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Trade-offs between foraging efficiency and pup feeding rate of lactating northern fur seals in a declining population

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ABSTRACT: Foraging strategies and their resulting efficiency (energy gain to cost ratio) affect animals' survival and reproductive success and can be linked to population dynamics. However, they have rarely been studied quantitatively in free-ranging animals. We investigated foraging strategies and efficiencies of wild northern fur seals *Callorhinus ursinus* during their breeding season to understand potential links to the observed population decline in the Bering Sea. We equipped 20 lactating females with biologgers to determine at-sea foraging behaviours. We measured energy expenditure while foraging using the doubly-labelled water method, and energy gained using (1) the types and energy densities of prey consumed, and (2) the number of prey capture attempts (from acceleration data). Our results show that seals employed 2 foraging strategies: one group (40%) fed mostly in oceanic waters on small, high energy-density prey, while the other (60%) stayed over the shallow continental shelf feeding mostly on larger, lower quality fish. Females foraging in oceanic waters captured 3 times more prey, and had double the foraging efficiencies of females that foraged on-shelf in neritic waters. However, neritic seals made comparatively shorter trips, and likely fed their pups ~20 to 25% more frequently. The presence of these strategies which either favor foraging efficiency (energy) or frequency of nursing (time) might be maintained in the population because they have similar net fitness outcomes. However, neither strategy appears to simultaneously maximize time and energy allocated to nursing, with potential impacts on the survival of pups during their first year at sea.

KEY WORDS: Foraging efficiency · Reproduction success · Northern fur seal · Pup growth · Diet

INTRODUCTION

A central concept in life-history theory is that organisms must optimally allocate resources between the competing demands of reproduction, body maintenance, and growth in order to maximize survival and reproductive success. Changes in environmental conditions that result in sub-optimal energy intake will likely result in animals modifying how they allocate energy. In long-lived animals, self-maintenance has priority over reproduction because fitness, and lifetime reproductive success depend on longevity

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(Bouten et al. 1994). Hence, reductions in energy intake or foraging efficiency are expected to affect breeding performances before they affect adult survival. However, the extent to which reproduction is impaired by food availability depends in part on the flexibility and capacity of animals to acquire energy (foraging efficiency) and allocate it (physiological plasticity). Therefore, the extent to which an animal can adjust to localized environmental changes and the availability and distribution of prey will determine its body condition, fitness and capacity to survive and successfully reproduce.

Numbers of northern fur seals *Callorhinus ursinus* breeding in the central Bering Sea (St. Paul and St. George Islands) have declined by 3.5% yr^{-1} for the last 20 yr (Muto et al. 2016), as have populations of other top predators in this area (Wanless et al. 2005). Female fur seals sustain high energy output while nursing their pups for 4 mo by making frequent foraging trips at sea. The breeding season is a crucial period in the life history of otariids, when foraging success of mothers determines the quantity of energy available to transfer to their pups and ensure their post-weaning survival (Reid & Forcada 2005). These marine predators use areas of high productivity that are linked to higher and more predictable prey concentrations (Costa 1993).

In the Bering Sea, major environmental changes starting in the mid-1970s altered the distribution and relative abundances of different species of fish (Conners et al. 2002, Grebmeier et al. 2006). The main prey available to top marine predators in the Bering Sea switched from high to low energy density prey (Trites et al. 2007, Sinclair et al. 2008). Free-ranging predators usually buffer changes in the availability of prey by changing their foraging areas. However, during the breeding season, mothers leave their pups on shore while they forage at sea, and the limited fasting abilities of a small, growing pup puts a cap on the upper end of how long the mother can spend at sea. The fasting constraints of pups and their need to grow rapidly strongly constrain the foraging range of lactating northern fur seals (central place foragers). Consequently, localised oceanographic variations within the foraging range of northern fur seals during the breeding season likely affect foraging success of fur seals and the rearing of their pups.

When faced with environmental changes, animals have to employ strategies that balance the energetic costs and time spent foraging, against the potential energy gained from prey. Foraging efficiency is the difference between the energy gained through prey intake and the energy spent to extract the prey from the environment. A decline in foraging efficiency of lactating fur seals could explain reduced reproductive success, and thus overall fitness, of the breeding females in the central Bering Sea compared to pre-1970s — and consequently the declining population trend of northern fur seals. There is therefore a need to determine, at the finest scale possible, the foraging and feeding behaviours of individual fur seals to determine links between specific environmental parameters, fitness, and demographic parameters.

Estimating foraging efficiency requires measuring foraging costs and energy gain at the same time. Traditionally, foraging efficiency has not been measured, but merely inferred by comparing the change in mass before and after a foraging trip (Gentry & Kooyman 1986, Lea et al. 2006, Luque et al. 2007), which lacks fine-scale precision (Insley et al. 2008). It is rare that both sides of the 'efficiency equation' have been studied on the same individuals (Costa 1993). The doubly-labelled water (DLW) method has been used and validated to estimate the field metabolic rate (FMR) of pinnipeds (Speakman 1997, Butler et al. 2004, Sparling et al. 2008), but has been seldom used on northern fur seals since the 1980s (Costa & Gentry 1986, Costa 1987).

Feeding success, or the energy gained while foraging, is more difficult to measure than energy expenditure. Proxies for feeding success such as changes in mass and body composition, changes in buoyancy during drift dives, transit rate, dive shape and bout structures have all been tried and validated to some extent on marine predators under certain conditions (Hindell et al. 2010), but none of these techniques provide a quantitative measure of energy gain itself. Changes in stomach temperatures have been used in seals and birds to detect feeding events (Grémillet & Plös 1994, Kuhn & Costa 2006), but have been less successful in wild otariids that tend to regurgitate the devices before going to sea. However, the development of high sampling rate sensors such as tri-axial accelerometers can provide indices of prey capture attempts in marine mammals. The use of these indices, when paired with an estimate of diet composition, allows calculation of quantitative estimates of energy gained during a feeding trip. Together with measures of foraging effort, foraging efficiency of individuals can then be calculated.

Comparing foraging efficiencies relative to foraging strategies can further shed light on the trade-offs animals make between the 2 main currencies in animal bioenergetics: time and energy. It is these trade-offs that likely impact the energy reserves that mothers have to feed their pups, which in turn influence

the chances of pups surviving their first year at sea (Doidge & Croxall 1989, Baker & Fowler 1992, Boltnev et al. 1998, McMahon et al. 2000, Hall et al. 2001, Beauplet et al. 2005). In this context, the objectives of our study were (1) to determine the foraging strategies of lactating northern fur seals, including locations, diving patterns and targeted prey, (2) to measure foraging efficiency (foraging gain/cost ratio) and determine how it relates to foraging strategy, and (3) to assess how different foraging strategies and efficiencies might be related to reproductive success and the declining trend of the fur seal population breeding in the central Bering Sea.

MATERIALS AND METHODS

Data collection

Data were collected from 20 lactating northern fur seals at Reef Rookery on St. Paul Island (Bering Sea, 57° 6' N to 170° 17' W) during the breeding season (Aug–Sep 2011) under US National Marine Fisheries Service (NMFS) permit #14329-01 and University of British Columbia (UBC) animal care permit #A10-0364. Data collection procedures were similar to those reported in Jeanniard du Dot et al. (2017b). In short, females were captured using a hoop net, anesthetized (gas anesthesia with isoflurane), measured and weighed, and equipped with biologgers. They were also injected with DLW, and had blood samples drawn to measure metabolic rates (see details below). Animals were released upon full recovery from anesthesia, and recaptured after a single foraging trip at sea using the same methods.

