P_{out} is flight power (i.e. increased rate of locomotor costs), U_f is the animal's speed during flight, and t_f is the flight time. P_{in} and P_{out} are species-specific values and t_d , U_f , and t_f are dependent on the individual's

- behavioral response to sonar exposure.
- Consumption power (P_{in})
- The rate of energy intake is the product of feeding rate (r_f) and prey energy per feeding event (E_p) :

$$P_{in} = r_f \times E_p \tag{2}$$

 r_f was calculated as the lunge rate for rorquals and buzz rate for odontocetes using tag sensors. E_p was derived using active acoustics (rorquals) and stomach contents (odontocetes) [ref?].

65 Flight power (P_{out})

The locomotor costs associated with fleeing the sonar source is the energetic cost of swimming at U_f relative to the cruising swim speed. Using the relationship between stroke frequency and swimming speed and a scaling relationship for mass-specific stroke costs, we calculate P_{out} as:

$$P_{out}(U_f) = (f_s(U_f) - f_s(U_c)) \times C_L \times m \tag{3}$$

Where f_s is a function relating stroke frequency to swimming speed, U_f and U_c are the swimming speeds during the flight response and cruising, C_L is the mass-specific locomotor cost of a stroke, and m is the animal's mass. Equation (2) assumes, during the flight response, cetaceans increase swimming speed and stroke frequency, but the mass-specific locomotor cost of a stroke remains the same regardless of speed. Although C_L increases with swimming speed, the scaling relationships do not hold for larger cetaceans (Williams et al., 2017). To

be conservative, we use the C_L scaling relationship for cruising speeds, estimated as $C_L = 1.46 + 0.0005m$ in $J \cdot stroke^{-1} \cdot kg^{-1}$. We chose 1.5 m/s for U_c based on size-invariant scaling of cruising speeds (Katsufumi et al., 2007). Assuming cetaceans maintain a Strouhal number of 0.3 and stroke amplitude is one fifth body length, stroking frequency as a function of body length is:

$$S_t = \frac{Af}{U}$$

$$f = \frac{1.5U}{L}$$
(4)

82 Case studies

We analyzed tag data from four controlled exposure experiments (CEEs) and applied the E_{sonar} model to estimate energetic costs of observed behavioral 84 responses to sonar exposure [BRS ref?]. As part of an on-going behavioral response study to naval sonar, in 2011-2015 (???) Blainville's beaked whales 86 Mesoplodon densirostris), Cuvier's beaked whales (Ziphius cavirostris), northern 87 minke whales (Balaenoptera acutorostrata), and blue whales (B. musculus) were tracked with multi-sensor tags and exposed to mid-frequency active sonar in 89 Southern California. Time displaced from foraging (t_d) , time in flight (t_f) , and 90 speed of flight (U_f) were selected based on tag data. These values should be 91 considered realistic scenarios, but not the typical or most common response. 92 Behavioral responses to sonar exposure are highly variable and seem to depend 93 on received sound level, distance to source, foraging behavior, and habituation (Southall et al., 2019; Wensveen et al., 2019).

96 Cross-species comparisons

To facilitate comparisons of energetic consequences across body size, we present E_{sonar} as 1) the energy cost (kJ), 2) the mass-specific energy cost

Table 1: Cetacean feeding rates

Group	Species	r_f	N
Odontocete	Phocoena phocoena	96.8	8
Odontocete	Grampus griseus	17.7	17
Odontocete	$Mesoplodon\ densirostris$	13.0	14
Odontocete	$Globicephala\ macrorhynchus$	2.5	2
Odontocete	$Globicephala\ melas$	7.9	9
Odontocete	Orcinus orca	9.7	10
Odontocete	$Ziphius\ cavirostris$	12.8	4
Odontocete	Berardius bairdii	4.6	1
Odontocete	$Physeter\ macrocephalus$	12.5	36
Rorqual	$Balaen optera\ bonaerensis$	44.1	3
Rorqual	$Megaptera\ novae angliae$	17.2	6
Rorqual	$Balaen optera\ physalus$	13.3	3
Rorqual	$Balae noptera\ musculus$	10.7	11

Note:

 r_f is lunges/hour (rorquals) or buzzes/hour (odon tocetes)

Species ordered by size within groups.

 99 $(kJ \ kg^{-1})$, and 3) the ratio of energy cost to daily basal metabolic requirements $(E_{sonar}:BMR)$. Kleiber's equation for mammals predicts basal metabolic rate as: $BMR=293.1m^{0.75}$ (kJ/day) (Kleiber, 1975). Given the uncertainty in BMR estimates for large cetaceans and the differences between basal and active metabolic rates, $E_{sonar}:BMR$ should be interpreted as an index of energetic consequence and not an absolute measure of impact.

