

UC Irvine

UC Irvine Previously Published Works

Title

Foraging ecology of short-tailed shearwaters near the Pribilof Islands, Bering Sea

Permalink

<https://escholarship.org/uc/item/1pv5j6j5>

Journal

Marine Ecology Progress Series, 141(1-3)

ISSN

0171-8630

Authors

Hunt, GL
Coyle, KO
Hoffman, S
et al.

Publication Date

1996

DOI

10.3354/meps141001

Copyright Information

This work is made available under the terms of a Creative Commons Attribution License, available at <https://creativecommons.org/licenses/by/4.0/>

Peer reviewed

Foraging ecology of short-tailed shearwaters near the Pribilof Islands, Bering Sea

George L. Hunt, Jr.^{1,*}, Kenneth O. Coyle², Susan Hoffman^{1,**}, Mary Beth Decker¹, Elizabeth N. Flint^{1,***}

¹Department of Ecology and Evolutionary Biology, University of California, Irvine, California 92697, USA

²Institute of Marine Sciences, University of Alaska, Fairbanks, Alaska 99775-1080, USA

ABSTRACT: We studied short-tailed shearwaters *Puffinus tenuirostris* foraging near the Pribilof Islands, Alaska, USA, during the summers of 1987, 1988, and 1989. Their foods were almost exclusively the euphausiid *Thysanoessa raschii*, which they obtained both from near-surface swarms and from epibenthic layers. Near-surface mating swarms of euphausiids occurred in areas of elevated phytoplankton standing stocks near inshore tidal fronts. Many of these euphausiids had attached spermatophores. Shearwaters also obtained euphausiids over shallow reefs and inshore of the fronts where euphausiids were trapped in water shallower than 40 m by irregularities in bottom topography ('bathymetric traps'). We hypothesize that the largely inshore distribution of shearwaters in the southeastern Bering Sea described by previous workers is the result of attraction of shearwaters to frontal areas where euphausiids may forage on phytoplankton stocks throughout the summer. These areas, when shallower than 40 m, would also permit shearwaters to access epibenthic aggregations of euphausiids during daylight, when euphausiids not engaged in mating swarms usually migrate to depth.

KEY WORDS: Short-tailed shearwater · *Puffinus tenuirostris* · Euphausiids · *Thysanoessa raschii* · Seabird foraging · Fronts · Bering Sea

INTRODUCTION

Short-tailed shearwaters *Puffinus tenuirostris* migrate annually from their breeding colonies in south-eastern Australia to spend the austral winter in the North Pacific (Shuntov 1964, Tanaka & Kajihara 1979, Guzman 1981, Gould et al. 1982). Millions of these birds forage in the Bering Sea between April and October (Schneider & Shuntov 1993), where they eat predominantly euphausiids, in particular, *Thysanoessa raschii* and *T. inermis* (Ogi 1973, Ogi et al. 1980, Hunt et al. 1981a, Schneider et al. 1986, Troy & Johnson 1989, Troy & Bradstreet 1991). In the Bristol Bay region of the southeastern Bering Sea, Ogi et al. (1980) esti-

mated that shearwaters consume at least 30000 t of euphausiids annually, a consumption roughly equivalent to the consumption of euphausiids by sockeye salmon *Oncorhynchus nerka* during the month that they reside in Bristol Bay before ascending rivers to spawn (Nishiyama 1974). To forage efficiently, shearwaters would profit from mechanisms that permit euphausiids to be predictably harvested. Discovery and understanding of these mechanisms are important because, as shearwaters are the greatest component of avian biomass in the eastern Bering Sea, their foraging distribution and high prey consumption impact the ecosystem (Schneider et al. 1986). Their prey consumption will have its greatest impact if it is highly concentrated in space or time and conflicts with the needs of other high trophic level organisms.

Key to understanding the foraging ecology of short-tailed shearwaters in the southeastern Bering Sea is identification of the circumstances in which they are able to capture euphausiids. Short-tailed shearwaters forage by pursuit diving from the surface, plunging

*E-mail: glhunt@uci.edu

Present addresses:

**Department of Biology, California State University, Long Beach, California 90840, USA

***Hawaiian Islands National Wildlife Refuge, US Fish and Wildlife Service, PO Box 50167, Honolulu, Hawaii 96850, USA

from above the surface, seizing prey at the surface, and hydroplaning, a foot-propelled movement at the surface in which the head and back may be beneath the surface while the wings are held out of the water (Ogi et al. 1980, Morgan 1982). In the eastern Bering Sea, short-tailed shearwaters frequently forage in flocks of tens of thousands or more, with major concentrations of these flocks occurring in an arc from the eastern passes of the Aleutian Islands along the coast into Bristol Bay and then northwestward over the nearshore waters to Nunivak Island (Guzman 1981, Hunt et al. 1981c, Gould et al. 1982, Troy & Johnson 1989). Schneider & Shuntov (1993) reviewed data showing that these large shearwater aggregations are associated with the 50 m isobath, near the inner front (Schumacher et al. 1979, Kinder & Schumacher 1981), throughout the eastern and western Bering Sea, as well as in the shallow passes of the Aleutian and Commander Islands.

