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Climatic effects on breeding seabirds of the northern Japan Sea

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ABSTRACT: Seabirds as marine top predators have been put forth as reliable indicators of ecosystem change. To understand climate–ecosystem change in the northern Japan Sea, we studied the timing of breeding, chick diets, and breeding success of 3 seabird species almost continuously over 26 yr on Teuri Island, 1984 through 2009. Key climate drivers in this region are: (1) westerly winter winds that cool the atmosphere and the ocean and (2) the northward flowing Tsushima Current (TC) that warms the ocean in late spring and summer. Chick diet showed decadal changes in coastal pelagic fish, with a switch from sardine *Sardinops melanostictus* to anchovy *Engraulis japonicus* in the late 1980s corresponding to intensification of the TC. There were no long-term trends in the timing of breeding or breeding success of rhinoceros auklet *Cerorhinca monocerata* and Japanese cormorant *Phalacrocorax filamentosus*, but these variables were affected by interannual variation in spring air and sea temperatures, as well as the timing of TC warm water intrusions within the foraging range (~60 km from the colony) of the birds. Effects include: (1) freezing of the breeding grounds by winter winds limiting access to nesting grounds and (2) availability of anchovy and sandlance *Ammodytes personatus* affecting chick diet, growth, and breeding success. We did not detect relationships between chlorophyll *a* concentrations and the availability of prey species. The timing and success of black-tailed gulls *Larus crassirostris*, which fed mainly on sandlance, however, could not be explained by these factors. Our study indicates potential links between the Pacific Decadal Oscillation index and the flow rate of the TC, and between the Arctic Oscillation index and local wind stress, and suggests that broad-scale atmospheric pressure fields influence local weather, oceanography, and seabirds indirectly through changes in prey availability.

KEY WORDS: Auklet · Breeding performance · Chick diet · Cormorant · Gull · Prey availability · Regime shift · Timing of breeding

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INTRODUCTION

Changes in wind stress (Aebischer et al. 1990), sea-ice extent (Loeb et al. 1997), sea-surface temperature (SST; Bertram et al. 2001, Frederiksen et al. 2006), and upwelling (Thayer & Sydeman 2007, Schroeder et al. 2009) affect the foraging and breeding of seabirds through changes in food web structure. In addition to these 'bottom-up' effects, changes in seasonal and spatial patterns of prey influences seabird diet

and breeding performance (Durant et al. 2005, Grémillet et al. 2008). Adverse local weather (low air temperature, heavy rain or snow, or strong winds) may limit accessibility to nesting grounds (reviewed by Schreiber 2002). Thus, various mechanisms may underlie the responses of seabirds to climate variability and change.

To understand climatic drivers of marine ecosystem change, it is necessary to examine how seabird diet and breeding performance varies with climatic fac-

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tors on multiple spatial and temporal scales. Seabirds may respond to climate fluctuations on immediate to interannual (Gaston et al. 2009, Moe et al. 2009), decadal (regime shift) (Anderson & Piatt 1999, Durant et al. 2004), and long-term (ocean warming) (Jenouvrier et al. 2009) scales. Responses may also vary between regions (Frederiksen et al. 2007), depending on regional geography, marine physics, and ecosystems, and vary between species depending on the species-specific constraints (Furness & Tasker 2000). Thus, long-term information from multiple species in multiple regions is useful for testing mechanisms.

The Japan Sea lies between the Eurasian continent and Japan (Fig. 1). Here, increases in air temperature (www.data.kishou.go.jp/climate/cpdinfo/temp/an_jpn) and SST (Yeh et al. 2010) have been reported and related to global climate change. In addition, the warm Tsushima Current (TC) heats the sea during the late spring and summer, while westerly winds cool the ocean in winter (Hase et al. 1999, Chiba & Saino 2003). These latter factors vary at 2 time scales. (1) A decadal change in the surface air pressure in Siberia has influenced wind stress and SST; the SST was lower between the late 1970s and late 1980s and has been higher since then (Minobe et al. 2004, Yeh et al. 2010). This climatic regime shift induced a change in phytoplankton and zooplankton (Chiba & Saino 2003, Chiba et al. 2005), as well as pelagic fish communities (Tian et al. 2008). (2) Annual changes in SST are related to air pressure anomalies in the northwestern Pacific represented by variation of the

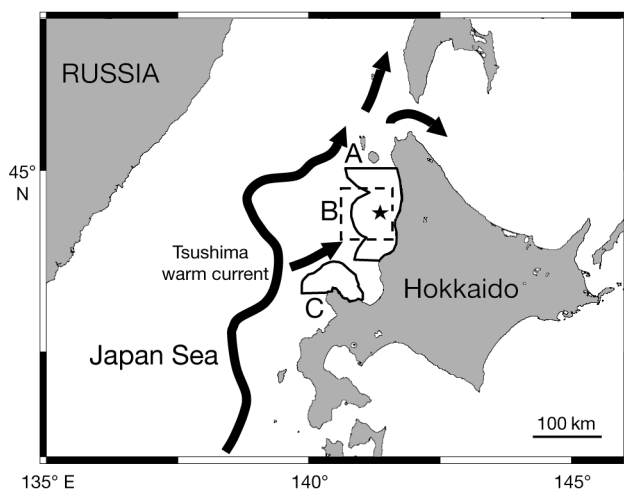


Fig. 1. Study area. ★ = Teuri Island. Around the colony, we measured local sea-surface temperature (SST) in Area A (60 × 150 km), chlorophyll *a* in Area B, and SST for determining the date of warm water arrival (WWA) within foraging range of rhinoceros auklets (Area C)

Aleutian Low, which drives the warm TC northward, from the Tsushima Strait in the southern Japan Sea (Minobe et al. 2004).

To understand the relationships between local and large-scale climate fluctuations and seabird biology in the Japan Sea we studied the chick diets and breeding performance of *Cerorhinca monocerata* (rhinoceros auklets, RHAU), *Phalacrocorax filamentosus* (Japanese cormorants, JCOM), and *Larus crasirostris* (black-tailed gulls, BTGL) on Teuri Island (Fig. 1) over 26 yr, 1984 through 2009. We examined long-term trends and the effects of interannual variation in air temperature, SST, other local climate factors, and prey availability on the timing of breeding, chick diets, and breeding success. We discuss how these local factors are influenced by broad-scale climate change.

MATERIALS AND METHODS

Study area and species

We conducted our study on Teuri Island (44° 25' N, 141° 52' E). Data were collected in 1984, 1985, and 1987, and from 1992 to 2009 (n = 21 yr). Additional data on the breeding of BTGL were collected in 1980. Table 1 summarizes the breeding biology and chick diets of the 3 studied species. The population sizes varied from year to year. RHAU feed on sand-lance *Ammodytes personatus*, juvenile Japan Sea greenling *Pleurogrammus azonus*, and krill *Thysanoessa longipes* and *T. inermis* in spring (Ito et al. 2009) and on sand-lance, sardine *Sardinops melanostictus*, and anchovy *Engraulis japonicus* in summer. JCOM feed on benthic fish in spring (M. Ito unpubl. data) and on epipelagic fish (sand-lance and anchovy), benthic rock fish *Sebastes* spp., flat fish (Pleuronectiformes), and epibenthic greenlings in summer. BTGL feed on krill *T. inermis* and fish in spring (Tomita et al. 2009) and mainly on sand-lance in summer. All 3 species spend winter around the Japan archipelago.