105 Results

- Consumption power (P_{in})
- Feeding rates were derived from tag data and tended to decrease with body size. Rorqual feeding rates ranged from 10.7 to 44.1 lunges/hour and odontocete feeding rates ranged from 2.5 to 96.8 buzzes/hour (Table 1).
- Prey energy per feeding event was empirically derived from acoustic backscatter (mysticetes) and stomach samples (odontocetes). Filter feeders consumed the

- most energy per feeding event. Generally, delphinids targeted more energy-rich prey than Ziphids (Fig. 1).
- Modeled consumption power covered four orders of magnitude: from 2.4e3 kJ/hr (*P. phocoena*) to 1.4e7 kJ/hr (*B. musculus*) (Fig. 2).
- Flight power (P_{out})
- Predicted stroke frequencies at cruising speed decreased with length, from
- 1.8 Hz for a 1.22 m *P. phocoena* to 0.09 for a 25 m *B. musculus* (Fig. 3).
- Across body sizes, P_{out} increased linearly with flight speed (Fig. 4).
- 120 Case studies
- We modeled the energetic consequence of sonar exposure for four observed
- behavioral responses. The beaked whale ($M.\ densirostris$ and $Z.\ cavirostris$)
- responses were more severe than the rorquals' (B. bonaerensis and B. musculus)
- but the mass-specific energetic costs and E_{sonar} : BMR ratio were greater for
- 125 rorquals (Table 2).

Table 2: Behavioral responses to sonar and estimated energetic consequences of exposure.

	Behavioral responses			Energetic consequences				
Species	$t_d \text{ (min)}$	t_f (min)	$U_f \text{ (m/s)}$	E_{out} (kJ)	E_{in} (kJ)	E_{sonar} (kJ)	E_{sonar} $(kJ$ $kg^{-1})$	$E_{sonar}:BMR_d$
M. densirostris	360	30	4.5	3210	20500	23700	27.5	0.509
Z. cavirostris	360	30	4.5	10400	46700	57000	19.7	0.492
B. bonaerensis	150	60	3.5	44600	949000	994000	148.0	4.58
B. musculus	60	5	2.5	79600	14200000	14300000	153.0	9.14

Note:

 t_d is the time displaced from foraging and t_f is the time fleeing the sonar source at the elevated speed, U_f . E_{out} and E_{in} are the energetic costs of increased locomotion and lost feeding, respectively. E_{sonar} is presented as a total cost, as a mass-specific cost, and as a ratio to daily basal metabolic requirements.

Figures Figures

127 References

- ¹²⁸ Cade, D.E., Friedlaender, A.S., Calambokidis, J., Goldbogen, J.A., 2016.
- 129 Kinematic diversity in rorqual whale feeding mechanisms. Current Biology 26,
- ¹³⁰ 2617–2624. https://doi.org/10.1016/j.cub.2016.07.037
- DeRuiter, S.L., Langrock, R., Skirbutas, T., Goldbogen, J.A., Calambokidis,
- ¹³² J., Friedlaender, A.S., Southall, B.L., 2017. A multivariate mixed hidden markov
- model for blue whale behaviour and responses to sound exposure. The Annals
- of Applied Statistics 11, 362–392. https://doi.org/10.1214/16-AOAS1008
- DeRuiter, S.L., Southall, B.L., Calambokidis, J., Zimmer, W.M.X., Sadykova,
- D., Falcone, E.A., Friedlaender, A.S., Joseph, J.E., Moretti, D., Schorr, G.S.,
- 137 Thomas, L., Tyack, P.L., 2013. First direct measurements of behavioural
- responses by cuvier's beaked whales to mid-frequency active sonar. Biology
- Letters 9, 20130223-20130223. https://doi.org/10.1098/rsbl.2013.0223
- England, G.R., Evans, D., Lautenbacher, C., Morrissey, S., Hogarth, W.,
- others, 2001. Joint interim report bahamas marine mammal stranding event of
- 15-16 march 2000. US Department of Commerce, US Secretary of the Navy.
- Frantzis, A., 1998. Does acoustic testing strand whales? Nature 392, 29–29.
- 144 https://doi.org/10.1038/32068
- Friedlaender, A.S., Hazen, E.L., Goldbogen, J.A., Stimpert, A.K., Calam-
- bokidis, J., Southall, B.L., 2016. Prey-mediated behavioral responses of feeding
- blue whales in controlled sound exposure experiments. Ecological Applications
- ¹⁴⁸ 26, 1075–1085. https://doi.org/10.1002/15-0783
- Goldbogen, J.A., Calambokidis, J., Croll, D.A., McKenna, M.F., Oleson,
- E., Potvin, J., Pyenson, N.D., Schorr, G., Shadwick, R.E., Tershy, B.R., 2012.
- Scaling of lunge-feeding performance in rorqual whales: Mass-specific energy
- expenditure increases with body size and progressively limits diving capacity.