A tidal front, similar to the inner front of the southeastern Bering Sea (Schumacher et al. 1979, Kinder & Schumacher 1981), occurs in the vicinity of the 50 m isobath around St. Paul and St. George Islands in the Pribilof Islands of Alaska, USA (Kinder et al. 1983). At St. Paul Island, this front separates well-mixed inshore waters from strongly stratified middle domain water; at St. George Island, the front demarks the transition between a well-mixed nearshore regime and the waters of the outer domain (Coyle & Cooney 1993).

Vertical mixing at fronts is likely to support local phytoplankton blooms over a longer period than would stratified waters of the shelf, which become nutrient depleted after the spring bloom (Sambrotto et al. 1986, Whitley et al. 1986; see also Pingree et al. 1975, Simpson et al. 1979, and Le Fèvre 1986 for a review). For example, in August 1987, high phytoplankton biomass was present above the thermocline just seaward of the fronts surrounding the Pribilof Islands, as were large concentrations of euphausiids (*Thysanoessa raschii* near St. Paul Island and *T. inermis* on the south side of St. George Island; Coyle & Cooney 1993). Elsewhere, frontal regions with enhanced phytoplankton production also support elevated and prolonged production of zooplankton (Kjørboe et al. 1988, Kjørboe 1993, Nielsen et al. 1993), and planktivorous predators such as fish larvae (e.g. Kjørboe et al. 1988) and seabirds (see Hunt & Schneider 1987 and Hunt 1991 for reviews) frequently aggregate in frontal regions to forage.

In this paper we investigate the roles of tidal fronts and irregularities in the bottom that can block the downward migration of euphausiids along a sloping bottom as features of the marine environment important for successful foraging by shearwaters in the Bering Sea. In the 1970s, few short-tailed shearwaters

were seen near the Pribilof Islands, and no feeding flocks were seen (Hunt et al. 1981b). In contrast, during surveys around the islands in 1987 and 1988 (Coyle & Cooney 1993, Hunt unpubl.), we saw high numbers of short-tailed shearwaters, including flocks of feeding birds. In 1989, we investigated the foraging ecology of these shearwaters by analyzing the distribution and behaviors of shearwaters in relation to prey biomass and bottom topography in the vicinity of St. Paul Island.

METHODS

The distribution and abundance of short-tailed shearwaters, acoustically determined biomass and phytoplankton were assessed in the vicinity of St. Paul and St. George Islands in 1987, between 28 July and 16 August, and in 1988, between 28 July and 20 August. Surveys extended approximately 40 nautical miles (n miles) (74 km), weather permitting, from these islands (Figs. 1 & 2; see also Coyle & Cooney 1993). In 1989, between 6 and 14 August, we focused our surveys in the region north of St. Paul where we encountered large numbers of foraging shearwaters in the vicinity of a frontal system (Coyle & Cooney 1993), and to the east of St. Paul where large flocks of foraging shearwaters were also encountered.

Counts of shearwaters and other seabirds were made from the bridge of the RV 'Alpha Helix' (eye height 7.7 m above the sea surface) while the ship was underway. Vessel speeds varied from about 15 km h⁻¹ when we were conducting echosounder surveys to about 19 km h⁻¹ when we were not deploying the transducer. Birds were counted continuously in a 300 m arc from directly ahead of the vessel to 90° off the side with the best visibility (least glare). The continuous count of flying birds (as opposed to a snap-shot method, see Tasker et al. 1984) overestimates the density of flying birds on the transect line (van Franeker 1994), but provides an estimate of avian movements in an area (Hunt et al. 1994). Data on flying birds are depicted as birds counted per 9.3 km of survey, the distance between oceanographic stations which required 0.625 h to cover at 15 km h⁻¹; counts of birds on the water or feeding are presented as numbers km⁻². Areas surveyed were determined by multiplying the distance traveled, as determined by satellite navigation, by the width of the transect surveyed.