Study plots and breeding biology

RHAU nest in burrows 1 to 2 m deep on cliff shoulder slopes. JCOM nest on cliff ledges and rock stacks, and BTGL on glassy slopes. A single RHAU study plot (ca. 20 × 50 m) containing ~1000 burrows was set-up on a gentle slope in 1984. To increase the number of JCOM samples while minimizing plot size

Table 1. *Cerorhinca monocerata*, *Phalacrocorax filamentosus*, *Larus crassirostris*. Total number of nests, body mass, foraging pattern, and foraging range in rhinoceros auklets (RHAU) Japanese cormorants (JCOM) and black-tailed gulls (BTGL). Range of the annual mean values of the mass proportion of fish prey in chick diet, the timing of breeding (hatching date for RHAU, hatching date of the first chick in a brood for JCOM, clutch initiation date for BTGL), clutch size, and the number of fledglings per active nest (nests with ≥ 1 hatching or egg)

	Rhinoceros auklet (RHAU)	Japanese cormorants (JCOM)	Black-tailed gull (BTGL)
No. of active nests	190,000–290,000 ^a	500–1600 ^a	2500–30,000 ^a
Body mass (kg)	0.5–0.6 ^b	2.3–3.2 ^e	0.5–0.6 ^h
Foraging mode	Wing-propelled diving	Foot-propelled diving	Surface feeding
Diving depth (m)	14.0 (median), 50 (max.) ^b	7.2–15.1 (mean), 26–39 (max.) ^e	–
Foraging range (km)	87 (0.5–164) ^c	10–22 ^f	Around the island ⁱ
Wintering area	Along Japan archipelago ^d	Along Japan archipelago ^g	Along Japan archipelago ^j
Percent chick diet in 1984–2009 (median)^l			
Sardine	0–24 (0)	0–19 (0)	0–53 (0)
Sandlance	2–57 (13)	3–92 (21)	8–86 (41)
Anchovy	1–95 (69)	0–49 (13)	0–65 (20)
Juvenile Japan Sea greenling	1–50 (9)	0 (0)	0 (0)
Greenlings	0 (0)	0–60 (22)	0–34 (0)
Demersal fish	0 (0)	3–69 (16)	0–12 (0)
Egg laying date ^l	1 Apr–24 Apr ^k	8 Apr–27 May ^k	5 May–28 May
Hatch date ^l	16 May–9 Jun	6 May–24 June	2 Jun–25 Jun ^k
Clutch size ^l	1	2.6–4.0	1.4–2.6
No. of fledglings per nest ^l	0.3–0.9 per nest with chick	0.5–2.6 per nest with egg	0.0–1.2 per nest with egg

^aOsa & Watanuki (2002), Watanuki et al. (unpubl. data), ^bKuroki et al. (2003), ^cKato et al. (2003), ^dA. Takahashi (pers. comm.), ^eWatanuki et al. (1996), ^fWatanuki et al. (2004), ^gNelson (2005), ^hChochi et al. (2002), ⁱOlssen & Larsson (2004), Y. Osa pers. comm., ^jWatanuki (1987a), ^kCalculated from incubation period: 46 d (median) for RHAU (Gaston & Jones 1998), 28 d (mean) for JCOM (E. Hayashi pers. comm.), and 28 d (mean) for BTGL (Niizuma et al. 2005), ^lPresent study

and thus disturbance of BTGL, we set 2 to 4 JCOM plots (ca. 20 × 30 m) on cliff faces by map and 2 to 4 BTGL plots (ca. 10 × 10 m) on slopes within stakes. Artificial nest boxes were set in part of the RHAU plot. Breeding success and chick growth did not differ between natural burrows and nest boxes (Kuroki et al. 1998).

For RHAU we walked in the plot and checked the presence of eggs or chicks in 10 to 66 nests annually, including nest boxes, every 5 d. As RHAU are sensitive to disturbance during egg laying and incubation, we monitored them primarily after hatching. Chicks that disappeared from the nest after 40 d were considered to have fledged (Takahashi et al. 2001). We monitored the contents of 26 to 62 JCOM nests annually every 5 d from ca. 50 m away using a 20× to 40× telescope due to difficulties in approaching the nests. Chicks that disappeared after 45 d were considered to have fledged (Kato et al. 2001). For BTGL we walked in study plots and marked each nest with a numbered stake when we found newly laid eggs and banded newly hatched chicks. We monitored 30 to 104 nests annually and checked the presence of banded chicks every 5 d. Chicks that disappeared after 30 d or after attaining 500 g were considered to have fledged (Watanuki 1987b).

As the indices of the timing of breeding in each year, we used the mean laying date of the first egg in BTGL nests, and the mean hatching date of RHAU (Fig. 2a). The mean hatching date of the first JCOM chick in nests was used as timing index for this species, as we did not identify egg laying by JCOM in some years. Development period of embryo, i.e. incubation period, is relatively constant in most homoeothermic bird species so both the timing of egg-laying and the timing of hatching can be used as an index for the timing of breeding. As an index of breeding success for RHAU, we used the number of fledglings produced per chick hatched per pair (Fig. 2b). For JCOM and BTGL, we used the number of fledglings produced per pair that had eggs.

Collection and analysis of seabird diet

RHAU bring back food cross-wise in their bills in the evening and night, and JCOM and BTGL, in the stomach during the day. We caught RHAU arriving with prey after sunset and collected 64 to 226 bill-loads each year. Chicks and adults of BTGL and JCOM sometimes vomited food when they were