Energy expenditure, foraging behaviours and prey capture attempts

Based on GPS tracks and on stable isotope $\delta^{13}C$ and $\delta^{15}N$ values (which indicate foraging habitat and trophic level, respectively) measured from these same animals for another study (Jeanniard du Dot et al. 2017a), 8 of the 20 tagged females went to the oceanic waters to forage in the deep basin of the Bering Sea (up to 3000 m deep), hereafter called 'oceanic' females, and the remaining 12 stayed over the shallow Bering Sea shelf (0 to 200 m deep), hereafter called 'neritic' females (Fig. 1). Analyses and calculations of the energy expenditure, foraging behaviours and prey capture attempts are detailed in Jeanniard du Dot et al. (2017c) and Jeanniard du Dot

et al. (2017b). In brief, measurements of daily energy expenditure (DEE, kJ d^{-1}) were performed using the DLW method, where a weighed dose of DLW (0.3 to 0.6 g kg^{-1} body mass) was intravenously injected via catheter into the hind-flipper before a second blood sample was taken following a 2-h period of equilibration. A final blood sample was drawn to determine isotope levels of ^2H and ^{18}O at the end of the foraging trip upon recapture. We used a 2-pool model and a plateau method to calculate initial dilution spaces, and the equation from Speakman et al. (1993) to account for evaporative water loss when calculating metabolic, as well as a respiratory quotient of 0.80 to convert CO_2 production rates into DEE (Sparling et al. 2008, Dalton et al. 2014). Energy spent during the 'non-foraging' time was corrected by subtracting on-land energy expenditure from the total estimate using previously determined values specifically for lactating northern fur seals (4.67 W kg^{-1}, in Gentry & Kooyman 1986). Error in individual DEE for each seal was calculated by generating 1000 values following a normal distribution of 1.8 ± 7.2% of the measured values of DLW, based on the error estimated by Dalton et al. (2014) when DLW was compared to respirometry on northern fur seals.

The fur seals were also equipped with Daily Diary tags (DDT, Wildlife Computers) recording tri-axial acceleration and tri-axial magnetic field at 16 Hz and depth, light level, and water temperature at 1 Hz. This tag was positioned as close as possible to the center of mass of the animal to optimize measurements of the overall dynamic body acceleration (Halsey et al. 2009), which has been calculated for these animals in another study. We used depth data to determine diving behaviours using a custom-made R programme previously developed for Antarctic fur seals. Dives were defined as periods of time that animals spent under water below a minimum depth of 3 m and for a minimum of 4 s until they returned to the surface. Dives were classified as 'benthic' when animals dove to 95% of the bathymetric depth. Fast-loc GPS MK10 loggers (Wildlife Computers) were glued slightly below the DDT along the spine and recorded GPS coordinates along the track of the animal at sea, as well as depth and water temperature at 1 Hz. Finally, GCDC X6 or X8 accelerometers glued to the fur on the head of the animals recorded tri-axial acceleration at 16 or 20 Hz to measure prey capture attempts (PrCA). PrCA have been traditionally measured using head accelerometers (Viviant et al. 2010, Gallon et al. 2013); however, we found that back acceleration can provide an accurate estimate of PrCA providing a correction factor is used (Jean-
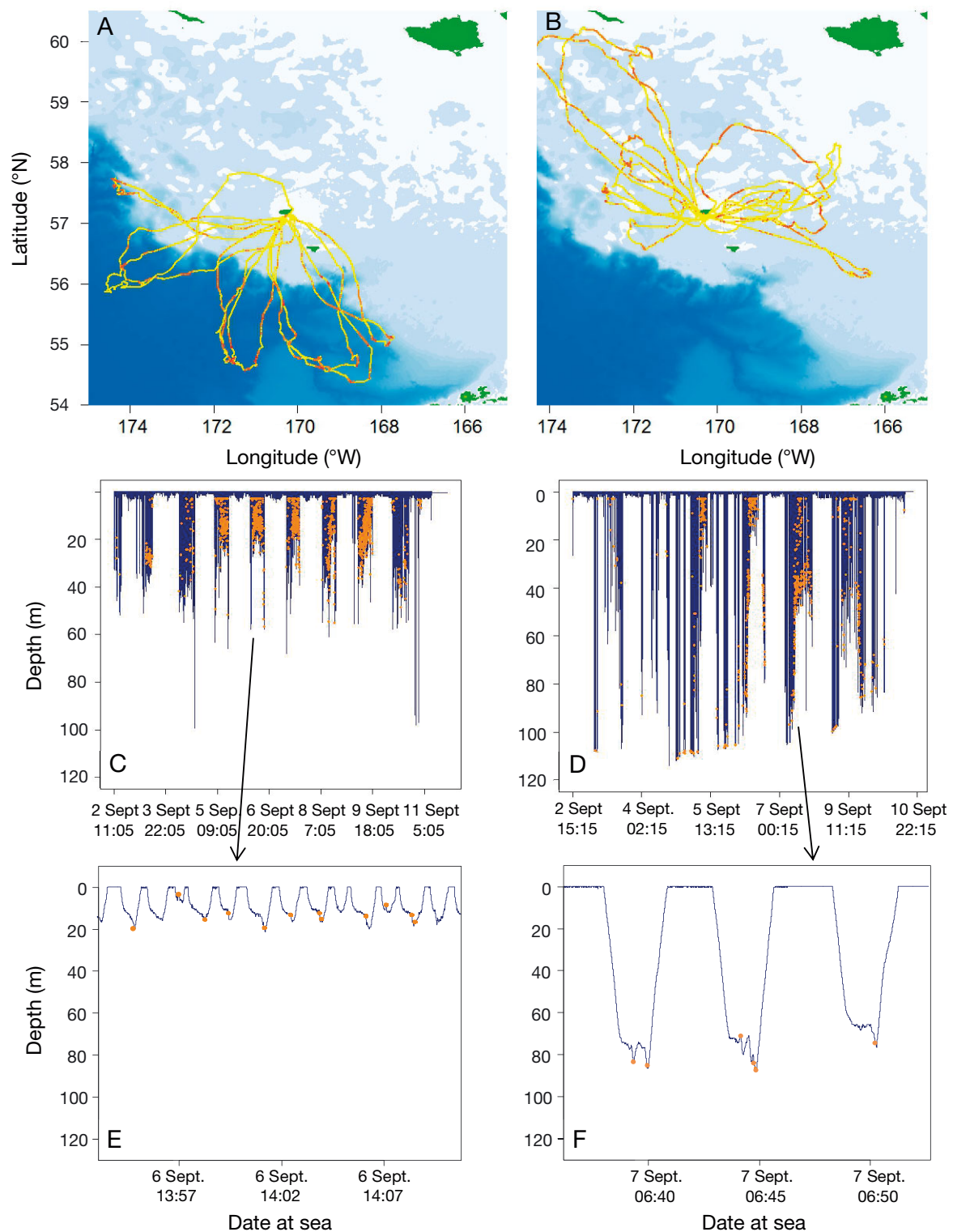
Fig. 1. (A,B) Foraging tracks and dive profiles over a (C,D) full foraging trip, and (E,F) a 5-min period of diving for lactating northern fur seals from Reef Rookery on St. Paul Island in the Bering Sea. Fur seals foraged either off the shelf in oceanic waters (left panels, n = 8) or on the shelf in neritic waters (right panels, n = 12). Orange dots show spatial distribution of prey capture attempts in Panels A and B (with yellow representing no prey capture attempt) and distribution of prey capture attempts over time and depth profiles in Panels C–F