Shearwater behaviors were recorded as flying (with flight direction), sitting on the water, and feeding. We defined feeding birds as those which were hydroplaning, plunge diving at shallow angles into the water, or making pursuit dives from the surface. For the purpose of this analysis, we assumed that birds sitting on the

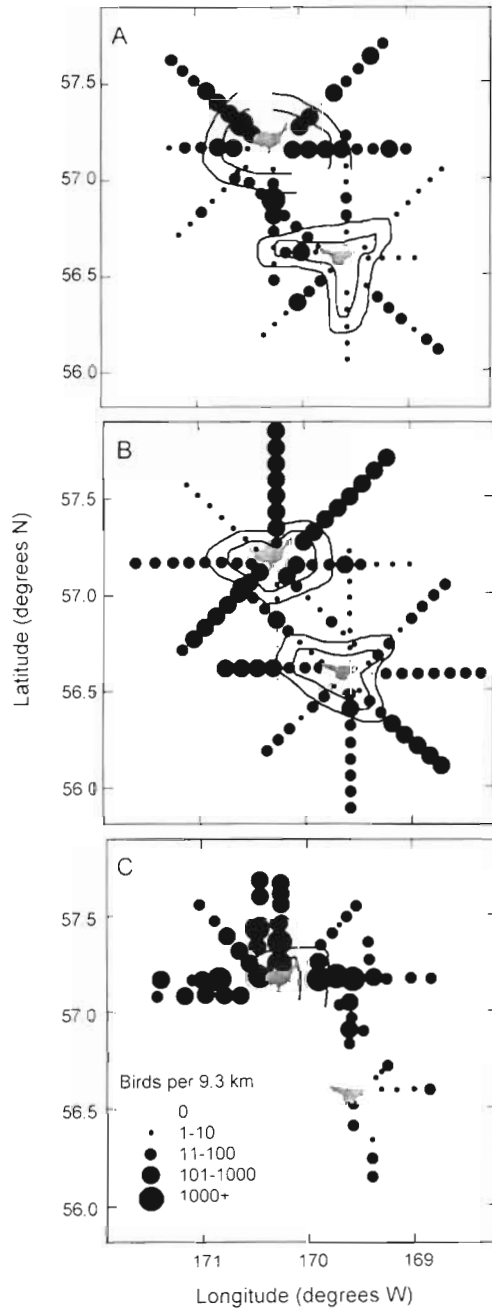


Fig. 1 Distribution and rate of encounter with short-tailed shearwaters in flight (birds 9.3 km^{-1} of survey track): (A) 28 July to 16 August 1987; (B) 28 July to 20 August 1988; and (C) 6 to 14 August 1989. Upper island is St. Paul, lower island is St. George (Alaska, USA); solid lines indicate approximate locations of the tidal fronts around each island, as determined by CTD casts

water had previously fed in the vicinity of where they were resting. We determined the shearwaters' prey by shooting 3 to 5 birds per flock from flocks of birds that were foraging. We limited our collections to foraging birds so that we could be certain that the birds had ob-

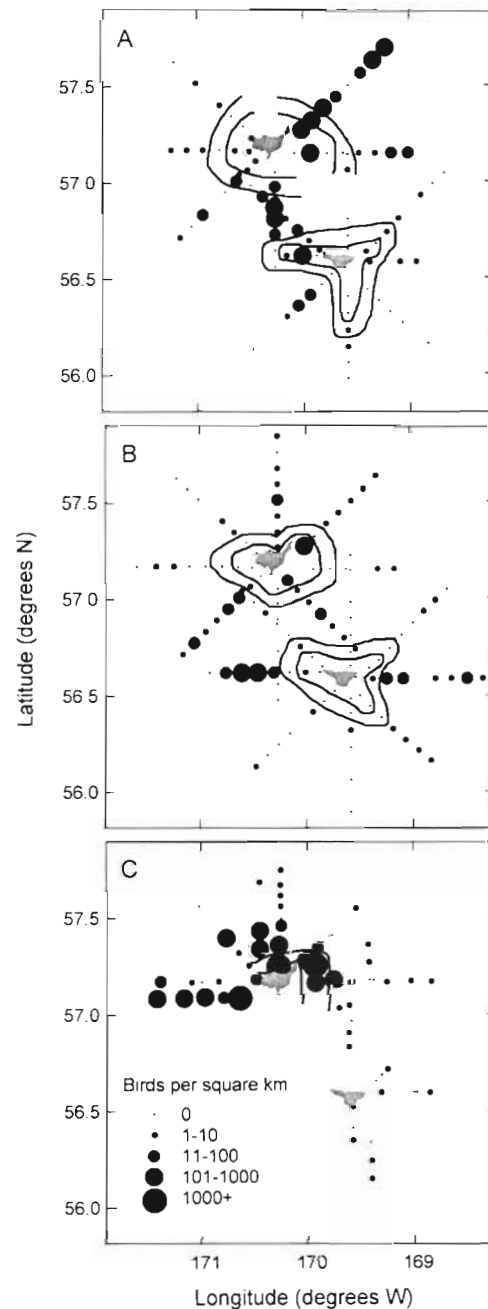


Fig. 2 Distribution and density (birds km^{-2}) of short-tailed shearwaters that were either feeding or resting on the water: (A) 28 July to 16 August 1987; (B) 28 July to 20 August 1988; and (C) 6 to 14 August 1989. Upper island is St. Paul, lower island is St. George; solid lines indicate the approximate locations of the tidal fronts around each island, as determined by CTD casts