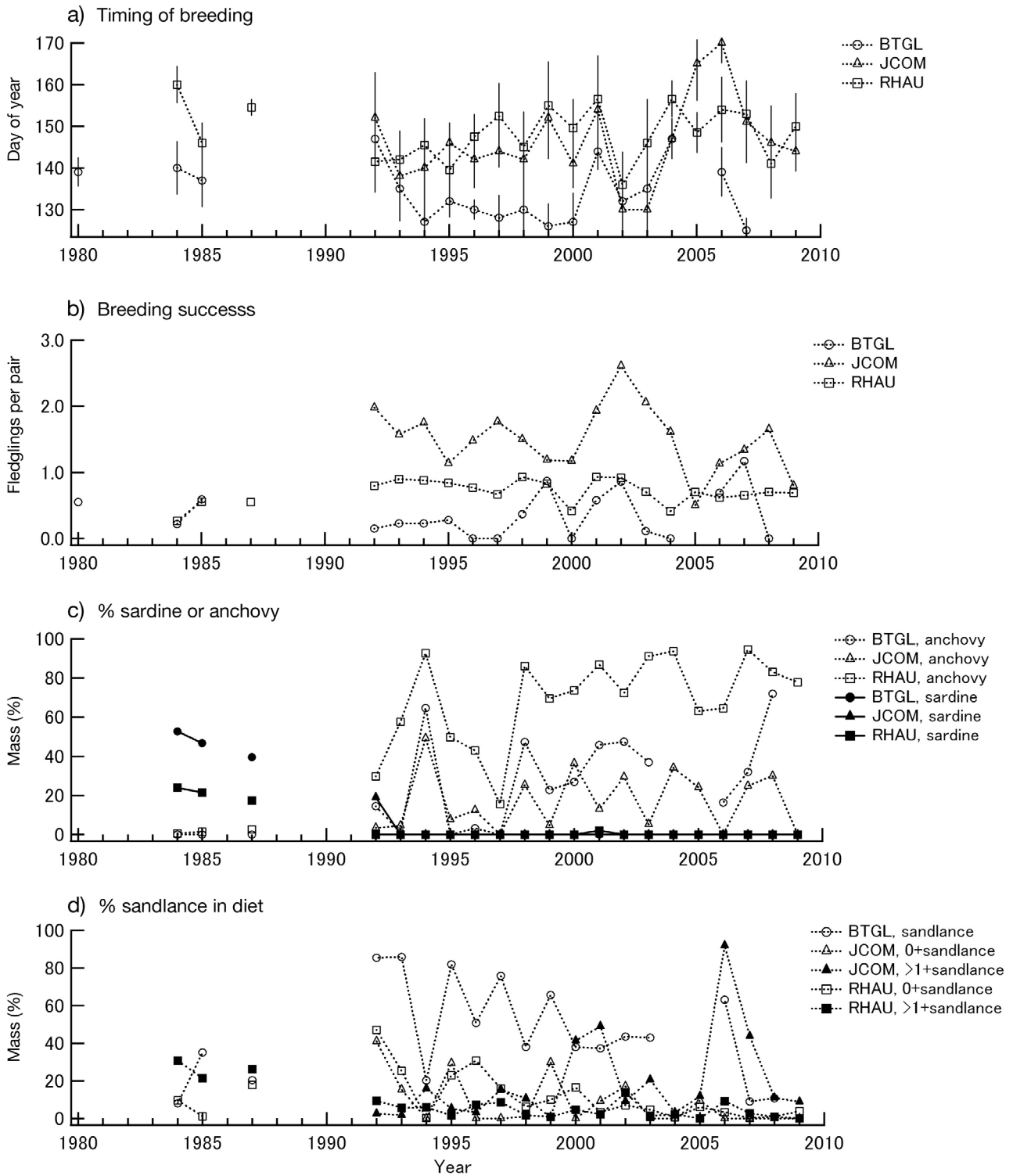


Fig. 2. *Larus crassirostris*, *Cerorhinca monocerata*, *Phalacrocorax filamentosus*. Interannual changes in (a) timing (mean \pm 1 SD) of breeding, (b) breeding success, and (c,d) dietary composition of black-tailed gulls (BTGL), rhinoceros auklets (RHAU), and Japanese cormorants (JCOM). (a) The timing is represented by egg laying for BTGL and hatching for RHAU and JCOM. (b) Breeding success is shown as the mean number of fledglings per pair. (c) Changes in the mass proportion of sardine and anchovy in the diet. (d) Changes in the mass proportion of 0+ sandlance and >1+ sandlance in the diet of RHAU and JCOM, and of all sandlance in the diet of BTGL

caught, and we collected 19 to 115 regurgitations from BTGL and 21 to 37 from JCOM each year.

Each food sample was weighed, and prey items were sorted and weighed separately. Prey items were identified to the lowest possible taxonomic level. We estimated the length of prey fish from otolith size (Ishikawa & Watanuki 2002) or we measured the total length of undigested fish. The sandlance were categorized into cohorts of 0+ yr (<110 mm total length) and >1+ yr (≥ 110 mm) (Takahashi et al. 2001), although it was sometimes difficult in food samples of BTGL, since these were degraded. We also calculated the percentage mass of important prey (sardine, anchovy, 0+ and >1+ sandlance) in the total each year (Fig. 2c,d).

Prey abundance

As indices of the availability of major prey in spring (krill, sandlance) and summer (anchovy, sandlance), we used the reported stock size of anchovy, catch per unit effort (CPUE) for krill, and the total annual catch of 0+ sandlance from fisheries (Fig. 3a). Krill and 0+ sandlance are harvested in March to April and May, respectively, around the island. We used the CPUE for krill (KrillCPUE, annual catch per fishing days as reported by fishermen on Yagishiri Island, ~4 km east of Teuri Island, in 1992 to 2007) and the annual catch of 0+ sandlance around Teuri and Yagishiri Islands (CatSL0, published by the Department of Fisheries and Forestry from 1980 to 2005, and on www.pref.hokkaido.lg.jp from 2006). The Tsushima anchovy stock is distributed from the west coast of Kyushu to the Japan Sea coast, and is seasonally available for seabirds breeding on Teuri Island when its distribution extends to the north. Its stock size (ENGstock) has been estimated annually by the Fisheries Agency since 1991, and its trend has been validated by local acoustic surveys (<http://abchan.job.affrc.go.jp>, in Japanese).

Local climate and broad-scale climate indices

Wind, rain, snow, ice or low air temperatures can cause freezing of the Teuri nesting grounds, metabolic stress, and difficulty in finding food. Changes in ocean temperature (indexed by SST) may influence the distribution and abundance of prey. We used monthly (February to July) average air temperature (Temp.) and monthly rainfall (Rain) recorded on Yagishiri Island, and the total snowfall in March (Snow),

when seabirds arrive at the colony, recorded at Haboro weather station (~30 km east of Teuri Island) (www.data.jma.go.jp, in Japanese) to measure local weather conditions. We used monthly mean SST in an area of 60 × 150 km around the island (Area A in Fig. 1, Fig. 3b) to index ocean temperature. Local SST in this sector was reported by Hakodate Kaiyou Kisyodai from 1985 (www.jma-net.go.jp/kahodate/menu/sea.html, in Japanese); it was based on MGDSST (merged satellite and *in situ* data global daily SST) collected by the AVHRR (advanced very high resolution radiometer) sensor on NOAA (US National Oceanic and Atmospheric Administration) satellites and the Japanese multi-functional transport satellite (MTSAT).

Anchovy is harvested from waters with a SST of 12 to 15°C around Hokkaido (Mihara 1998), and seasonally expands its distribution northward during summer. RHAU switches prey from sandlance and juvenile greenling to anchovy when warm water (SST = 13°C) arrives at the southern edge of its maximum foraging range (164 km; Table 1, Area C in Fig. 1) in late April to late June (Watanuki et al. 2009). We used the date of warm water arrival (WWA, www.jma-net.go.jp/kahodate/menu/sea.html, in Japanese) in the region as an index of the seasonal availability of anchovy to breeding seabirds.

We used an index of the flow rate of the TC (area of the water at 100 m depth with temperature >10°C) (www.data.kishou.go.jp/kaiyou/shindan/e_2/maizuru_tsushima/maizuru_tsushima.html, in Japanese; Fig. 3c) to understand the effects of the flow on SST and the timing of the northern expansion of anchovy distribution. Surface chlorophyll *a* concentration (chl *a*) in spring was hypothesized to relate to the timing of breeding through the availability of local prey during the spring (krill and sandlance) and to the chick diets through availability of sandlance in summer. We used monthly (March to May) chl *a* in an area of 81 × 81 km around Teuri Island (Area B in Fig. 1, Fig. 3b) that was obtained from Ocean Color Web (<http://oceancolor.gsfc.nasa.gov/>) at a spatial resolution of 9 km by the SeaDAS 6.2 data analysis system reported after 1998 (<http://oceancolor.gsfc.nasa.gov/seadas/>).

Broad-scale climate indices are known to influence local climate and then marine ecosystems in the North Pacific (Chiba & Saino 2003, Minobe et al. 2004, Chiba et al. 2005). We used the Pacific Decadal Oscillation (PDO) index (www.data.kishou.go.jp/shindan/b-1/pdo/pdo.html), which reflects changes in air pressure and SST over the Pacific; the North Pacific Index (NPI; www.data.kishou.go.jp/db/)