niard du Dot et al. 2017b), as has also been found to be the case for elephant seals (C. Guinet pers. obs.). PrCA were measured while animals were diving below 3 m depth using acceleration data recorded on the head or back of animals using the same method as described in Jeanniard du Dot et al. (2017b). In brief, the heave (z) and surge (x) raw acceleration channels were filtered using a 3rd order high-pass filter at 3 Hz. The resulting dynamic acceleration signals were then summed and a running variance was applied over a 2-s window on which a cluster analysis was applied (k-mean function in R, 2 clusters), which provided each animal with an individual threshold, above which the signal corresponded to a PrCA. PrCA represents all attempts at capturing prey, i.e. success could not be verified. We nevertheless used PrCA as a proxy for feeding rate (Volpov et al. 2015) while accounting for error (under- or overestimation of effective prey captures) in individual PrCA for each seal. The detection rate (true positive rate) of PrCA is known to range from 68 to 97% (i.e. differnce from 100% representing the degree of underestimation of true PrCA), while the false positive rate ranges from 6 to 48% (representing the degree of overestimation of true PrCA) in Steller sea lions and Australian fur seals (Viviant et al. 2010, Volpov et al. 2015). We thus modeled uncertainties around the measured individual PrCA value using these true positive and false positive errors between the ranges mentioned above generated by a uniform distribution using a Monte Carlo approach (1000 values generated).

Each seal performed a different number of dives per trip. The dive depth and dive duration for each individual was calculated as an average over their foraging trip (dives are nested within individuals). To account for the multiple measures within an individual over the trip and obtain means for the group weighted by number of dives that each individual performed, the statistical difference between groups (oceanic versus neritic) for dive depth and dive duration were calculated using linear mixed effect models. We used group as a fixed effect and individual as a random effect. Model assumptions were verified and any autocorrelation or heterogeneity of variances of within-group residuals were corrected as outlined in Pinheiro & Bates (2000). Statistical differences between groups in other parameters (that were single measures or counts over the foraging trips, or measures normalized by a time period) from Table 1 were tested with 2-sample *t*-tests ($\alpha = 0.05$) or Mann-Whitney tests depending on normality.

Diet estimates

Based on analyses of scats, of $\delta^{13}C$ and $\delta^{15}N$ values in blood and of GPS locations of the females we tracked (detailed in Jeanniard du Dot et al. 2017a), northern fur seals were found to have 2 different diets depending on their foraging strategy, i.e. whether they fed on-shelf in neritic waters or off-shelf in oceanic waters. We thus assigned one or the other of these 2 diet compositions to individual females based on their respective foraging strategies. Stable isotope $\delta^{13}C$ and $\delta^{15}N$ values of the oceanic females of this study are not significantly different from the values found for females breeding on Bogoslof Island in the southern Bering Sea (see Fig. 3 in Zeppelin & Orr 2010 and Fig. 2 in Jeanniard du Dot et al. 2017a), which indicate similar foraging habitat and diet. We thus assumed that oceanic females fed on a diet similar to the one estimated on Bogoslof Island in 2009 (Trites et al. 2015), as thoroughly discussed in Jeanniard du Dot et al. (2017a). We assigned the diet composition determined by scats collected on reef rookery in 2011 to neritic females as scats represent prey consumed within the previous 24 to 48 h, i.e. on the shelf. To determine the size of prey consumed, we used scat hard-part remains collected on Reef Rookery in the 2011 group (representative of prey sizes that particular year) and hard parts recovered from scats collected at Bogoslof Island in 2009 whenever fish specific to the oceanic diet were not recovered in the collected scats on Reef Rookery in 2011 (Trites et al. 2015, Jeanniard du Dot et al. 2017a).

For each prey species found in the 2 diets (*i*), the energy density (*ED_i*) expressed in wet mass was estimated from published values (Perez 1994, Van Pelt et al. 1997, Paul & Paul 1999, Logerwell & Christiansen 2000, Iverson et al. 2002, Gende et al. 2004, Andrews et al. 2009, Whitman 2010, Vollenweider et al. 2011). Energy density of walleye pollock differs by age class (Paul & Paul 1999), so we obtained the overall energy density of pollock by accounting for the proportion of each age class (determined from their size) in the diet, i.e. 85% of age-0 at 4.00 kJ g^{-1} and 15% of age-1+ at 4.73 kJ g^{-1} (Whitman 2010, Heintz et al. 2013). Salmonids were identified at the species level from the DNA meta-barcoding method that provides diet information at a finer taxonomic level than the hard part method. We thus calculated the proportion of each salmon species in the overall diet of fur seals. Then, the energy density of salmon species found in Alaskan waters (Hendry & Berg 1999, Anthony et al. 2000, Rosen & Trites 2000, Gende et al. 2004) was averaged and weighted by the proportion of different

Table 1. Mean foraging and metabolic parameters of female northern fur seals (NFS) foraging in neritic waters on the Bering Sea shelf (n = 12) or in oceanic waters in the Bering Sea basin (n = 8) for 1 trip at sea. Values are mean ± SE, and letters in superscript indicate significant differences between the groups (p < 0.05). Values of similar parameters for female Antarctic fur seals (AFS) are added for comparison (from Jeanniard du Dot et al. 2017b). EE: energy expenditure; PrCA: prey capture attempts; ED: energy density; EC_p: energy content prey; EGain: energy gain