tained their prey near where we collected them. We restricted our collections to 5 birds per flock because initial inspection showed that all birds collected had taken similar prey and thus larger samples would not have significantly increased our knowledge of prey use.

Table 2. Distribution of short-tailed shearwater numbers with respect to an oceanographic frontal system north of St. Paul Island in 1989. Proportion of effort or counts in parentheses

	Location					Total
	Inshore	2 n miles inshore of front	4 n miles frontal zone	2 n miles offshore of front	Offshore	
Minutes of survey effort	52 (0.095)	72 (0.132)	152 (0.279)	72 (0.132)	196 (0.360)	544 (0.998)
Shearwaters observed flying	413 (0.059)	1989 (0.283)	2661 (0.380)	564 (0.080)	1388 (0.198)	7015 (1.000)
Shearwaters observed on water or feeding	424 (0.011)	7236 (0.187)	30431 (0.785)	535 (0.014)	131 (0.003)	38757 (1.000)

Table 3. Comparison of expected and observed distributions (proportions) of short-tailed shearwaters, by behavior, with respect to an oceanographic front north of St. Paul Island in 1989. Expected proportion based on time spent surveying zone. Confidence intervals on the proportion of shearwaters observed were constructed for a per family Type I error rate of $\alpha = 0.05$ (Family = flying, feeding or on water); p_i = true proportion of birds in that behavior category in a particular zone

	Location				
	Inshore	2 n miles inshore of front	4 n miles in frontal region	2 n miles offshore of front	Offshore
Expected proportion of birds	0.095	0.132	0.279	0.132	0.360
Confidence interval around true proportion of flying birds	$0.052 \leq p_i \leq 0.066$	$0.270 \leq p_i \leq 0.297$	$0.364 \leq p_i \leq 0.394$	$0.077 \leq p_i \leq 0.081$	$0.183 \leq p_i \leq 0.213$
Confidence interval around true proportion of birds on water or feeding	$0.009 \leq p_i \leq 0.0123$	$0.182 \leq p_i \leq 0.192$	$0.780 \leq p_i \leq 0.791$	$0.0123 \leq p_i \leq 0.0153$	$0.003 \leq p_i \leq 0.004$

shelf-dwelling *Thysanoessa raschii* are vertical migrators, descending to depth during the day (Ponomareva 1963). During the summer months, members of the genus *Thysanoessa* may form daytime near-surface and surface swarms for mating (Smith & Adams 1988, Hanamura et al. 1989). Breeding by *T. raschii* appears to occur in the presence of elevated phytoplankton densities (Paul et al. 1990). It is of interest to examine the predictability of where shearwaters are likely to encounter euphausiids vulnerable to predation. We suggest that 3 mechanisms may support the vast majority of short-tailed shearwater foraging in the southeastern Bering Sea: foraging at euphausiid mating swarms in frontal regions, foraging at depth on euphausiids trapped on the bottom inshore of tidal fronts, and foraging where tidal currents force euphausiids to remain in shallow water, such as may occur in the Aleutian passes.

Fluorescence measurements in the vicinity of the tidal fronts surrounding the Pribilof Islands indicate the presence of elevated levels of phytoplankton (Schneider et al. 1990, Coyle & Cooney 1993) relative

to the stratified waters of the southeastern Bering Sea, where, beginning in June, nutrient limitation leads to low phytoplankton production and standing stocks (Sambrotto et al. 1986, Whitledge et al. 1986). In summer, tidal fronts, such as the front on the north side of St. Paul Island or the inner front of the southeastern Bering Sea, may have elevated levels of phytoplankton in comparison to stratified offshore waters as a result of enhanced vertical flux of nutrients and *in situ* production (Pingree et al. 1975, Simpson et al. 1979, see Le Fèvre 1986, Kiørboe 1993 for reviews). In other frontal systems, enhanced zooplankton production has been associated with enhanced phytoplankton production in the frontal region (Richardson 1985, Kiørboe et al. 1988, Kiørboe 1993).