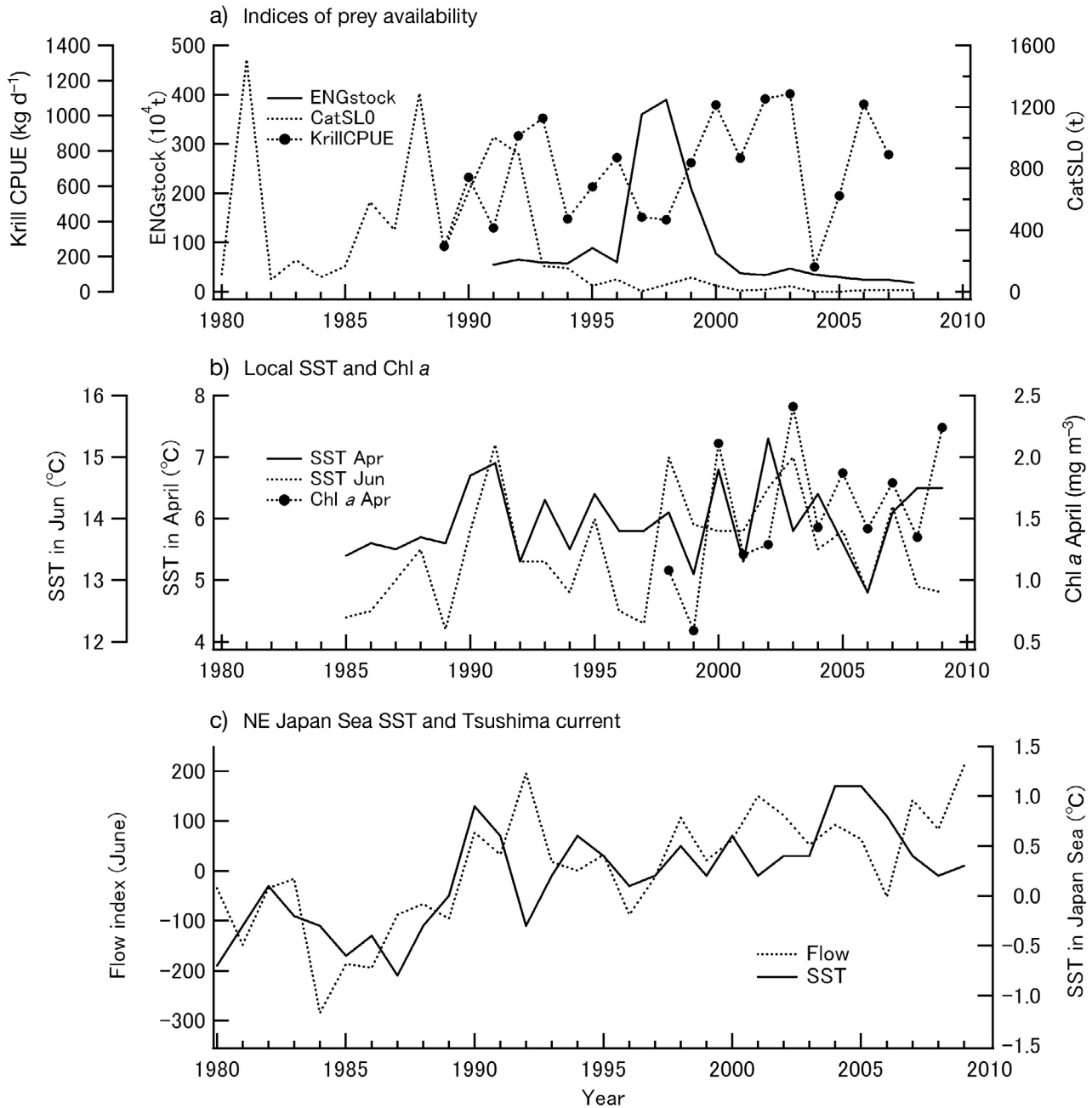


Fig. 3. (a) Estimated Tsushima anchovy *Engraulis japonicus* stock size (ENGstock), annual fishery harvest of 0+ sand lance *Ammodytes personatus* (CatSL0), and catch of krill *Thysanoessa longipes* and *T. inermis* per day (KrillCPUE) around Teuri Island. (b) Sea-surface temperature (SST) in Area A (Fig. 1) in April and June and surface chlorophyll *a* concentration in Area B (Fig. 1) in April. (c) Annual mean SST over the northeastern Japan Sea (www.data.jma.go.jp) and the Tsushima Current flow index in June

climate/pdo/npwin.txt), which reflects the strength of the Aleutian low pressure; and the Arctic Oscillation (AO) index (www.cpc.ncep.noaa.gov/products/precip/CWlink/daily_ao_index), which reflects the difference in air pressure between the Arctic and mid-latitudinal area of the North Pacific.

Analyses

To test for long-term interannual trends in seabird diet and breeding, we used simple linear regression analyses. To identify effects of local climatic factors and prey availability indices on variation in the tim-

ing of breeding, diet, and breeding success (the dependent variables), we used path analysis with Amos V. 6.0 software (Amos Development Corporation). For Temp., Rain, and SST in February to June, we used those in the month in which each had the strongest correlation (largest Pearson's r) with the dependent variables. In addition, we used annual values of Snow in March, CatSL0, KrillCPUE, and WWA. To simplify the analysis, indirect effects such as those of the TC flow on SST were not included in the models but examined separately. If correlation between potential independent variables was significant, co-variance between these was included in the potential path models. Year was included as a potential independent factor to test for trends in seabird diet and breeding performance. The model giving the smallest value of the Akaike Information Criterion (AIC) was selected as the best fitting model, and the top 3 models ($\Delta\text{AIC} < 2.0$) were considered as potential models (Burnham & Anderson 1998). Following Frederiksen et al. (2004), we examined whether the slopes of factors were significant using Type III sums of squares, and used the R^2 value (coefficient of determination) as the proportion of variance explained by each model.

The TC flow can indirectly affect the breeding biology and diets of seabirds through changes in SST and WWA (Watanuki et al. 2009). To look at the effects of the flow index on SST and WWA, we performed linear regression analyses using 1985 to 2009 data. To understand how broad-scale climate relates to the local climate factors, we performed path analyses using long-term data (1943 to 2008) in which PDO, NPI, and winter average AO were independent and Temp. in April, Snow, mean February wind speed recorded at Yagishiri Island, and the TC flow index in June were dependent variables.

Chl *a* data were collected from 1998. Because of the small sample size (12 yr), we used simple linear regression analyses to examine the effects of chl *a* on the availability of krill and sandlance that were harvested around the island and the proportion of these species in the diet.

RESULTS

Trends in climate and seabird parameters

We found no significant 1 yr lagged autocorrelations in the timing of breeding or breeding success of any species ($p > 0.05$). No significant inter-annual linear trends were found in the timing of breeding or

the breeding success in all 3 seabird species, or in the mass proportion of 0+ sandlance, >1+ sandlance, and sandlance in the diet of RHAU, JCOM, and BTGL, respectively ($p > 0.05$; Fig. 2). There were increasing trends in the mass proportion of anchovy in the diet of RHAU ($r = 0.788$, $n = 21$, $p < 0.01$) and BTGL ($r = 0.614$, $n = 18$, $p < 0.01$), though that for JCOM showed no trend ($p > 0.05$), presumably because no JCOM diet data were available before 1991 (Fig. 2c). Negative trends were found in the mass proportion of 0+ sandlance in the diet of JCOM ($r = -0.510$, $n = 18$, $p < 0.05$) and that of >1+ sandlance in the diet of RHAU ($r = 0.767$, $n = 21$, $p < 0.001$; Fig. 2d).

During the study period, there were increasing linear trends in Temp. ($r = 0.448$ to 0.539 , $n = 21$, $p < 0.05$), except in June ($p > 0.05$) and in the TC flow index ($r = 0.477$ to 0.658 , $n = 20$, $p < 0.05$). No trends were observed in Rain ($p > 0.05$), except in May ($r = 0.546$, $n = 21$, $p = 0.009$), Snow, SST and WWA ($p > 0.05$). ENGstock and KrillCPUE showed no significant trends ($p > 0.05$), but CatSL0 showed a decreasing trend ($r = 0.492$, $n = 19$, $p = 0.03$).

