Landscape-mediated variation in diet is associated with egg size and maculation in a generalist forager

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Human impacts alter landscapes with consequences for the distribution and availability of high-quality food resources to populations inhabiting those landscapes, which may impact on the reproductive output of individuals in those populations. The sensitivity of wild populations to changes in food resources may vary among stages of the annual cycle. For example, in birds, effects are likely to be greater during costly stages such as egg production. Here we compare assimilated diet (from stable isotope analysis of chick feathers) and egg traits (egg size, shape, eggshell colour and maculation, using pattern-analysis software) in Herring Gulls Larus argentatus, across seven colonies in southwest Scotland and Northern Ireland. The Herring Gull is an opportunistic, generalist forager on both marine and terrestrial resources which frequently exploits anthropogenic food sources such as fishery discards and human refuse. We found that larger eggs were laid in colonies where females consumed either a higher proportion of marine resources or terrestrial resources; smaller eggs were laid in colonies where females had an intermediate diet. In colonies where females consumed more marine items, they also laid eggs with higher maculation (intensity and size of spots) compared with colonies where females mainly consumed terrestrial food. We also found smaller and more pointed eggs, suggestive of resource shortages, in larger colonies. Generalist foragers are often thought to have the capacity to buffer themselves against changes in the food web, provided that enough alternative food is available. However, this study highlights that specializing on the most profitable or available resources has consequences for egg traits even in an opportunistic generalist forager exploiting a large range of habitats. If variation in egg traits is related to reproductive output, then understanding the impact of assimilated diet on reproduction early in the breeding season can provide important insights into how populations will respond to landscapes altered by human impact.

Keywords: conservation, gull, intertidal, Larus, marine, pattern-analysis software, resource use, stable isotopes.

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Human impact on the landscape can affect the quantity, quality and configuration of resources on which animals depend (Benton et al. 2003, Chace & Walsh 2006, Halpern et al. 2008). Landscape composition can affect fitness-related traits with potential consequences for demographic parameters (e.g. Kurki et al. 2000, Penteriani et al. 2002, Bond et al. 2004). Understanding the relationship...
between landscape characteristics and fitness-related traits can therefore provide insights into the conservation and management of populations inhabiting human-impacted landscapes.

Human-impacted landscapes can have reduced or degraded foraging habitat but may also provide new foraging opportunities. Consequent changes in resource availability may influence a consumer’s diet and thus the composition of energy and nutrients consumed (Amar & Redpath 2005, Palma et al. 2006). The quantity and quality of consumed resources can have important consequences for survival and reproductive performance (Alverson 1992, Grémillet et al. 2008, Österblom et al. 2008, White 2008). Optimal foraging theory predicts that energy gain, and associated fitness, is maximized by specializing on the single most abundant and/or highest quality food (Stephens & Krebs 1986). However, specialist foragers may be negatively affected when landscape changes impact their food availability (Montevecchi 1993, Davoren & Montevecchi 2003, Vucetich & Peterson 2004, Millon & Bretagnolle 2008). Conversely, generalist foragers are thought to buffer themselves against changes in food availability by switching to alternatives (Schoener 1971, Pyke et al. 1977). However, how human-impacted landscapes influence generalists’ resource consumption and hence their reproduction is not known in many wild populations (Yohannes et al. 2016).

Sensitivity to resource availability may vary among stages of the annual cycle (Lindstrom et al. 1993, Perrins 1996, Wikelski & Tarlow 2003) and is likely to be greater during nutritionally and energetically demanding periods. Egg formation in birds is a particularly demanding stage (Robbins 1981, Monaghan & Nager 1997). Egg size is positively related to the abundance and quality of resources during egg formation in birds (Bolton et al. 1992, Christians 2002, Nager 2006, Sorensen et al. 2009, Barrett et al. 2012, Kouwenberg et al. 2013, Bennett et al. 2017). Egg shape may also depend on resource availability during egg formation (Ardendt 2004, Górski et al. 2015). Furthermore, food availability and quality has been shown to influence variation in egg pigmentation (eggshell colour and maculation) in several bird species, with less pigment deposited at low availability of overall (Avilés et al. 2006, Duval et al. 2013) or specific resources (i.e. calcium, Gosler et al. 2005). There is evidence that offspring development and survival can depend on egg size (reviewed by Krist 2011) and egg pigmentation (Maurer et al. 2011, Lahti & Ardia 2016, Hargitai et al. 2018). Thus, female birds living in human-impacted landscapes may find their preferred food reduced or may need to rely on alternatives of different quality which may affect egg quality and, ultimately, demographic rates.