Summertime near-surface foraging of shearwaters may be associated with euphausiid mating swarms. Spermatophores occurred on many of the euphausiids found in the stomachs of the shearwaters, some of which were foraging at or just beneath the water's surface. In these cases, we conclude that the shearwaters were obtaining euphausiids that were or had recently

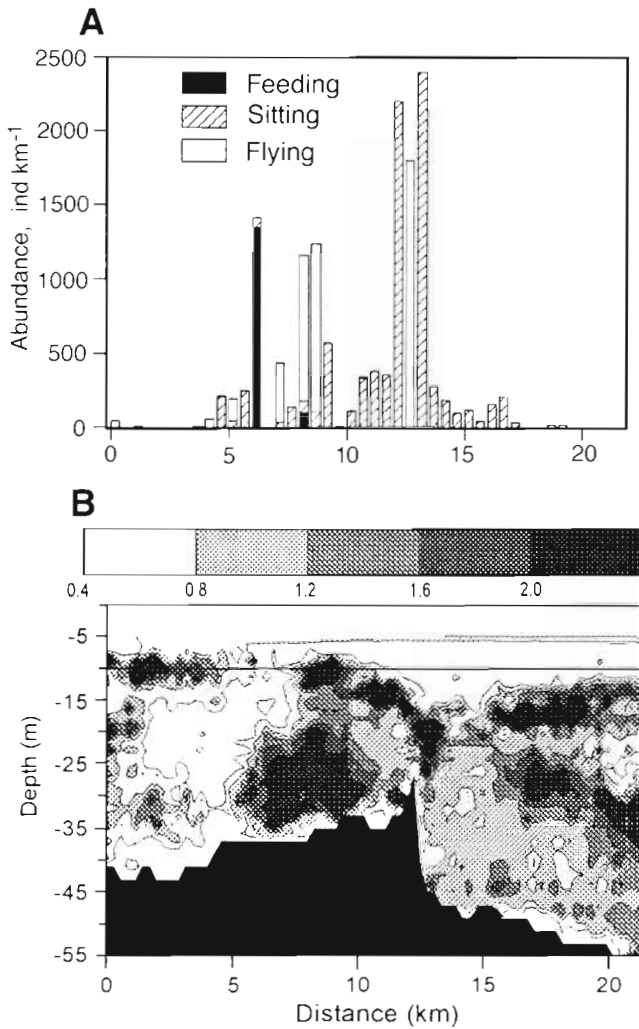


Fig. 5. Observations east of St. Paul Island, 14 August 1989. (A) distribution of short-tailed shearwaters in birds km^{-1} between 1252 and 1412 h ADT; and (B) acoustically estimated biomass in g m^{-3} over a shallow underwater reef

been in mating swarms. We hypothesize that these mating swarms would be most frequently encountered near fronts. We hypothesize further that patches of

near-surface phytoplankton associated with the tidal fronts in the southeastern Bering Sea provide the food necessary to sustain mating swarms of euphausiids long after near-surface phytoplankton concentrations away from the front have decreased. These concentrations of surface swarms of euphausiids could result in the aggregation of foraging shearwaters in the vicinity of fronts. The predominately coastal distribution of short-tailed shearwater flocks in Bristol Bay and the southeastern Bering Sea (Guzman 1981, Hunt et al. 1981c, Gould et al. 1982, Troy & Johnson 1989) fits with this prediction, although we lack the data necessary to demonstrate the validity of the proposed mechanism for determining this distribution.

Shearwater foraging at the Pribilof frontal systems is of particular interest because it is the first documented case of marine birds foraging at a front when the proposed mechanism for enhanced prey availability was the result of prey attraction to elevated phytoplankton concentrations at the front. In other cases of birds foraging at fronts, either the mechanism was not identified, or the zooplankton had been aggregated by physical forcing associated with frontal processes (Hunt 1991, Franks 1992).

We also found short-tailed shearwaters foraging over epibenthic patches of acoustically located biomass which net tows revealed to contain euphausiids (Coyle & Cooney 1993). These patches were at depths of between 30 and 45 m. Epibenthic patches of euphausiids at depths greater than 40 m were rarely attended by foraging or sitting shearwaters. We observed what we interpreted as the bubble tracks of diving shearwaters down to depths of about 30 m, considerably deeper than the 20 m maximum dive depth estimated by Morgan (1982) for short-tailed shearwaters in the Southern Hemisphere. Thus, we would expect that short-tailed shearwaters could capture euphausiids between the surface and 30 or 40 m depth.