Parameter	Neritic NFS	Oceanic NFS	AFS
Mass pre-departure (kg)	39.4 ± 2.0[a]	35.7 ± 1.2[a]	31.1 ± 0.9[b]
Trip duration (d)	7.1 ± 2.4 [a]	9.7 ± 1.8[b]	7.6 ± 3.8[a]
Distance travelled (km)	655 ± 63 [a]	892 ± 51[b]	635 ±77[a]
Number of dives	1831 ± 374[a]	3631 ± 313[b]	3949 ± 597[b]
Dive depth (m)	26.8 ± 3.3[a]	16.9 ± 2.0[b]	19.9 ± 2.7[b]
Dive duration (s)	79.8 ± 8.5[b]	37.3 ± 6.4[a]	42.6 ± 4.5[a]
Benthic dives (%)	15.2 ± 2.6[a]	2.2 ± 0.4[b]	0 ± 0[b]
Time diving (%)	33.1 ± 3.1[a]	24.7 ± 0.8[b]	29.0 ± 0.7[a]
Time transiting (%)	28.3 ± 2.3	32.0 ± 2.2	26.4 ± 1.6
Time diving (h)	55.8 ± 7.9	55.1 ± 3.5	51.3 ± 5.9
Time transiting (h)	49.0 ± 7.2[a]	72.2 ± 7.8[b]	49.8 ± 7.9[a]
At-sea metabolic rate (MJ d^{-1} kg^{-1})	0.56 ± 0.05	0.54 ± 0.07	0.59 ± 0.04
Total EE (MJ kg^{-1})	3.7 ± 0.4	4.9 ± 0.8	4.1 ± 0.6
EE diving (MJ kg^{-1})	2.0 ± 0.3	2.0 ± 0.2	2.2 ± 0.3
EE transiting (MJ kg^{-1})	1.0 ± 0.2[a]	1.6 ± 0.2[b]	1.3 ± 0.2[ab]
PrCA per minute of diving	0.43 ± 0.37 [a]	1.87 ± 38[b]	1.37 ± 0.22[b]
PrCA per capturing dive	1.77 ± 0.23	1.75 ± 0.17	2.04 ± 0.11
PrCA d^{-1}	100 ± 26[a]	275 ± 39[b]	336 ± 38[b]
PrCA per trip	789 ± 163[a]	2404 ± 338[b]	2328 ± 387[b]
ED$_{Diet}$ (kJ g^{-1})	4.28 ± 0.72	5.75 ± 1.44	7.75 ± 2.47
EC_p (kJ)	330.60 ± 11.20	194.94 ± 8.13	152.46 ± 1.08
EGain per dive (kJ)	179.6 ± 49.8	188.5 ± 34.4	177.7 ± 21.4
EGain per time diving (kJ min^{-1})	84.2 ± 16.5[a]	145.9 ± 24.4[b]	130.6 ± 16.3[b]
EGain per time at sea (kJ min^{-1})	26.2 ± 6.2[a]	35.21 ± 5.4[b]	37.6 ± 4.6[b]
Foraging efficiency	1.51 ± 0.69[a]	3.03 ± 0.60 [b]	3.44 ± 0.45[b]

species identified in the same scats with the DNA meta-barcoding method (Jeanniard du Dot et al. 2017a) to obtain an overall salmonid energy density representative of the fur seal diet. The variations in prey body mass and energy density values were estimated from their reported ranges converted as mean ± SD. Whenever information was missing for prey with low frequencies of occurrence in the diet, we replaced it with the energy density of the closest related prey item or with the average of the energy content for the specific prey group.

We calculated energy density in diet (ED_{Diet}) by averaging ED_i values, weighted by their proportion of the corresponding prey item within the diet (P_i), over the number of prey in the diet (N):

$$ED_{Diet} = \frac{\sum (ED_i \times P_i)}{N} \qquad (1)$$

We then estimated the gross energy intake (kJ) for each prey ingested by deriving the size of prey ingested from hard-part remains in the collected scats. We relied on all identifiable bones to estimate number of prey items in each scat, but only used unbroken

otoliths in good or fair condition to estimate fish length (Tollit et al. 2004). Lengths of otoliths were measured using the method described in Harvey et al. (2000), and correction factors were applied to these measured lengths to account for digestion whenever a species-specific correction factor was available (i.e. 1.1691 for Atka mackerel and 1.1593 for walleye pollock; Tollit et al. 2004). Measured otoliths of the same size (±0.1 mm) were paired and assumed to come from the same fish (Hunt et al. 1980, Dragoo 1991).

Squid beaks were measured using guidelines from Wolff (1984) but we were not able to identify them to the species level (including using the DNA meta-barcoding method; see Jeanniard du Dot et al. 2017a). As northern fur seals are known to mostly consume squids from the gonatid family in the Bering Sea (*Gonatopsis borealis* and *Berryteuthis magister*; Gudmundson et al. 2006) we used the general size equation available for this family (Clarke 1962, 1986) with no correction factor (Tollit et al. 1997).

Fish otolith lengths (OL) or squid lower beak rostrum lengths (LRL, mm) were converted to fish/squid length (FL, cm), and then body mass (BM, g) using the species-specific allometric equations listed in the

section 'Equations to estimate body length and mass of fish and squid' in the Supplement at www.int-res.com/articles/suppl/m600p207_supp.pdf. For species of fish found in the scats that were identified by other hard parts than otoliths (vertebrae, gill rakers, etc.), the size of hard parts was compared with those of reference skeleton from Pacific ID Inc. and allocated to a range of sizes summarized in both Table 1 and Fig. 6 of Trites & Calkins (2008). We estimated average size ± SE of these fish species by simulating random selection of fish sizes from the range of sizes in each category using a uniform distribution (1000 simulations).

Once the mass (BM_i, g) and energy density (ED_i, kJ g^{-1}) of each prey item were estimated, the average energy content of a prey from a specific species (EC_i, kJ) was calculated as:

$$EC_i = BM_i \times ED_i \tag{2}$$

and the average energy content of a non-specific 'average' prey (EC_p, kJ) in the diets of neritic and oceanic fur seals was calculated by weighting the energy content of a specific prey item by its relative proportion in the diet (P_i):

$$EC_p = \sum (EC_i \times P_i) \tag{3}$$

We calculated mean ± SD of EC_i by generating 1000 values of mass and 1000 values of ED_i using normal distributions of their respective means ± SD. Means ± SD of ED_{Diet}, and EC_p in the 2 diets were calculated by generating values of EC_i and ED_i for each prey type in proportion to their respective importance in the diet (i.e. P_i) out of 1000 values from normal distributions using their respective mean ± SD. Prey species with no ED_i or EC_i values were substituted by the average ED or EC from their respective prey group (e.g. gadid, cephalopod, etc.) weighted by their own proportion within the diet. As the prey group 'other' did not have values for mass or energy density, we used an average of the rest of the diet weighted by its relative importance in the diet for prey species in this group.

Foraging efficiency

For each seal (s), the foraging efficiency (FE_s) was calculated using the method detailed in (Jeanniard du Dot et al. 2017b). In brief, FE_s is the ratio between the daily energy expenditure at sea for each animal (DEE_s) obtained from the DLW measures and energy gained while foraging at sea. For each seal, we estimated energy gained as the energy content of an 'average' prey in the diet (i.e. EC_p) multiplied by the number of times the seal attempted to capture a prey ($PrCA_s$):

$$FE_s = \frac{EC_p \times PrCA_s}{DEE_s} \tag{4}$$

There are a number of potential biases that could affect our analyses. These include the potential for DLW to either over- or under-estimate metabolic rate at the individual level, and for there to be errors in estimated diet composition, fish size and energy density from scat hard parts due to differential digestion and retention rates. Similarly, $PrCA_s$ values reflect capture attempts and not actual captures. As a result of such potential errors, we applied correction factors when available, and included uncertainties in the final calculations. Uncertainty around FE_s was calculated using 1000 Monte Carlo simulations that reflected errors in the 3 equation parameters, i.e. the error in individual DEE_s, EC_p and $PrCA_s$ for each seal, as mentioned in the respective sections above and described in detail in Jeanniard du Dot et al. (2017b). Mean ± SD of FE_s for each seal was calculated over the 1000 generated $PrCA_s$, DEE_s and EC_p values using Eq. (4). We calculated uncertainty over the average FE using the bootstrap method over 1000 simulations, where the random sampling with replacement was drawn from the 1000 values of FE_i generated per animal in the study.

A sensitivity analysis was performed to estimate the contribution of each of the input variables uncertainty to the overall variance in the resulting foraging efficiency. This was done by computing the standardized regression coefficients (SRC), bias and 95% confidence intervals for each of the input variables using the src function in R (*sensitivity* library, R3.0.3) over 1000 simulated values.