Timing of breeding

RHAU bred later in years with cold Temp. in April (Fig. 4), heavy Snow, and little Rain in April in the top 3 models, though ΔAIC were small among models (Table 2). JCOM bred later in years with cold air temperature and lower SST in April in the top 3 models, though as was seen for RHAU, ΔAIC were small among models (Fig. 4, Table 2). For the laying date of BTGL, ΔAIC were too small among top models to pick consistently significant factors (Table 2). Chl *a* peaked in April in 10 out of 12 yr and in May in the other years. Simple linear regression analyses indicate that Chl *a* in March to May did not relate to the timing of breeding for any species ($p > 0.05$).

Chick diet

In the 2 top models, RHAU fed more on anchovy in years when WWA was earlier, though ΔAIC were small among models (Table 3). However, surprisingly ENGstock was not included in the top 3 models (Table 3). JCOM fed more on 0+ sandlance in years with high CatSL0 in 3 top models (Table 3). The null model was selected as top model, and no factors explained the proportion of sandlance in the diet of BTGL (Table 3). Chl *a* in March to May did not relate to the spring prey availability indices (KrillCPUE:

$p > 0.05$; CatSL0: $p > 0.05$). Chl *a* in these months did not relate to the proportion of 0+ or >1+ sandlance in the diet of RHAU and BTGL ($p > 0.05$) either. There was significant negative correlation between Chl *a* in April and the proportion of 0+ sandlance in the diet of JCOM ($r = -0.629$, $n = 12$, $p = 0.03$), though its biological significance was unclear.

Breeding success

For RHAU, the top 3 models showed that breeding success was greater in years with earlier onset of breeding and earlier WWA (Table 4). For JCOM, the top 3 models showed that breeding success was greater in years with earlier breeding, lower Temp. and less Rain in June, earlier WWA, and greater CatSL0 (Table 4). The breeding success in BTGL varied widely between years and appeared to be greater in years with earlier WWA.

Interactions of local and large-scale climate

The TC flow index in May and June was negatively related to WWA ($r = -0.588$ to -0.495 , $n = 17$, $p = 0.03$ to 0.01), indicating that warm waters arrived later when the flow rate was reduced. June SST increased when the June TC flow was stronger ($r = 0.493$, $n = 17$, $p = 0.027$). Temp. in April was positively associated with the winter AO index in the top 3 models (Table 5). The mean wind speed in February was negatively associated with the winter AO index and positively associated with the NPI in the top 3 models (Table 5). Snow in March was negatively associated with winter AO and increased interannually in the top 2 models. The TC flow index in June was negatively associated with the PDO index and increased interannually (see Fig. 3c also) in the top 3 models. This indicates that anomalously low SST on the western side of the Pacific related to a weak TC in June.

Table 3. *Cerorhinca monocerata*, *Phalacrocorax filamentosus*, *Larus crassirostris*. Fitted path models relating the mass proportion of anchovy (for rhinoceros auklet [RHAU] and Japanese cormorant [JCOM]), 0+ and >1+ sandlance (for JCOM) or sandlance (for black-tailed gulls [BTGL]) in bird's diet to local sea-surface temperature (SST) in June, the date of arrival of 13°C water within the RHAU foraging range (WWA), the stock size of anchovy (ENGstock), and the annual catch of 0+ sandlance (CatSL0). Year is also included as a factor. Covariance between WWA and SST in June was significant ($r = -0.664$, $p < 0.05$) and was included in all models. Models having $\Delta AIC < 2.0$ but up to the third-best models are shown. ΔAIC , parameter estimates (with SE and significance in parentheses), and the coefficient of determination (R^2) are shown. – = independent variable not selected

Species	ΔAIC	Year	SST (Jun)	WWA	ENGstock	CatSL0	R^2
RHAU (anchovy)	0.000	3.258 (0.506, 0.001)	–	–1.787 (0.609, 0.003)	–	–	0.722
	1.087	2.988 (0.495, 0.001)	–	–1.874 (0.593, 0.002)	–	–	0.714
	1.203	3.023 (0.519, 0.001)	13.181 (4.765, 0.006)	–	–	–0.019 (0.018, 0.282)	0.679
JCOM (0+ sandlance)	0.000	–	–	–	–	0.039 (0.011, <0.001)	0.415
	0.366	–	3.789 (2.884, 0.189)	–	–	0.041 (0.011, <0.001)	0.484
	0.389	–	–	–0.481 (0.370, 0.194)	–	0.037 (0.011, <0.001)	0.442
JCOM (>1+ sandlance)	0.000	1.688 (0.979, 0.085)	–	–	–	–	0.148
	0.743	–	–	–	–	–	0.000
	1.644	–	–	–	–	–0.028 (0.026, 0.289)	0.065
JCOM (anchovy)	0.000	–	–	–	–	–	0.000
	0.918	–	–	–0.585 (0.559, 0.296)	–	–	0.062
BTGL (sandlance)	0.000	–	–	–	–	–	0.000
	0.631	–	–	–	–	0.034 (0.029, 0.249)	0.072
	1.053	–	–	–	0.057 (0.058, 0.330)	–	0.061

Table 4. *Cerorhinca monocerata*, *Phalacrocorax filamentosus*, *Larus crassirostris*. Fitted path models relating the number of fledglings per nest to the timing of breeding (hatching for rhinoceros auklet [RHAU] and Japanese cormorant [JCOM], laying for black-tailed gulls [BTGL]), monthly mean air temperature (Temp.) in June, total rain (Rain) in June, local sea-surface temperature (SST) in June, the date of arrival of 13°C water within the RHAU foraging range (WWA), the stock size of anchovy (ENGstock) and the annual catch of 0+ yr sand lance (CatSLO). Year is also included as a factor. Covariance between Temp. in June and SST in June ($r = 0.398$, $p < 0.05$) and between SST in June and WWA ($r = -0.664$, $p < 0.05$) were significant and included in all models. Models having $\Delta AIC < 2.0$ but up to the third-best models are shown. ΔAIC , parameter estimates (with SE and significance in parentheses), and the coefficient of determination (R^2) are also shown. – = independent variable not selected

Species	ΔAIC	Year	Timing	Temp. (Jun)	Rain (Jun)	SST (Jun)	WWA	ENGstock	CatSLO	R^2
RHAU	0.000	–	–0.013 (0.005, 0.005)	–	–	–	–0.012 (0.005, 0.015)	–	–	0.423
	0.116	–	–0.013 (0.004, 0.002)	–	–	–	–0.012 (0.005, 0.012)	0.000 (0.000, 0.146)	–	0.490
	0.138	–	–0.013 (0.004, 0.002)	–	–	–0.071 (0.046, 0.128)	–0.018 (0.006, 0.002)	0.000 (0.000, 0.076)	–	0.554
JCOM	0.000	0.052 (0.009, <0.001)	–0.024 (0.005, <0.001)	–0.380 (0.061, <0.001)	–0.004 (0.002, 0.061)	–	–0.031 (0.007, <0.001)	–	0.001 (0.000, <0.001)	0.902
	1.865	0.051 (0.009, <0.001)	–0.025 (0.004, <0.001)	–0.370 (0.060, <0.001)	–0.003 (0.002, 0.077)	–0.039 (0.081, 0.635)	–0.033 (0.010, 0.001)	–	0.001 (0.000, <0.001)	0.914
	1.880	0.049 (0.009, <0.001)	–0.024 (0.005, <0.001)	–0.389 (0.061, <0.001)	–0.004 (0.002, 0.069)	–	–0.031 (0.007, <0.001)	0.000 (0.000, 0.675)	0.001 (0.000, <0.001)	0.909
BTGL	0.000	–	–	–	–	–	–0.024 (0.013, 0.060)	–	–	0.179
	1.144 1.146	– –	– –0.048 (0.051, 0.345)	– –	– –	– –	–0.026 (0.012, 0.037)	– –	– –	0.000 0.241

DISCUSSION

We conducted an analysis of seabird–climate relationships for the northern Japan Sea, thereby updating previous long-term studies of this region, prey and seabirds (Takahashi et al. 2001, Deguchi et al. 2004, Ito et al. 2009, Watanuki et al. 2009). In the present paper, we focused on examining trends in climatic factors and seabird parameters as our time series is now of sufficient duration to investigate directional climate change as well as climate variability. The present study is significant, as relatively few long-term seabird studies have been carried out in the western North Pacific (Kitaysky & Golubova 2000, Watanuki et al. 2009). Our approach was to test for trends in large-scale and local climate indices and seabird parameters, and to relate factors using regression and AIC model selection procedures. We first tested the seabird parameters and found no evidence of autocorrelation.