We hypothesize that depressions in the tops of reefs and ridges across the slope of the escarpment surrounding the islands may trap euphausiids as they

Table 4. Presence of aggregations of short-tailed shearwaters in relation to prey depth or depth of bottom, and whether prey was detected under the flock, north and east of St. Paul Island in 1989

	Depth of top of prey concentration or of the seafloor, when no prey detected (m)			
	20–29	30–39	40–49	50–59
Feeding shearwaters	1 flock, no prey 2 flocks with prey	1 flock, no prey 3 flocks with prey	No flocks	No flocks
Resting shearwaters	1 flock with prey	3 flocks, no prey 3 flocks with prey	1 flock with prey	1 flock, no prey
Prey concentration detected, no shearwater concentration present	No prey concentrations without shearwaters	No prey concentrations without shearwaters	4 prey concentrations without shearwaters	4 prey concentrations without shearwaters

attempt to migrate downward in the morning. These depressions or 'bathymetric traps' can prevent downward-migrating euphausiids from descending farther because to do so, the euphausiids would have to reverse their downward migration and move toward the surface to clear the edge of the depression or the ridge. Where euphausiids are trapped against the bottom at depths of 40 m or less, they are vulnerable to predation by shearwaters. Euphausiids are unlikely to be trapped by bottoms with extremely steep slopes, such as those on the south side of St. George Island, because they will be able to move deeper while 'in contact' with the bottom. Where the bottom is flat, as it is on the reefs east of St. Paul Island, or slopes gently with ridges and gullies normal to the slope, as it does on the north side of St. Paul Island, there is a greater likelihood of the euphausiids being trapped at a shallow depth.

In the present study, concentrations of foraging or sitting short-tailed shearwaters north of St. Paul Island were at or within 2 n miles inshore of the tidal front. Because most bathymetrically trapped euphausiids attended by short-tailed shearwaters will be at depths of less than 40 m, these aggregations when near the coast are likely to be close to, but inshore of, the tidal fronts which occur parallel to and between the 50 and 70 m isobaths in the southeastern Bering Sea (Schumacher et al. 1979, Kinder et al. 1983, Coyle & Cooney 1993). If enhanced levels of phytoplankton are present in the vicinity of these inshore fronts as they were around the Pribilof Islands (Coyle & Cooney 1993), these frontal areas may be particularly attractive to foraging euphausiids. Thus, although the fronts are not necessary for the entrapment of vertically migrating euphausiids, the presence of a front and its associated concentrations of phytoplankton may lead to more elevated numbers of euphausiids in shallow water than might have occurred in the absence of the fronts.

Coyle et al. (1992) described a third mechanism whereby euphausiids can be forced to remain in shallow water where they are vulnerable to avian predators. At a reef east of St. George Island, murre (*Uria* spp.) foraged on euphausiids that were prevented from descending to depth by tidal currents that impinged upon the reef and counteracted the downward swimming of the euphausiids. A similar combination of a tidal pump and shallow bathymetry may be responsible for the large aggregations of foraging shearwaters in the passes of the Aleutian Islands (Troy & Bradstreet 1991, Schneider & Shuntov 1993, Hunt unpubl. data). Elsewhere, Vermeer et al. (1987) and Brown & Gaskin (1988) described situations in which extremely strong tidal currents interacting with bathymetry force euphausiids to the surface where they are fed upon by surface-foraging birds.

Acknowledgements. We thank the graduate students and volunteers from the University of California, Irvine, The University of California Research Expeditions Program, and the University of Alaska, Fairbanks, and the Captain and crew of the 'Alpha Helix' for their assistance during the cruises. R. T. Cooney aided in field collections, planning and data interpretation. R. Russell aided with the statistical analysis. We thank W. Montevecchi and 2 anonymous reviewers for helpful comments on an earlier version of this paper. This research was supported by National Science Foundation, Division of Polar Programs grant DPP85-09843, and by the University of California Research Expeditions Program.