RESULTS

Foraging behaviours

The foraging tracks of the 8 oceanic females and 12 neritic females are shown on Fig. 1. Overall, females weighed (±SE) 37.9 ± 1.34 kg prior to departure (range 30.8–55.6 kg, n = 20). Twelve females gained mass (0.7–5.2 kg gained), and 8 lost mass during their foraging trip (0.2–4.2 kg lost). This corresponds to an overall gain of 3.5 ± 1.8% of initial body mass for the 20 females (range from 9.9% lost to 16.1% gained, and an average net gain of 3.2 ± 2.5% for neritic females and 3.9 ± 2.7% for oceanic fe-

males). Foraging trips lasted 7.96 ± 2.17 d (range 4.26–12.03 d) and average horizontal (surface) distance travelled was ~750 ± 50 km (range 391–1200 km). Oceanic females spent on average 2.5 d more at sea during their foraging and travelled 200 km more than neritic females (both p < 0.009; Fig. 1, Table 1).

Neritic females regularly performed benthic dives (15.2 ± 2.6% of all dives) and occasionally foraged during the day, while oceanic females rarely did so (2.2 ± 0.4% of all dives and 0% when off the shelf) and were nocturnal divers (Table 1, Fig. 1C,D). Neritic females also performed significantly fewer foraging dives than the oceanic ones (p < 0.001), but their dives were almost twice as deep and lasted twice as long (p < 0.05; see details in Table 1). In addition, ~73% of oceanic female dives and ~55% of neritic female dives were shallower than 15 m. Overall, oceanic seals spent significantly less time diving and more time transiting over the full foraging trip time scale compared with neritic females (both p < 0.05; Table 1). In other words, oceanic females travelled ~30 h more than neritic females, for the same number of hours spent diving over the — on average longer — foraging trip (Fig. 1, Table 1). In addition, females that were in better body condition (mass/length) prior to starting a foraging trip made shorter foraging trips compared to females in poorer body condition (slope, p = 0.006, R^2 = 0.58; Fig. 2). In the model (Fig. 2), the slope was similar for both groups (i.e. same relation between body condition and subsequent time at sea) but the intercept was higher for the oceanic group (+1.98 d, p = 0.015), which reflects the longer trip durations of females foraging off-shelf.

Energetic cost of foraging

Rates of energy expenditure per day at sea were not statistically different between foraging strategies whether mass-corrected or not (p > 0.07; Table 1). This translated into oceanic females spending slightly, but not significantly more energy per trip than neritic females overall (p > 0.09), even though they spent more energy transiting to and from feeding grounds (p = 0.06; Table 1).

Prey capture attempts

Oceanic females attempted to capture prey more often per dive (0.76 ± 0.09 PrCA per dive) than neritic females (0.46 ± 0.09 PrCA per dive, p = 0.004). Since
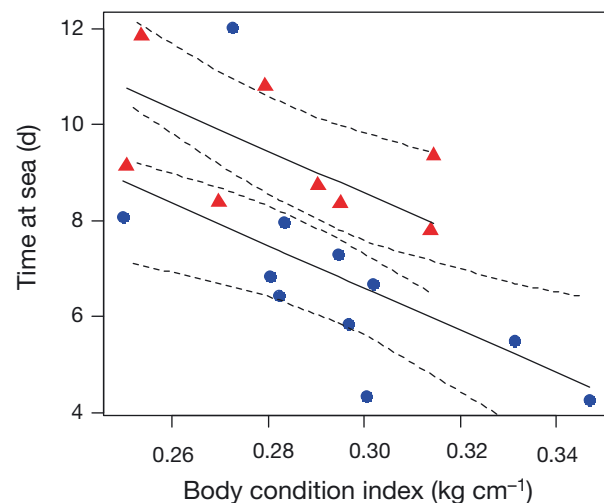

Fig. 2. Time spent at sea by female northern fur seals (n = 20) during a foraging trip from Reef Rookery on St. Paul Island as a function of their own body condition (mass/length) prior to leaving the rookery, for oceanic females foraging off the shelf (red triangles) and neritic females foraging on the shelf (blue circles). Solid lines show the results of the linear regression fitted between the 2 parameters (slopes not different between the group, i.e. all seals will increase their time at sea to the same extent for a given decrease in body condition no matter what foraging group [oceanic or neritic] they belong to) and the dashed lines represent the 95% CI of the predicted values (F = 10.95, p = 0.001, R^2 = 0.58)

oceanic females dove more often at shallower depths and for shorter duration, this translated into a greater rate of capture per min spent diving for oceanic (1.87 ± 0.38 PrCA min⁻¹ spent diving) than neritic females (0.43 ± 0.37 PrCA min⁻¹, p = 0.016). However, the number of attempts to capture prey was similar for both groups (~1.76 PrCA per dive, p = 0.73) when only dives in which at least one PrCA occurred were considered. Overall, neritic females attempted to capture prey nearly 3 times less often than oceanic females per day of foraging (100 ± 26 vs. 275 ± 39 PrCA d⁻¹, respectively; p = 0.024) or per trip (789 ± 163 vs. 2404 ± 338 PrCA per trip, respectively; p = 0.003; Fig. 1, Table 1). Most of the PrCA occurred during nighttime (92.9 ± 2.9% for neritic females and 99.0 ± 0.4% for oceanic females; p = 0.05). Visually, PrCA for oceanic females were localized to small and specific areas (orange dots on Fig. 1A), while neritic females had greater geographic and temporal spread in feeding areas (Fig. 1B).

Diet and prey energy contents

Diet composition of the females depended on their foraging strategy (Table 2) as shown in Jeanniard du Dot et al. (2017a). Neritic females mostly consumed

Table 2. Proportion, calculated as split-sample frequency of occurrence (SSFO), average prey mass, prey energy density (ED) and prey energy content (EC) for different prey groups in diets of female northern fur seals (NFS) breeding on rookeries on St. Paul Island (representative of the 'neritic' group), and on Bogoslof Island (representative of the 'oceanic' group; Jeanniard du Dot et al. 2017a). Prey groups labelled 'Other' were assigned average diet values. SSFO values were obtained from morphological identification of hard-part remains in our study and from collection of samples of Bogoslof Island rookeries in 2009. Mass was calculated from the size of hard-part remains. NA: data not available