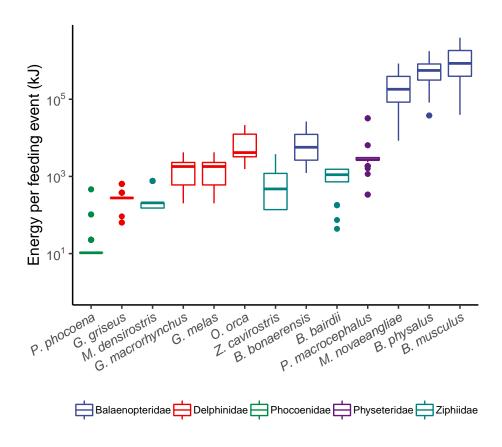


Figure 1: Energy per feeding event. Note \log scale on y-axis.

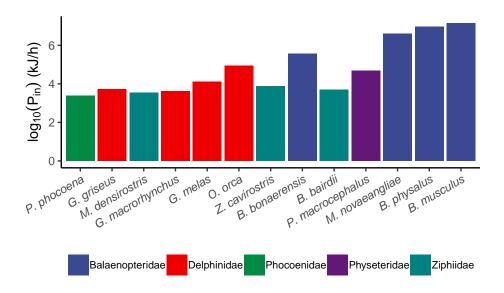


Figure 2: Modeled consumption power (P_{in}) .

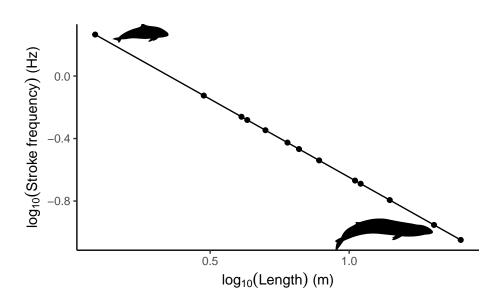


Figure 3: Predicted cruising speed stroke frequencies (f_s) .

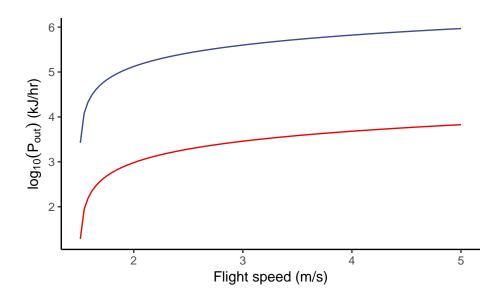


Figure 4: Flight power (P_{out}) by flight speed for B. musculus (blue) and Z. cavistrostris (red). Note log scale on y-axis.

Functional Ecology 26, 216–226. https://doi.org/10.1111/j.1365-2435.2011.01905.