LITERATURE CITED

- Brown RGB, Gaskin DE (1988) The pelagic ecology of the grey and red-necked phalaropes *Phalaropus fulicarius* and *P. lobatus* in the Bay of Fundy, eastern Canada. *Ibis* 130:234–250
- Coyle KO, Cooney RT (1993) Water column sound scattering and hydrography around the Pribilof Islands, Bering Sea. *Cont Shelf Res* 13:803–827
- Coyle KO, Hunt GL Jr, Decker MB, Weingartner TJ (1992) Murre foraging, epibenthic sound scattering and tidal advection over a shoal near St. George Island, Bering Sea. *Mar Ecol Prog Ser* 83:1–14
- Franks PJS (1992) Sink or swim: accumulation of biomass at fronts. *Mar Ecol Prog Ser* 82:1–12
- Gould PJ, Forsell DJ, Lensink CJ (1982) Pelagic distribution and abundance of seabirds in the Gulf of Alaska and the eastern Bering Sea. US Fish and Wildlife Service FWS/OBS-82/48, Washington, DC
- Green CH, Wiebe PH, Burczynski J (1989) Analyzing zooplankton size distribution using high-frequency sound. *Limnol Oceanogr* 34:163–178
- Guzman J (1981) The wintering of sooty and short-tailed shearwaters (genus *Puffinus*) in the North Pacific. PhD thesis, University of Calgary
- Hanamura Y, Kotori M, Hamaoka S (1989) Daytime surface swarms of the euphausiid *Thysanoessa inermis* off the west coast of Hokkaido, northern Japan. *Mar Biol* 102:369–376
- Haney JC, Solow AR (1992) Testing for resource use and selection by marine birds. *J Field Ornithol* 63:43–52
- Hunt GL Jr (1991) Occurrence of polar seabirds at sea in relation to prey concentrations and oceanographic factors. In: Sakshaug E, Hopkins CCE, Øritsland NA (eds) Proceedings of the Pro Mare Symposium on Polar Marine Ecology, Trondheim, 12–16 May 1990. *Polar Res* 10(2):553–559
- Hunt GL Jr, Burgeson B, Sanger GA (1981a) Feeding ecology of seabirds of the eastern Bering Sea. In: Hood DW, Calder JA (eds) The eastern Bering Sea: oceanography and resources. NOAA/BLM, Washington, DC, p 629–647
- Hunt GL Jr, Croxall JP, Trathan PN (1994) Marine ornithology in the southern Drake Passage and Bransfield Strait during the BIOMASS programme. In: El-Sayed SZ (ed) Southern Ocean ecology: the BIOMASS perspective. Cambridge University Press, Cambridge, p 231–245
- Hunt GL Jr, Eppley Z, Burgeson B, Squibb R (1981b) Reproductive ecology, foods, and foraging areas of seabirds nesting on the Pribilof Islands, 1975–1979. Environmental assessment of the Alaskan continental shelf; final reports of principal investigators, Vol 12. NOAA/OMPA, Washington, DC, p 1–257
- Hunt GL Jr, Gould PJ, Forsell DJ, Peterson H Jr (1981c) Pelagic distribution of marine birds in the eastern Bering