Prey group	SSFO (%)	Mass (g)	ED (kJ g^{-1})	EC (kJ)
Neritic NFS				
Gadid	69.75	15.49 ± 0.13	4.09 ± 0.35	67.859 ± 0.28
Cephalopod	12.28	3.74 ± 1.16	4.76 ± 0.11	17.79 ± 0.04
Salmon	6.03	797.78 ± 35.65	5.53 ± 0.30	4425.00 ± 9.80
Hexagrammid	3.64	18.67 ± 2.48	3.90 ± 0.23	75.19 ± 0.31
Mesopelagic	2.42	3.87 ± 0.25	5.67 ± 0.25	21.91 ± 0.05
Forage	1.53	10.69 ± 0.19	4.54 ± 0.66	49.61 ± 1.34
Flatfish	0.28	3.78 ± 0.01	5.14 ± 0.65	19.45 ± 0.08
Other	4.08	NA	4.28 ± 0.72	330.60 ± 11.02
Total diet			4.28 ± 0.72	330.60 ± 11.02
Oceanic NFS				
Gadid	0.86	16.48 ± 0.11	4.12 ± 0.33	67.87 ± 0.24
Cephalopod	42.70	3.74 ± 1.16	4.76 ± 0.11	17.79 ± 0.04
Salmon	3.87	797.78 ± 282.00	5.53 ± 0.30	4425.00 ± 9.80
Hexagrammid	0.60	18.69 ± 2.37	4.02 ± 0.08	75.19 ± 0.29
Mesopelagic	48.97	3.77 ± 0.52	5.85 ± 0.88	21.68 ± 0.01
Forage	1.55	16.47 ± 0.51	5.47 ± 0.57	92.28 ± 3.14
Other	1.46	NA	5.75 ± 1.44	194.94 ± 8.13
Total diet			5.75 ± 1.44	194.94 ± 8.13

gadids (~70%), cephalopods (~12%) and salmon (~6%), while diets of oceanic females consisted mostly of mesopelagic fish (~49%), cephalopods (~43%) and salmon (~4%) (Trites et al. 2015, Jeanniard du Dot et al. 2017a). Analyses of prey size from otoliths, beaks and other bone structures in the scats showed that adult female fur seals preyed mostly on small fish (3 to 20 g). For example, the sizes of ingested pollock corresponded mostly to age-0 and age-1 fish (Fig. 3). The only exception was salmonids, which were ~31 cm long and weighed ~800 g. Energy density of each major fish species ranged from ~3.5 kJ g^{-1} for flatfish, to ~9 kJ g^{-1} for northern lampfish (Table S1 in the Supplement). These sizes translated into energy content of prey groups ranging from 20 to 75 kJ, except for larger salmon that reached ~4500 kJ (Table 2). Mass, energy density and energy content of individual prey species are detailed in Table S1. The total energy density of diet (i.e. ED_{Diet}) for neritic females was 4.28 ± 0.72 kJ g^{-1}. Salmonids and gadids had the highest energy content per prey ingested (i.e. EC_p), with average values of 330.60 ± 11.02 kJ per prey. The energy density of the diet (i.e. ED_{Diet}) for oceanic females was greater than that for neritic fe-

males at 5.75 ± 1.44 kJ g^{-1} (p < 0.03). However, due to the overall smaller size of prey ingested, the energy content per prey ingested (i.e. EC_p) was only 194.94 ± 8.13 kJ (p < 0.001; Table 2).

Foraging efficiency

The calculated energy gains per dive were similar between oceanic and neritic groups (p = 0.82; Table 1), but this translated into a lower rate of energy gain per minute of diving (p = 0.02; Table 1) for neritic females (84.2 ± 16.5 kJ min^{-1}) compared to oceanic females (145.9 ± 24.4 kJ min^{-1}). Similarly, neritic females gained less energy per minute spent at sea over the foraging trip (26.2 ± 6.2 kJ min^{-1}) than oceanic females (35.21 ± 5.4 kJ min^{-1}, p = 0.05).

Based on the foregoing, the resulting average foraging efficiency of all individuals for which simultaneous measures of energy expenditure and prey capture attempts were available (4 animals were missing DLW measurement and/or acceleration data) ranged from 0.42 to 5.40 (Table 3). Foraging efficiency was lower for neritic females (1.51 ± 0.69, 95% CI: 0.86–2.42) than for oceanic females (3.03 ± 0.60, 95% CI: 2.17–4.02; p = 0.05).

Sensitivity analyses showed that the largest contributor to the uncertainty around FE was related to PrCA estimates (Table S2), This was not surprising given that different validation studies have not consistently agreed on their detection rates and false positive rates (Viviant et al. 2010, Volpov et al. 2015). Errors associated with mass of the prey affected FE more than energy density or relative proportion of prey in the diet, at least given the errors we estimated using the bootstrap method. Ultimately, all parameters related to foraging success (PrCA and diet-related parameters) tended to overestimate foraging efficiency, but this was partially compensated by the error associated with the DLW method, which tended to underestimate foraging efficiency. These tendencies were similar between the 2 diet groups (Table S2), as well between this study and the Antarctic fur seal study (Jeanniard du Dot et al. 2017b).
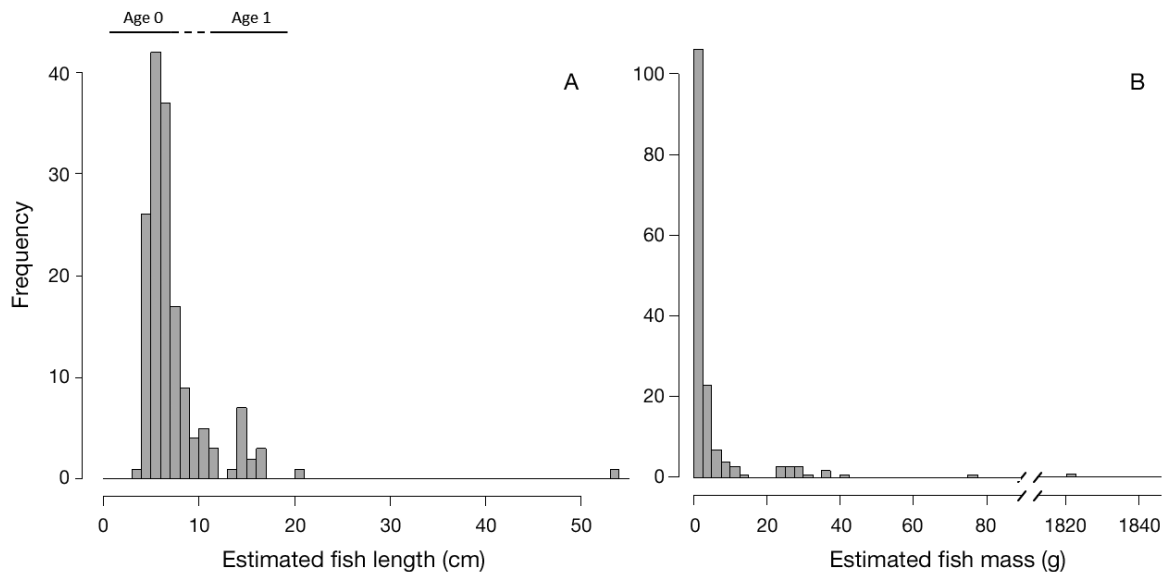
Fig. 3. Frequencies of walleye pollock (A) length and (B) mass consumed by female northern fur seals in the Bering Sea esti-mated from otolith size (n = 173) found in 98 scats collected on Reef Rookery, St. Paul Island, in August 2011 (Jeanniard du Dot et al. 2017a)

DISCUSSION

Food is not evenly distributed in space and time in the ocean, so acquiring it requires animals to make behavioural 'decisions' as to where, when, and for how long to search for food. Such decisions or forag-ing strategies will affect energy expenditure as well as the amount of energy animals manage to extract from their habitat. Since energy is one of the funda-mental currencies of foraging ecology underlying behavioural, physiological, reproductive and devel-opmental traits and capacities, the ratio between energy expenditure and gain from foraging is a piv-otal tool to estimate the growth, survival and repro-ductive fitness of animals.