One group of generalist foragers that increasingly use human-impacted landscapes are the large gulls, Laridae (Rock 2005). Even when breeding, and constrained to a central nesting site, gulls can move over a large area and exploit a variety of resources from marine food, consisting of pelagic fish and intertidal invertebrates, to terrestrial food sources, including those derived from human refuse and farming (Hunt 1972, Götmark 1984, O’Hanlon et al. 2017). As their marine food webs have been simplified and impoverished by human impact (Halpern et al. 2008), many European gull populations are consuming anthropogenic terrestrial food from farmland and built-up areas (Pons & Migot 1995, Coulson & Coulson 2008, Schwemmer et al. 2008, Garthe et al. 2016, Gyimesi et al. 2016). Marine foods, in particular fish, are thought to best meet the high energy and nutrient requirements of laying female gulls (Hiom et al. 1991), but some populations now rely on anthropogenic, terrestrial food resources (e.g. Pons & Migot 1995, Steigerwald et al. 2015, Gyimesi et al. 2016). Across southwest Scotland and Northern Ireland, there is a strong relationship between the local landscape and diet of breeding Herring Gulls Larus argentatus. Gulls predominantly forage on the locally most available resources, and colonies with a higher proportion of marine resources in their diet have a higher breeding success (O’Hanlon et al. 2017). The marine diet consists mainly of intertidal invertebrates (crustaceans and bivalves) with few marine fish, consistent with the absence of any significant fishery in the region, and therefore limited access to discards. Terrestrial food consists mainly of grain, although terrestrial invertebrates and anthropogenic food waste are also consumed (O’Hanlon et al. 2017). Here we test the relationship between assimilated diet during egg formation, based on stable isotope analyses of chick down feathers, and egg traits from seven Herring Gull colonies across southwest Scotland and Northern Ireland. These colonies occupy different landscapes, with varying availability of marine and terrestrial resources. We predict that colonies consuming a more marine diet, at a higher
trophic level, produce larger and more pigmented eggs, measured as more intense eggshell colour and maculation (intensity and size of spots). This study aims to contribute to our understanding of how altered landscapes impact egg formation of a generalist seabird.

**METHODS**

Fieldwork was carried out between 5 May and 11 July 2014 across seven Herring Gull colonies located in southwest Scotland and Northern Ireland covering an approximate area of 200 km x 250 km (Fig. 1, Table 1). In this region, we selected colonies that had within their foraging ranges low (1–2 km² built-up area > 10 km away) to high human population density (over 400 km² built-up area as close as 2–3 km), and rocky intertidal habitat with low wave fetch, supporting a diverse and abundant invertebrate community (Burrows 2012), to resource-poorer, exposed coastlines with high wave fetch (Table 1). The colonies typically contained several gull species and we identified Herring Gull nests by direct observations of attending adults. Within our study colonies, clutch initiation spread over approximately 3 weeks starting in early May, with the incubation period lasting 4 weeks. We visited each colony for a few days every 2 weeks (Table 1). Thus, at each visit only a snapshot of clutches and chicks were available that were suitable for taking egg measurements during incubation and for sampling down of chicks in their first week of life for stable isotope analysis. Consequently, we could not take assimilated diet and egg measurements from the same nests. There was no bias that a colony visited during early egg-laying was also visited during early chick-rearing (Spearman rank correlation: \( r_s = 0.16, n = 7, P = 0.73 \)).

**Egg measurements**

During short colony visits during the incubation period between 5 May and 3 June, we took digital photographs of Herring Gull clutches in situ, under standardized conditions, to minimize the duration of disturbance (Fig. S1). The modal clutch size of Herring Gulls is three eggs (Pierotti 1982). In this study it was unknown whether smaller clutches had been subject to partial nest predation, so we did not know clutch size at laying. Thus, we include clutch size in data analyses. From these digital photographs we measured four egg traits: size, shape, eggshell colour and maculation (a more detailed account of how we did this is provided in Appendix S1). Egg size, measured as egg volume, and shape were measured using the Egg Measurement Tool plugin developed by Troscianko (2014) in IMAGEJ (Schneider et al. 2012). We measured egg length (\( l \)) and width (\( w \)) to calculate egg volume (\( V \)), using \( V = klw^2 \) (Hoyt 1976) where \( k \) is a species-specific shape constant, here taken as 0.000476 (Harris 1964). Egg shape was calculated as the deviation from a perfect ellipse, with a higher shape index indicating more pointed eggs (Fig. S2). We measured eggshell colour from the background, excluding maculation, as red (R), blue (B) and green (G) values in IMAGEJ. Eggshell colour was represented by the green/blue (G/B) ratio, with a high G/B ratio indicating brown-coloured and a low G/B ratio paler, green-blueish coloured eggs (Fig. S3). We also measured maculation, which is a complex trait comprising the distribution, intensity and size of spots (Gosler et al. 2000). A range of approaches have been developed to measure maculation (Gosler et al. 2005, Stoddard et al. 2014, Wegmann et al. 2015, Brulez et al. 2016, Gómez & Liñán-Cembrano 2016). We used NaturePatternMatch (Stoddard et al. 2014), which determined the similarity of an egg to an unmaculated reference (duck) egg. This measure considers the intensity and size of spots but not distribution, with the less maculated eggs (fewer and lighter spots) having a higher similarity score (Fig. S4). The number of sampled nests varied among colonies owing to differences in colony size, the proportion of nests that were accessible and the time available to take photographs (Table 1). Egg size, shape and eggshell colour were determined for a total of 514 eggs from 194 clutches. Maculation could only be determined for 318 eggs from 128 clutches; 66 clutches (34%) were excluded as one or more eggs within the clutch were dirty or had started to pip, which would have biased the estimate of the similarity score, or contained eggs that returned no maculation score, as the software rejected images due to insufficient contrasts. The proportion of eggs rejected for maculation scoring did not differ across colonies (Chi-square test: \( \chi^2 = 0.62, P = 0.996 \)).