- Sea. In: Hood DW, Calder JA (eds) The eastern Bering Sea shelf: oceanography and resources. NOAA/BLM, Washington, DC, p 689-718
- Hunt GL Jr, Schneider DC (1987) Scale-dependent processes in the physical and biological environment of marine birds. In: Croxall JP (ed) Seabirds: feeding biology and role in marine ecosystems. Cambridge University Press, Cambridge, p 7-41
- Kinder TH, Hunt GL Jr, Schneider D, Schumacher JD (1983) Correlation between seabirds and oceanic fronts around the Pribilof Islands, Alaska. *Estuar Coast Shelf Sci* 16: 309-319
- Kinder TH, Schumacher JD (1981) Hydrographic structure over the continental shelf of the southeastern Bering Sea. In: Hood DW, Calder JA (eds) The eastern Bering Sea shelf: oceanography and resources. NOAA/BLM, Washington, DC, p 31-52
- Kjørboe T (1993) Turbulence, phytoplankton cell size, and the structure of pelagic food webs. *Adv Mar Biol* 29:1-72
- Kjørboe T, Munk P, Richardson K, Christiansen V, Paulsen H (1988) Plankton dynamics and herring larval growth, drift and survival in a frontal area. *Mar Ecol Prog Ser* 44: 205-219
- Le Fèvre J (1986) Aspects of the biology of frontal systems. *Adv Mar Biol* 23:163-299
- Morgan WH (1982) Feeding methods of the short-tailed shearwater *Puffinus tenuirostris*. *Emu* 82:226-227
- Neu CW, Byers CR, Peek JM (1974) A technique for analysis of utilization-availability data. *J Wildl Manage* 38:541-545
- Nielsen TG, Løkkegaard B, Richardson K, Pederson FB, Hansen L (1993) Structure of plankton communities in the Dogger Bank area (North Sea) during a stratified situation. *Mar Ecol Prog Ser* 95:115-131
- Nishiyama T (1974) Energy requirement of Bristol Bay sockeye salmon in the central Bering Sea and Bristol Bay. In: Hood DW, Kelly EJ (eds) Oceanography of the Bering Sea with emphasis on renewable resources. *Inst Marine Sci Occas Publ 2*, University of Alaska, Fairbanks, p 321-343
- Ogi H (1973) Preliminary examination of stomach contents of murre (*Uria* spp.) from the eastern Bering Sea and Bristol Bay, June-August, 1970 and 1971. *Jpn J Ecol* 23:201-209
- Ogi H, Kubodera TK, Nakamura K (1980) The pelagic feeding ecology of the short-tailed shearwater *Puffinus tenuirostris* in the subarctic Pacific region. *J Yamashina Inst Ornithol* 12:157-182
- Paul AJ, Coyle KO, Ziemann DA (1990) Timing of spawning of *Thysanoessa raschii* (Euphausiacea) and occurrence of their feeding stage larvae in an Alaskan bay. *J Crust Biol* 10:69-78
- Pingree R, Pugh P, Holligan PM, Forster G (1975) Summer phytoplankton bloom and red tides in the approaches to the English Channel. *Nature* 258:672-677
- Ponomareva LA (1963) Euphausiids of the North Pacific Ocean: the distribution and ecology of the major species. *Akademiya Nauk, SSSR. Institut Okeanologii, Moscow* [In Russian]
- Richardson K (1985) Plankton distribution and activity in the North Sea/Skagerrak-Kattegat frontal area in April 1984. *Mar Ecol Prog Ser* 26:233-244
- Sambrotto RN, Niebauer HJ, Goering JJ, Iverson RL (1986) Relationships among vertical mixing, nitrate uptake, and phytoplankton growth during the spring bloom in the southeast Bering Sea middle shelf. *Cont Shelf Res* 5:161-198
- Schneider DC (1982) Fronts and seabird aggregations in the southeastern Bering Sea. *Mar Ecol Prog Ser* 10:101-103
- Schneider DC, Harrison NM, Hunt GL Jr (1990) Seabird diet at a front near the Pribilof Islands, Alaska. *Stud Avian Biol* 14:61-66
- Schneider DC, Hunt GL Jr, Harrison NM (1986) Mass and energy transfer to seabirds in the southeastern Bering Sea. *Cont Shelf Res* 5:241-257
- Schneider DC, Shuntov VP (1993) The trophic organization of the marine bird community in the Bering Sea. *Rev Fish Sci* 1:311-335
- Schumacher JD, Kinder TH, Pasinski DJ, Charnell RL (1979) A structural front over the continental shelf of the eastern Bering Sea. *J Phys Oceanogr* 9:79-87
- Shuntov VP (1964) Transequatorial migrations of the thin-billed shearwater *Puffinus tenuirostris* (Temm.). *Zool Zh* 43:590-598 (in Russian)
- Simpson JH, Edlestein DJ, Edwards A, Morris NCG, Tett PB (1979) The Islay Front: physical structure and phytoplankton distribution. *Estuar Coast Mar Sci* 9:713-726
- Smith SE, Adams PB (1988) Daytime surface swarms of *Thysanoessa spinifera* (Euphausiacea) in the Gulf of the Farallones, California. *Bull Mar Sci* 42:76-84
- Springer AM (1992) A review: walleye pollock in the North Pacific — how much difference do they really make? *Fish Oceanogr* 1:80-96
- Tanaka Y, Kajihara T (1979) The distribution of *Fulmaris glacialis* and *Puffinus tenuirostris* in the North Pacific and the Okhotsk Sea during the summer. *J Yamashina Inst Ornithol* 11:79-86
- Tasker ML, Hope Jones P, Dixon T, Blake BF (1984) Counting seabirds from ships: a review of methods employed and a suggestion for a standardized approach. *Auk* 101:567-577
- Troy DM, Bradstreet MSW (1991) Marine bird abundance and habitat use. In: Marine birds and mammals of the Unimak Pass Area: abundance, habitat use and vulnerability. LGL Alaska Research Associates, Inc. Final Report to Mineral Management Services, Anchorage, Contract MMS14-35-001-30564, p 5-1 to 5-70
- Troy DM, Johnson SR (1989) Marine birds, Section 6. In: Truett JC (ed) Environmental characterization and biological utilization of the North Aleutian Shelf nearshore zone. Outer Continental Shelf Environmental Assessment Program, Final Reports of Principal Investigators 60. NOAA, Washington, DC, p 355-453
- van Franeker JA (1994) A comparison of methods for counting seabirds at sea in the Southern Ocean. *J Field Ornithol* 65: 96-108
- Vermeer K, Szabo I, Greisman P (1987) The relationship between plankton-feeding Bonaparte's and mew gulls and tidal upwelling at Active Pass, British Columbia. *J Plankton Res* 9:483-501
- Whitledge TE, Reeburgh WS, Walsh JJ (1986) Seasonal inorganic nitrogen distributions and dynamics in the southeastern Bering Sea. *Cont Shelf Res* 5:109-132

This article was presented by D. C. Schneider (Senior Editorial Advisor), St. John's, Newfoundland, Canada

Manuscript first received: May 26, 1995
Revised version accepted: July 10, 1996