Timing of breeding

Climate factors affected the timing of breeding, but differently between species. RHAU bred earlier in years with a warm and wet spring and little snow, probably because frozen soil and snow prevent them from excavating their burrows (Watanuki 1987a, Watanuki et al. 2009). JCOM bred earlier in years with higher spring SST and air temperature. Catches of adult Japan Sea greenlings in the study region tended to be higher in years with a warmer winter (Hoshino et al. 2009), and flatfish come to shore during early spring and spawn earlier in warmer regions (Nagasawa & Torisawa 1991, Minami 1995). Thus, the abundance and seasonality of these prey species of JCOM in spring (M. Ito unpubl. data) might be related to the timing of JCOM. BTGL laid eggs earlier in years when the spring SST was from 3.0 to 4.9°C (Tomita et al. 2009), since swarms of krill, i.e. the main prey of BTGL in spring, occur at the surface with this range of SSTs in this region (Hanamura et al. 1989). However, we could not find any consistent linear effects of spring SST and krill CPUE on BTGL timing (Fig. 4, Table 2), so that longer term studies including other climate factors and more appropriate prey availability measures should be carried out before conclusions can be drawn.

Table 5. Fitted path models relating local physical factors (air temperature in April at Yagishiri, Temp. [Apr]; average wind speed at Yagishiri in February, Wind [Feb]; total snowfall in March at Haboro, Snow [Mar]; Tsushima current flow index in June, Flow [Jun]; to Pacific Decadal Oscillation [PDO], North Pacific Index [NPI] and average winter Arctic Oscillation [AOav; December to February]) index. Year is also included as a factor. Covariance between Year and PDO ($r = 0.310$, $p < 0.05$), between year and AO (winter) ($r = 0.345$, $p < 0.001$), and between PDO and NPI ($r = -0.531$, $p < 0.01$) were significant and included in all models. Models having $\Delta AIC < 2.0$ but up to the third-best models are shown. ΔAIC , parameter estimates (with SE and significance in parentheses), and the coefficient of determination (R^2) are shown. – = independent variable not selected

	ΔAIC	Year	PDO	NPI	AOav	R^2
Temp. (Apr)	0.000	–	–	–	1.071 (0.335, 0.001)	0.142
	0.924	0.008 (0.007, 0.285)	–	–	0.901 (0.368, 0.014)	0.148
	1.201	–	–	0.144 (0.161, 0.370)	1.014 (0.334, 0.002)	0.142
Wind (Feb)	0.000	–	–	0.312 (0.130, 0.017)	–0.687 (0.288, 0.017)	0.170
	1.639	–	–0.066 (0.154, 0.668)	0.269 (0.154, 0.080)	–0.680 (0.288, 0.018)	0.165
	1.974	0.001 (0.006, 0.849)	–	0.319 (0.308, 0.014)	–0.708 (0.308, 0.022)	0.174
Snow (Mar)	0.000	0.478 (0.211, 0.023)	–	–	–24.739 (10.329, 0.017)	0.124
	1.071	0.543 (0.218, 0.013)	–4.372 (4.521, 0.334)	–	–26.775 (10.295, 0.009)	0.144
Flow (Jun)	0.000	4.383 (0.664, <0.001)	–55.414 (14.794, <0.001)	–	–	0.499
	0.535	4.078 (0.708, <0.001)	–52.700 (14.654, <0.001)	–	40.033 (33.173, 0.228)	0.515
	1.271	4.296 (0.666, <0.001)	–46.814 (16.878, <0.001)	14.138 (15.849, 0.372)	–	0.505

Using path analyses the effect of year on the timing of JCOM was significant (Table 2) presumably because JCOM timing was extraordinary late in 2005 and 2006 (Fig. 2a). Thus, at least between 1992 and 2009, there did not appear to be long-term linear trends in the timing of breeding in seabirds in our region. During the past 30 to 50 yr, the timing of seabird breeding has advanced in the Arctic (Gaston et al. 2009, Moe et al. 2009), but has been delayed in the Antarctic (Barbraud & Weimerskirch 2006). In the temperate zone, results are more variable; laying or hatching occurs later in the western North Sea (Fredriksen et al. 2004, Wanless et al. 2008), where the North Atlantic Oscillation index has been decreasing, but earlier in the Canadian Pacific (Bertram et al. 2001, but see Hipfner at www.dfo-mpo.gc.ca/CSAS/Csas/publications/resdocs-docrech/2010/2010_053_e.pdf), where SST has been increasing. No apparent trends were observed for auklets off the California coast (Thayer & Sydeman 2007, Schroeder et al. 2009), but murrelets have been observed earlier (W. Sydeman pers. comm.). No change has been observed for Southeast Alaska (Slater & Byrd 2009). Thus, the impact of global warming on the timing of

seabird breeding appears to be variable among regions, presumably depending on the local ecosystem and climate system. There could also be age-dependent differences (Pinaud & Weimerskirch 2002) in seabird responses to climate variability and change. We cannot evaluate this hypothesis for our unmarked populations in the Japan Sea, but the variability observed in global patterns of seabird timing in relation to climate certainly may be related to age-specific breeding phenology and the age structure of studied seabird populations.

Chick diet

Decadal and interannual variability of local climate factors and prey availability affected the seabird diet. RHAU and BTGL fed on sardine and sandlance from 1984 to 1987, but RHAU fed on anchovy and BTGL on sandlance after 1992 (see also Deguchi et al. 2004). These decadal changes in chick diet reflected the cold-to-warm regime shift in the late 1980s in the TC region (see also Fig. 3c), with a steep decrease in the catch of cold-water sardine and an increase in the

catch of warm-water anchovy (Tian et al. 2008). Anchovy spawn in the western North Pacific at a SST of 15 to 28°C, and sardine, at 13 to 20°C (Takasuka et al. 2008). Cumulative mortality through the early life stages of sardine between 1988 and 1991, when SST was higher, apparently caused the stock to crash (Watanabe et al. 1995).

Interannual variation in the proportion of anchovy in the diet of RHAU was explained by the timing of the arrival of warm water (SST = 13°C) within the bird's foraging range. In years with a strong TC, RHAU switched prey earlier from cold-water species (sandlance and juvenile greenlings; Nagasawa & Torisawa 1991) to anchovy and hence fed more on anchovy in these years (Watanuki et al. 2009). The annual catch of 0+ sandlance was positively related to the proportion of 0+ sandlance in the diet of JCOM, supporting the relationship between the seabird diet and local prey availability. The proportion of sandlance in the diet of BTGL, however, could not be explained by either local climate factors or the catch of 0+ sandlance. This is surprising as BTGL are surface feeders and feed mainly on sandlance (Deguchi et al. 2004; Table 1, Fig. 2d). Possibly because of a narrow foraging habitat, BTGL might not be able to switch to anchovy when they become available.