Foraging efficiency varies with foraging strategy

The 20 female northern fur seals we tracked from Reef Rookery employed 2 foraging strategies that were indicative of different prey bases and distribu-tions (Boyd 1996, Forcada & Staniland 2009). A mi-nority (40%) of the females foraged in oceanic waters over the deep basin. They travelled greater distances, spent more time at sea and less time diving, and made shallower dives of shorter durations (~15 m for ~40 s) compared to the remaining 60% of the females that foraged in neritic waters (~40 m for ~80 s). The oceanic females fed at night on primarily mesopelagic fish such as northern smoothtongue or myctophids,

i.e. on small energy-dense schooling fish that migrate diurnally towards the surface at night (Jeanniard du Dot et al. 2017a). In contrast, neritic females dove benthically to the relatively shallow bottom of the Bering Sea shelf (~150 m). On average, the neritic fe-males spent 15% of their time diving to the bottom (defined as within 95% of bathymetric depth) during night and day, and fed mostly on gadids or other shelf-associated prey. Neritic females tended to ob-tain less energy per gram of prey ingested because they fed on a diet that was 44% lower in energy den-sity than that of oceanic females. However, the neritic females fed on bigger prey, which resulted in ~60% more energy per average prey item ingested com-pared to the average prey consumed by oceanic fe-males. These 2 foraging strategies based on location and diving behaviours have been observed in female fur seals from Russian and Alaskan breeding islands (Gentry 1998), but energetic costs associated with the 2 strategies within a single year have not been previ-ously quantified.

Oceanic females spent less time diving overall, and attempted to capture 4 times more prey per minute spent diving compared to neritic females. Both stra-tegies (oceanic and neritic) resulted in ~1.76 attempts to capture prey per dive in which at least one capture occurred. This value is within the range of prey cap-ture attempts per dive measured in Antarctic fur seals, i.e ~1.88 recorded by Viviant et al. (2016), and ~2 by Jeanniard du Dot et al. (2017b). In addition to performing more (shorter) dives than neritic females,

oceanic females also attempted to capture prey 3 times more often per day and had fewer unsuccessful dives (i.e. dives with no PrCA detected). They also had higher rates of energy gain than their neritic counterparts. A consequence of feeding further from the breeding beaches is that oceanic individuals expended slightly more energy during their foraging trip to and from foraging grounds. However, this increased cost of travelling further was compensated for by the relative energetic advantage that feeding in the deep basin offered them. Consequently, oceanic seals were more energetically efficient at foraging (3.03) compared to neritic seals (1.51).

Despite assumptions and uncertainties in our calculations (detailed in Jeanniard du Dot et al. 2017b), our estimates of foraging efficiency (from 1 to 3) are within in the range of 1 to 4 calculated for other birds and mammals (Hainsworth 1977, Stacey 1986, Parker et al. 1996, Weimerskirch et al. 2003a). However, all of these estimates are much lower than the foraging efficiency of 23 reported for northern fur seals in the 1980s (Costa 1993). This extremely high efficiency was derived using an isotopic method to estimate changes in body water content, and was based on the assumption that the seals were mainly eating squids.

Squids have a high water content and may not have been the dominant prey of the seals studied by Costa (1993), based on the results of more recent studies (Antonelis et al. 1997, Gudmundson et al. 2006, Trites et al. 2015). In addition, the resolution of depth sensors used to record diving behaviours in the 1980s (10 m resolution; Costa 1993) likely resulted in a large proportion of short dives being missed (~40 dives d^{-1} recorded per day in 1980s compared to ~300 dives d^{-1} in this study), and thus in an underestimation of feeding events. As such, energy gain estimated in the in this 1980s study may have been inflated by a factor of 5 or more, thereby overestimating the true foraging efficiency of fur seals.

Conflict between time and energy currencies in maternal investment

Both neritic and oceanic strategies had average foraging efficiencies greater than 1, indicating that the lactating females obtained enough energy to meet their own physiological needs (although more than half the females in the neritic group had FE < 1; Table 3). However, lactation and the cost of pup

Table 3. Measured and corrected energy expenditure (EE), prey capture attempts (PrCA), energy gain, and foraging efficiency during single foraging trips of individual female northern fur seals (NFS) breeding on rookeries on St. Paul Island and foraging in the Bering Sea. Seals foraged either in neritic waters on the shelf (upper rows, n = 9) or in ocean waters in the basin (lower rows, n = 7). Mean values (±SD), as well as average energy content per average prey (EC_p), are also shown for each group. The corrected values of EE and PrCA (calculated as explained in 'Materials and methods; Foraging efficiency') were used with the estimated EC_p to calculate the energy animals gained while foraging at sea and their foraging efficiency (i.e. the ratio of energy gain to energy expended)

Seal ID	Meas. EE (MJ)	Corr. EE (MJ)	Meas. PrCA	Corr. PrCA	EC_p (kJ)	Energy gain (MJ)	Foraging efficiency
Neritic NFS							
1	367.43	360.55	1008	921	330.60 ± 11.02	303.65 ± 52.13	0.89 ± 0.15
5	177.37	174.16	346	319		104.91 ± 17.62	0.64 ± 0.11
6	115.74	113.58	321	295		97.83 ± 16.54	0.89 ± 0.16
8	115.07	113.01	152	140		46.24 ± 7.81	0.42 ± 0.07
12	205.27	201.40	718	660		218.73 ± 37.50	1.13 ± 0.20
13	152.06	149.36	1681	1538		513.94 ± 86.87	3.48 ± 0.64
14	440.71	432.94	635	580		192.28 ± 33.15	0.51 ± 0.08
17	146.65	143.96	549	503		165.83 ± 28.66	1.19 ± 0.22
19	122.60	120.56	1693	1567		517.47 ± 88.11	4.32 ± 0.79
Mean ± SD	204.77 ± 39.40	201.06 ± 38.68	789 ± 189	725 ± 174		240.10 ± 57.77	1.51 ± 0.69
Oceanic NFS							
2	249.57	244.92	2727	2505	194.94 ± 8.49	486.06 ± 85.02	2.02 ± 0.38
3	156.54	153.81	1474	1351		265.36 ± 45.81	1.73 ± 0.32
4	428.07	420.02	3576	3252		641.52 ± 113.62	1.54 ± 0.29
10	73.97	72.67	1996	1839		361.00 ± 62.35	4.96 ± 0.94
11	120.03	117.89	1518	1397		272.40 ± 47.53	2.33 ± 0.44
15	125.03	122.81	3654	3378		656.14 ± 112.83	5.40 ± 1.00
20	136.22	133.86	2381	2194		419.78 ± 74.39	3.23 ± 0.61
Mean ± SD	184.20 ± 45.40	180.85 ± 44.53	2475 ± 339	2274 ± 310		443.18 ± 53.51	3.03 ± 0.60

growth is substantial (Arnould 1997, Goldsworthy et al. 2004). Mothers must therefore have higher foraging efficiencies to ensure their own survival as well as that of their pups (Bouten et al. 1994). As such, oceanic females that gain 3 times more energy than what they spend obtaining it are likely to have more energy overhead to invest in their pups than neritic females that obtain only 1.5 times more energy.