Goldbogen, J.A., Southall, B.L., DeRuiter, S.L., Calambokidis, J., Friedlaen-

der, A.S., Hazen, E.L., Falcone, E.A., Schorr, G.S., Douglas, A., Moretti, D.J.,

Kyburg, C., McKenna, M.F., Tyack, P.L., 2013. Blue whales respond to simulated

mid-frequency military sonar. Proceedings of the Royal Society B: Biological

Sciences 280, 20130657–20130657. https://doi.org/10.1098/rspb.2013.0657

Jepson, P.D., Arbelo, M., Deaville, R., Patterson, I.A.P., Castro, P., Baker,

J.R., Degollada, E., Ross, H.M., Herráez, P., Pocknell, A.M., Rodríguez, F.,

Howie, F.E., Espinosa, A., Reid, R.J., Jaber, J.R., Martin, V., Cunningham,

A.A., Fernández, A., 2003. Gas-bubble lesions in stranded cetaceans. Nature

¹⁶⁴ 425, 575–576. https://doi.org/10.1038/425575a

156

Katsufumi, S., Yutaka, W., Akinori, T., J.O, M.P., Hideji, T., Ryo, K., J,

66 P.P., Yves, H., Tomonari, A., Yuuki, W., Yoko, M., P, C.D., Charles-André, B.,

- Kagari, A., Masao, A., Phil, T., Ari, S., Yasuhiko, N., 2007. Stroke frequency, but
- not swimming speed, is related to body size in free-ranging seabirds, pinnipeds
- and cetaceans. Proceedings of the Royal Society B: Biological Sciences 274,
- ¹⁷⁰ 471–477. https://doi.org/10.1098/rspb.2006.0005
- Kleiber, M., 1975. The fire of life: An introduction to animal energetics.
- Robert E. Krieger, Huntington.
- Kvadsheim, P.H., DeRuiter, S., Sivle, L.D., Goldbogen, J., Roland-Hansen,
- R., Miller, P.J., Lam, F.-P.A., Calambokidis, J., Friedlaender, A., Visser, F.,
- Tyack, P.L., Kleivane, L., Southall, B., 2017. Avoidance responses of minke
- whales to 1-4 kHz naval sonar. Marine Pollution Bulletin 121, 60-68. https:
- 177 //doi.org/10.1016/j.marpolbul.2017.05.037
- Pirotta, E., Booth, C.G., Costa, D.P., Fleishman, E., Kraus, S.D., Lusseau,
- D., Moretti, D., New, L.F., Schick, R.S., Schwarz, L.K., Simmons, S.E., Thomas,
- L., Tyack, P.L., Weise, M.J., Wells, R.S., Harwood, J., 2018. Understanding the
- population consequences of disturbance. Ecology and Evolution 8, 9934–9946.
- 182 https://doi.org/10.1002/ece3.4458
- Potvin, J., Goldbogen, J.A., Shadwick, R.E., 2012. Metabolic expenditures
- of lunge feeding rorquals across scale: Implications for the evolution of filter
- feeding and the limits to maximum body size. PLOS ONE 7, e44854. https:
- 186 //doi.org/10.1371/journal.pone.0044854
- Rohr, J.J., Fish, F.E., 2004. Strouhal numbers and optimization of swimming
- by odontocete cetaceans. Journal of Experimental Biology 207, 1633–1642.
- 189 https://doi.org/10.1242/jeb.00948
- Simmonds, M.P., Lopez-Jurado, L.F., 1991. Whales and the military. Nature
- 191 351, 448. https://doi.org/10.1038/351448a0
- Southall, B.L., DeRuiter, S.L., Friedlaender, A., Stimpert, A.K., Goldbogen,
- 193 J.A., Hazen, E., Casey, C., Fregosi, S., Cade, D.E., Allen, A.N., Harris, C.M.,

- Schorr, G., Moretti, D., Guan, S., Calambokidis, J., 2019. Behavioral responses
- of individual blue whales (balaenoptera musculus) to mid-frequency military
- sonar. The Journal of Experimental Biology 222, jeb190637. https://doi.org/10.
- 197 1242/jeb.190637
- Sumich, J.L., 1983. Swimming velocities, breathing patterns, and estimated
- costs of locomotion in migrating gray whales, eschrichtius robustus. Canadian
- ²⁰⁰ Journal of Zoology 61, 647–652. https://doi.org/10.1139/z83-086
- Tyack, P.L., Zimmer, W.M.X., Moretti, D., Southall, B.L., Claridge, D.E.,
- Durban, J.W., Clark, C.W., D'Amico, A., DiMarzio, N., Jarvis, S., McCarthy, E.,
- Morrissey, R., Ward, J., Boyd, I.L., 2011. Beaked whales respond to simulated
- and actual navy sonar. PLoS ONE 6, e17009. https://doi.org/10.1371/journal.
- 205 pone.0017009
- Wensveen, P.J., Saana, I., Hansen, R.R., Benda-Beckmann, A.M. von,
- Kleivane, L., IJsselmuide, S. van, Lam, F.-P.A., Kvadsheim, P.H., DeRuiter, S.L.,
- ²⁰⁸ Curé, C., Narazaki, T., Tyack, P.L., Miller, P.J., 2019. Northern bottlenose
- whales in a pristine environment respond strongly to close and distant navy sonar
- 210 signals. Proceedings of the Royal Society B: Biological Sciences 286, 20182592.
- 211 https://doi.org/10.1098/rspb.2018.2592
- Werth, A.J., Potvin, J., 2016. Baleen hydrodynamics and morphology of cross-
- flow filtration in balaenid whale suspension feeding. PLOS ONE 11, e0150106.
- 214 https://doi.org/10.1371/journal.pone.0150106
- Williams, T.M., Friedl, W.A., Haun, J.E., 1993. The physiology of bottlenose
- dolphins (tursiops truncatus): Heart rate, metabolic rate and plasma lactate
- concentration during exercise. Journal of Experimental Biology 179, 31–46.
- Williams, T.M., Kendall, T.L., Richter, B.P., Ribeiro-French, C.R., John,
- J.S., Odell, K.L., Losch, B.A., Feuerbach, D.A., Stamper, M.A., 2017. Swimming
- 220 and diving energetics in dolphins: A stroke-by-stroke analysis for predicting

- the cost of flight responses in wild odontocetes. The Journal of Experimental
- ${}^{222} \quad \text{Biology 220, 1135-1145. https://doi.org/10.1242/jeb.154245}$