Breeding success

There were no trends in the breeding success of seabirds on Teuri Island. Breeding success of RHAU was determined by the arrival of warm water in the northern Japan Sea. Increasing breeding success was correlated with earlier arrival dates of the warm TC because anchovy were advected into this region with the warm waters and provided more energy to chicks than other prey (Watanuki et al. 2009). Yet, we found no evidence that breeding success was significantly influenced by the anchovy stock size (Table 4). The seasonal expansion of the anchovy distribution to the Teuri seabird foraging area might be important rather than overall abundance of this Tsushima anchovy stock, which was measured in the southern areas of the Japan Sea. In years when the local availability of epipelagic fish (anchovy and sandlance) was low, JCOM fed on benthic and epibenthic fishes in coastal areas 30 to 40 km from the colony, and, hence, their feeding frequency was reduced (Watanuki et al. 2004), ultimately affecting chick growth and survival (Kato et al. 2001). Similar to RHAU, breeding success of JCOM was greater in years when warm waters arrived earlier, i.e. high local anchovy availability. An-

chovy stock size had no significant relationship with this species either. The local annual catch of 0+ sandlance (important prey of JCOM; Table 1) positively affected breeding success, also indicating the importance of the availability of this prey.

Why the breeding success of RHAU and JCOM was higher in years when they bred earlier, as reported in common murre *Uria aalge* in the North Sea (Votier et al. 2009) and cassin's auklet and common murre in California (Schroeder et al. 2009), is unclear. RHAU and JCOM feed on different prey during the pre-laying, laying, and chick-rearing periods (Kato et al. 2001, Ito et al. 2009, M. Ito unpubl. data), yet both species showed an inverse relationship between timing and success. The effects of SST on prey availability could differ between spring (egg laying) and summer (chick rearing). With simple regression analyses, no annual trends were observed in the breeding success of JCOM, though positive effects of year on it (Table 4) could be related to interannual trends in prey availability and timing of breeding.

No climate factors or prey availability indices explained the large interannual variations in the breeding success of BTGL. The gulls fed mostly on sandlance (Fig. 2d). As found in the North Sea (Furness & Tasker 2000), BTGL that feed mainly on sandlance might be sensitive to changes in the availability of sandlance, but the catch of 0+ sandlance was not correlated with their breeding success. The potential impact by feral cats on nesting BTGL (Watanuki 2010) might make it difficult to detect the climate factors and prey availability that are associated with their breeding performance.

In regions where seabirds feed mainly on sandlance, climate factors have affected their breeding success through bottom-up effects (Aebischer et al. 1990, Frederiksen et al. 2006). In our region, however, variability of local chl *a* concentration did not relate to the availability indices of local seabird prey (krill and 0+ sandlance), the proportion of these prey in the diet, or the seabird timing of breeding. Rather, interannual change in the availability of anchovy, an important alternative prey to sandlance, that appeared to be driven by the warm TC seemed to be important in this area. Alternatively, chl *a* could be a poor indicator of the abundance of availability of forage fish prey.

CONCLUSIONS

Interannual variations in broad-scale atmospheric indices, the AO and the Aleutian Low (manifested as

the PDO), are linked to winter wind stress and the summer flow rates of the TC (Minobe et al. 2004, present study), respectively. These factors influence the chick diets and breeding performance of 2 species of seabirds in northern Japan Sea, but not a third that may be responding, mainly, to terrestrially based factors (invasive cats). In this area the seasonal range expansion of warm waters and anchovy plays a key role in driving chick diet and breeding performance of seabirds. Our study indicates that the mechanisms of response from climate to seabirds are specific to the regional ecosystem under study as well as variation in the life histories of the seabirds under consideration.

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LITERATURE CITED

- Aebischer NJ, Coulson JC, Colebrook JM (1990) Parallel long-term trends across four marine trophic levels and weather. *Nature* 347:753–755
- Anderson PJ, Piatt JF (1999) Community reorganization in the Gulf of Alaska following ocean climate regime shift. *Mar Ecol Prog Ser* 189:117–123
- Barbraud C, Weimerskirch H (2006) Antarctic birds breed later in response to climate change. *Proc Natl Acad Sci USA* 103:6248–6251
- Bertram DF, Mackas DL, McKinnell SM (2001) The seasonal cycle revisited: interannual variation and ecosystem consequences. *Prog Oceanogr* 49:283–307
- Burnham KP, Anderson DR (1998) Model selection and inference: a practical information-theoretic approach. Springer, New York, NY
- Chiba S, Saino T (2003) Variation in mesozooplankton community structure in the Japan/East Sea (1991–1999) with possible influence of the ENSO scale climatic variability. *Prog Oceanogr* 57:317–339
- Chiba S, Hirota Y, Hasegawa S, Saino T (2005) North–South contrasts in decadal scale variations in lower trophic-level ecosystems in the Japan Sea. *Fish Oceanogr* 14: 401–412
- Chochi M, Niizuma Y, Takagi M (2002) Sexual differences in the external measurements of black-tailed gulls breeding on Rishiri Island, Japan. *Ornithology Sci* 1: 163–166
- Deguchi T, Watanuki Y, Niizuma Y, Nakata A (2004) Interannual variations of the occurrence of epipelagic fish in the diets of the seabirds breeding on Teuri Island, northern Hokkaido, Japan. *Prog Oceanogr* 61:267–275
- Durant JM, Anker-Nilssen T, Hjermmann DO, Stenseth NC (2004) Regime shift in the breeding of an Atlantic puffin population. *Ecol Lett* 7:388–394
- Durant JM, Hjermmann DO, Anker-Nilssen T, Beaugrand G, Myrsterud A, Pettorelli N, Stenseth NC (2005) Timing and abundance as key mechanisms affecting trophic interactions in variable environments. *Ecol Lett* 8:952–958
- Frederiksen M, Harris MP, Daunt F, Rothery P, Wanless S (2004) Scale-dependent climate signals drive breeding phenology of three seabird species. *Glob Change Biol* 10:1214–1221
- Frederiksen M, Edwards M, Richardson AJ, Halliday NC, Wanless S (2006) From plankton to top predators: bottom-up control of a marine food web across four trophic levels. *J Anim Ecol* 75:1259–1268
- Frederiksen M, Edwards M, Mavor RA, Wanless S (2007) Regional and annual variation in black-legged kittiwake breeding productivity is related to sea surface temperature. *Mar Ecol Prog Ser* 350:137–143
- Furness RW, Tasker ML (2000) Seabird–fishery interactions: quantifying the sensitivity of seabirds to reductions in sandeel abundance, and identification of key areas for sensitive seabirds in the North Sea. *Mar Ecol Prog Ser* 202:253–264
- Gaston AJ, Jones IL (1998) The auks. Oxford University Press, Oxford
- Gaston AJ, Gilchrist HG, Mallory ML, Smith PA (2009) Changes in seasonal events, peak food availability, and consequent breeding adjustment in a marine bird: a case of progressive mismatching. *Condor* 111:111–119
- Grémillet D, Lewis S, Drapeau L, van Der Lingen CD and others (2008) Spatial match–mismatch in the Benguela upwelling zone: Should we expect chlorophyll and sea-surface temperature to predict marine predator distribution? *J Appl Ecol* 45:610–621
- Hanamura Y, Kotori M, Hamada S (1989) Daytime surface swarms of the euphausiid *Thysanoessa inermis* off the west coast of Hokkaido, northern Japan. *Mar Biol* 102: 369–376
- Hase H, Yoon JH, Koterayama W (1999) The current structure of the Tsushima warm current along the Japanese coast. *J Oceanogr* 55:217–235
- Hoshino N, Takashima T, Watanobe M, Fujioka T (2009) Age-structures and catch fluctuations of arabesque greenling (*Pleurogrammus azonus*) in the coastal area of southern Hokkaido, Japan. *Sci Rep Hokkaido Fish Exp Stn* 76:1–11
- Ishikawa K, Watanuki Y (2002) Sex and individual differences in foraging behavior of Japanese cormorants in years of different prey availability. *J Ethol* 20:49–54
- Ito M, Minami H, Tanaka Y, Watanuki Y (2009) Intra- and inter-annual oceanographic changes as inferred from the diet of a piscivorous seabird. *Mar Ecol Prog Ser* 393: 273–284
- Jenouvrier S, Caswell H, Barbraud C, Holland M, Stroeve J, Weimerskirch H (2009) Demographic models and IPCC climate projections predict the decline of an emperor penguin population. *Proc Natl Acad Sci USA* 106:1844–1847
- Kato A, Watanuki Y, Naito Y (2001) Foraging and breeding performance of Japanese cormorants in relation to prey. *Ecol Res* 16:745–758
- Kato A, Watanuki Y, Naito Y (2003) Foraging behaviour of chick-rearing rhinoceros auklets at Teuri Island, Japan, determined by acceleration-depth recording micro data loggers. *J Avian Biol* 34:282–287
- Kitaysky AS, Golubova EF (2000) Climate change causes contrasting trends in reproductive performance of planktivorous and piscivorous alcids. *J Anim Ecol* 69:248–262