The greater energy surplus available to allocate to reproduction would suggest that oceanic females should produce bigger pups on average and have greater reproductive success than neritic females. However, energy gain is not the factor influencing overall allocation of energy to reproduction (Boggs 1992, Ydenberg et al. 1994). Maximizing the rate of energy delivery to pups (i.e. the frequency of nursing bouts) tends to produce bigger pups (Gentry & Kooyman 1986, Costa 1993). In our study, where 2 different strategies coexisted within a population, neritic females foraging on the shelf made significantly shorter foraging trips compared to oceanic females foraging in the deep basin. As foraging trip duration does not affect time spent ashore nursing pups between foraging trips (Antonelis et al. 1990), the females that adopted the neritic strategy likely fed their pups ~20 to 25% more frequently (considering a consistent 2-d difference in trip duration over the breeding season). This dichotomy in strategies was also observed in California sea lions *Zalophus californianus*, with deep-diving individuals that travelled shorter distances tending to spend more time at the rookery in presence of pup (McHuron et al. 2016). Under this scenario, neritic females likely increase their nursing bout rate, and might thus provide a growth advantage to their pup compared to oceanic females.

On the other hand, additional energy acquired by oceanic females has to be reflected in tissue and/or milk accretion. There are evidences that lactating Australian and Antarctic fur seals that spend more time at sea produce energy richer milk to compensate for the longer time their pups fasted (Arnould & Boyd 1995, Arnould & Hindell 1999). This is suggested by the limited milk lipid content data we collected on our tracked females (Fig. S1 in the Supplement). In addition, there is a general agreement that the composition of fur seal milk reflects what mothers fed on during the days or week prior to the nursing bout (Goebel 2002) and that oceanic females have significantly greater milk lipid content than neritic females (Hayden 2012). Consequently, oceanic females could direct their higher energy overhead to greater energy output per unit of time nursing. However, whether it is physiologically realistic to increase

milk quality to such levels as to provide an identical yield of milk energy as neritic females and compensate for additional days of absence and fewer nursing bouts over the reproductive season (i.e. longer fasting time and greater weight loss of their pup) remains unclear (Gentry & Holt 1986, Trillmich 1990).

The foraging strategies employed by northern fur seals are surprisingly similar to those of Antarctic fur seals feeding in similar habitats in the southern hemisphere (Jeanniard du Dot et al. 2017b). Time budgets (i.e. time at sea, distance travelled, and allocation of time to different activities) of fur seals in the sub-Antarctic were similar to those of neritic northern fur seals. However, energy acquisition (i.e. prey captures, diet and foraging efficiency etc.) of Antarctic fur seals was more similar to oceanic than neritic females. This indicates that northern fur seal individuals compromised on one of the limiting factors, i.e. time allocated to nursing versus energy overhead, whereas Antarctic fur seals from an increasing population did not have to do so (short foraging trips with high foraging efficiencies).

Foraging efficiency of Antarctic fur seal mothers is positively related to offspring growth (Jeanniard du Dot et al. 2017b), which means that, within the context of a single type of foraging strategy and diet composition within a population, females that forage more successfully than others will produce bigger pups at weaning that have greater chances of surviving their first year at sea (Lescroël et al. 2010). The fact that pup mass at weaning is directly linked to pup survival in seals (Doidge & Croxall 1989, Baker & Fowler 1992, Boltnev et al. 1998, McMahon et al. 2000, Hall et al. 2001, Beauplet et al. 2005) implies that female pinnipeds with the highest foraging efficiencies will likely have higher individual fitness. Unfortunately, we could not obtain pup growth data from mother–pup pairs to determine whether this relationship holds for northern fur seals (i.e. whether there was a difference in reproductive success between these 2 foraging and nursing strategies. However, we did collect mass-at-age data for pups (Fig. S2) as a population index of pup growth and global maternal investment to compare to data collected when the population was not declining, 15 or 30 yr previously (Boltnev et al. 1998, Donohue et al. 2000, Angliss & Allen 2009). Analysis of variances on the slopes of these different models showed that pup growth in 2011 was slower than in the 1980s or 1996 (p < 0.001 in all cases, see the section 'Pup growth' in the Supplement). The current reduced growth rates of pups might reflect the fact that this compromise ultimately results in lower provisioning from mothers

(whether by decrease of time feeding pups or energy available for lactation), but this needs confirmation using more robust and standardized data collection.

Foraging strategies within the population

The declining population of northern fur seals has experienced significant environmental changes around their breeding islands (Anderson & Piatt 1999, Napp et al. 2000, Overland & Stabeno 2004). Shifts in ocean climate have been hypothesized to be responsible for decreases in reproduction success of seabirds (Weimerskirch et al. 2003b) and marine mammals (Ferguson et al. 2005, Trites et al. 2007) by changing the availability, type and distribution of prey. In 2011, sea surface temperature on the shelf was colder, compared to the average over the previous 30 yr, but near-bottom temperature was warmer and bottom temperature was close to the average (Lauth & Conner 2014). This data does not indicate a particularly difficult year for walleye pollock body condition (Heintz et al. 2013) or abundance (Lauth & Conner 2014). There was no survey off the shelf in 2011 but water temperatures did not indicate outlier conditions.

The northern fur seal population on St. Paul Island might maintain 2 foraging strategies because they have similar net fitness outcomes, resulting in alternative evolutionary stable strategies (Hines 1987). This is the case for thick-billed murres *Uria lomvia* breeding in the Bering Sea that display similar neritic and oceanic strategies with no apparent difference in chick-feeding rates and fledging success (Paredes et al. 2015). In the case of fur seals, environmental conditions can affect foraging costs between years (Costa 2008), so the difference between both strategies in the population might be averaged in terms of fitness over several generations. It is possible that one strategy does better under particular environmental conditions, and having these 2 strategies in the population provides a buffer to changes in prey distribution/abundance under changing environmental conditions, as suggested for California sea lions (McHuron et al. 2016).

Our study provides a narrow window into the foraging ecology of northern fur seals. Even though there is evidence that the foraging behaviours (Call et al. 2008) and foraging efficiencies (Jeanniard du Dot et al. 2017b) of females are consistent within a year, extending conclusions beyond the timescale of our measurements should be done with caution. Antarctic fur seals in better body condition seem consistently more efficient over the breeding season at exploiting their uniform foraging habitat (Jeanniard du Dot et al. 2017b), but it might not be the case for northern fur seals that can alternate between 2 different habitats and diets. Flexibility or adaptability of females to environmental conditions likely operate at a longer timescale than that of our study. Therefore, our results in the reported short time window only allow educated inferences and conjectures regarding the consequences of foraging efficiencies on the fitness of individuals.

CONCLUSION

Despite the limitations of our study, our results provide interesting clues to the potential energy transfers to pups on St. Paul Island, and highlight the importance of investigating the series of events that link environmental conditions, animal behaviours, energy balance, fitness of individuals, and demographic parameters. It is noteworthy that females from an increasing population (Antarctic fur seals) can on average simultaneously 'optimize' foraging efficiency and frequency of feeding their pups, while females from a declining population (northern fur seals) are faced with a trade-off between these 2 objectives. There is a need to investigate whether one foraging strategy yields a lower reproductive success via lower survival of the young compared to the other, as this could ultimately lead to northern fur seals producing a significant portion of smaller pups at weaning, with lower survival during their first year and thus a lower recruitment into the population (Baker & Fowler 1992). A possible next step could thus be to estimate the survival rate of juveniles over the first few years of their pelagic life, and determine how it relates to the foraging strategies of their mothers during their first 4 mo of life.

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