- Kuroki M, Kato A, Watanuki Y, Takahashi A (1998) Artificial nest boxes for the study of breeding ecology of rhinoceros auklets *Cerorhinca monocerata*. J Yamashina Inst Ornithol 30:40–46 (in Japanese with English abstract)
- Kuroki M, Kato A, Watanuki Y, Niizuma Y, Takahashi A, Naito Y (2003) Diving behavior of an epipelagically feeding alcid, the rhinoceros auklet (*Cerorhinca monocerata*). Can J Zool 81:1249–1256
- Loeb V, Siegel V, Holm-Hansen O, Hewitt R, Fraser W, Trivelpiece W, Trovelpiece S (1997) Effects of sea-ice extent and krill or salp dominance on the Antarctic food web. Nature 387:897–900
- Mihara Y (1998) Distribution of the Japanese anchovy, *Engraulis japonicus*, off southeastern Hokkaido. Sci Rep Hokkaido Fish Exp Stn 53:9–14 (in Japanese with English abstract)
- Minami T (1995) Spawning season of pleuronectid flatfishes in the coastal waters of Hokkaido, Japan (review). Bull Hokkaido Natl Fish Res Inst 59:69–80
- Minobe S, Sako A, Nakamura M (2004) Interannual to interdecadal variability in the Japan Sea based on a new gridded upper water temperature dataset. J Phys Oceanogr 34:2382–2397
- Moe B, Stempniewicz L, Jakubas D, Angelier F and others (2009) Climate change and phenological responses of two seabird species breeding in the high-Arctic. Mar Ecol Prog Ser 393:235–246
- Nagasawa K, Torisawa M (1991) Fish and marine invertebrates of Hokkaido: biology and fisheries. Kita-nihon Kaiyo Center, Sapporo (in Japanese)
- Nelson JB (2005) Pelicans, cormorants, and their relatives. Pelecaniformes. Oxford University Press, Oxford
- Niizuma Y, Takagi M, Senda M, Chochi M, Watanuki Y (2005) Incubation capacity limits maximum clutch size in black-tailed gulls *Larus crassirostris*. J Avian Biol 36:421–427
- Olssen KM, Larsson H (2004) Gulls of North America, Europe and Asia. Princeton University Press, Princeton
- Osa Y, Watanuki Y (2002) Status of seabirds breeding in Hokkaido. J Yamashina Inst Ornithol 33:107–141
- Pinaud D, Weimerskirch H (2002) Ultimate and proximate factors affecting the breeding performance of marine top predator. Oikos 99:141–150
- Schreiber EA (2002) Climate and weather effects on seabirds. In: Schreiber EA, Burger J (eds) Biology of marine birds. CRC Press, Boca Raton, FL, p 179–215
- Schroeder ID, Sydeman WJ, Sarkar N, Thompson SA, Bograd SJ, Schwing FB (2009) Winter pre-conditioning of seabird phenology in the California Current. Mar Ecol Prog Ser 393:211–223
- Slater L, Byrd GV (2009) Status, trends, and patterns of covariation of breeding seabirds at St Lazaria Island, southeast Alaska, 1994–2006. J Biogeogr 36:465–475
- Takahashi A, Kuroki M, Niizuma Y, Kato A, Saito A, Watanuki Y (2001) Importance of the Japanese anchovy *Engraulis japonicus* to breeding rhinoceros auklets *Cerorhinca monocerata* on Teuri Island, Sea of Japan. Mar Biol 139:361–371
- Takasuka A, Oozeki Y, Kubota H, Lluch-Cota SE (2008) Contrasting spawning temperature optima: Why are anchovy and sardine regime shifts synchronous across the North Pacific? Prog Oceanogr 77:225–232
- Thayer JA, Sydeman WJ (2007) Spatio-temporal variability in prey harvest and reproductive ecology of a piscivorous seabird, *Cerorhinca monocerata*, in an upwelling system. Mar Ecol Prog Ser 329:253–265
- Tian Y, Kidokoro H, Watanabe T, Iguchi N (2008) The late 1980s regime shift in the ecosystem of Tsushima warm current in the Japan/East Sea: evidence from historical data and possible mechanisms. Prog Oceanogr 77:127–145
- Tomita N, Niizuma Y, Takagi M, Ito M, Watanuki Y (2009) Effect of interannual variations in sea-surface temperature on egg-laying parameters of black-tailed gulls (*Larus crassirostris*) at Teuri Island, Japan. Ecol Res 24:157–162
- Votier SC, Hatchwell BJ, Mears M, Birkhead TR (2009) Changes in the timing of egg-laying of a colonial seabird in relation to population size and environmental condition. Mar Ecol Prog Ser 393:225–233
- Wanless S, Harris MP, Lewis S, Frederiksen M, Murray S (2008) Later breeding in northern gannets in the eastern Atlantic. Mar Ecol Prog Ser 370:263–269
- Watanabe Y, Zenitani H, Kimura R (1995) Population decline of the Japanese sardine *Sardinops melanostictus* owing to recruitment failures. Can J Fish Aquat Sci 52:1609–1616
- Watanuki Y (1987a) Breeding biology and foods of rhinoceros auklets on Teuri Island, Japan. Proc NIPR Symp Polar Biol 1:175–183
- Watanuki Y (1987b) Inter-species, inter- and intra-colony differences in food habits and breeding of *Larus* gulls. PhD thesis, Hokkaido University, Hakodate (in Japanese)
- Watanuki Y (2010) Cat, human and seabirds on Teuri Island. Nature of Teuri Island 31:1–2 (in Japanese)
- Watanuki Y, Kato A, Naito Y (1996) Diving performance of male and female Japanese cormorants. Can J Zool 74:1098–1109
- Watanuki Y, Ishikawa K, Takahashi A, Kato A (2004) Foraging behavior of a generalist marine top predator, Japanese cormorants, in years of demersal vs. epipelagic prey. Mar Biol 145:427–434
- Watanuki Y, Ito M, Deguchi T, Minobe S (2009) Climate-forced seasonal mismatch between the hatching of rhinoceros auklets and the availability of anchovy. Mar Ecol Prog Ser 393:259–271
- Yeh SW, Park YG, Min H, Kim CH, Lee JH (2010) Analysis of characteristics in the sea surface temperature variability in the East/Japan Sea. Prog Oceanogr 85:213–223

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