**Assessment of assimilated diet**

We collected samples of several down feathers from different areas of the body (head, back and sides) from chicks from accessible nests in each
colony at the appropriate developmental stage (see Table 1 for sample sizes). Sampled chicks needed to be less than 1 week old based on known age or judged from chick development (Kadlec et al. 1969) to ensure that we collected only feather material that was grown by the chicks while developing in the egg. Thus, their carbon and nitrogen stable isotope signature reflects the resources that the female gulls consumed during egg formation (Klaassen et al. 2004).

Collected feathers were washed in liquid detergent (Ecover) diluted with deionized water (approximate 1 : 99 dilution), followed by a wash in a 2 : 1 mixture of chloroform/methanol. Dried samples were then homogenized. Where more than one chick from the same brood was sampled, feather material from all chicks of that brood were homogenized together to provide one sample per brood. Homogenized samples of feathers were weighed (mass between 0.7 and 0.8 mg) and sealed in tin capsules, combusted and analysed for carbon and nitrogen isotopes by continuous-flow isotope ratio mass spectrometry (using a Costech Elemental Analyser (Milan, Italy) linked to a Thermo Finnigan Delta Plus XP Mass Spectrometer (Bremen Germany)) at the NERC Life Sciences Mass Spectrometry Facility in East Kilbride (UK). The stable isotope ratios are expressed as $\delta^{13}C$ ($^{13}C/^{12}C$) and $\delta^{15}N$ ($^{15}N/^{14}N$) in parts per thousand ($\%$) relative to the international references Vienna PeeDee belemnite (V-PDB) marine fossil limestone for carbon, and atmospheric N$_2$ for nitrogen, respectively. Measurement accuracy was $\pm0.09\%$ for $\delta^{13}C$ and $\pm0.12\%$ for $\delta^{15}N$ based on the standard deviation of the repeated analyses of tryptophan as a lab standard. The nitrogen ratio reflects trophic level while the carbon ratio varies along a gradient from marine, coastal to terrestrial habitats (Hobson & Wassenaar 1999). For our study area, the isotopic analysis of

Figure 1. Study region and location of the seven Herring Gull breeding colonies across southwestern Scotland and Northern Ireland: 1 Islay, 2 Oronsay, 3 Pladda, 4 Lady Isle, 5 Portpatrick, 6 Copeland Islands, 7 Strangford.
these two elements captured the continuum from marine to terrestrial resource use by gulls (O’Hanlon et al. 2017). Although down feathers could only be obtained from two broods at Portpatrick (Fig. 1), we are confident that these feathers provide a representative reflection of resource use at this colony, as pellets and stable isotope analyses of feathers from older chicks from this colony indicated a similar diet (O’Hanlon 2016).

Table 1. Information on the land use around the Herring Gull study colonies across southwest Scotland and Northern Ireland (Fig. 1), sample sizes of eggs and clutches photographed, and broods from which down feathers were collected for stable isotope analyses.

| Colony     | Dates egg photographs were taken | Latitude leq  (°N) | Longitude leq (°W) | Amount of built-up area within 50 km of the colony (km²) and nearest distance (km) in parentheses | Average wave fetch within 50 km of the colony (km) | Colony size | Number of eggs (clutches) | Number of broods down feathers were sampled | Dates down feathers were sampled |
|------------|---------------------------------|-------------------|-------------------|-------------------------------------------------|-----------------------------------------------|------------|--------------------------|------------------------------------------|-------------------------------|
| 1. Islay   | 18 May 2014                     | 55.8              | 6.48              | 1 (12)                                          | 1044                                          | 25         | 24 (9)                   | 11                                       | 27 May 2014                   |
| 2. Oronsay | 3 June 2014                     | 54.43             | 5.12              | 2 (28)                                          | 743                                           | 95         | 75 (31)                  | 33                                       | 6 June 2014, 17 June 2014, 2 July 2014 |
| 3. Pladda  | 31 May 2014                     | 54.84             | 5.13              | 89 (29)                                         | 840                                           | 150        | 38 (15)                  | 24                                       | 20 June 2014, 28 June 2014    |
| 4. Lady Isle | 13 May 2014                 | 55.53             | 4.73              | 473 (3.5)                                       | 957                                           | 830        | 72 (24)                  | 28                                       | 30 May 2014, 11 June 2014, 18 June 2014 |
| 5. Portpatrick | 5 May 2014                  | 54.43             | 5.57              | 222 (8)                                         | 1243                                          | 175        | 58 (28)                  | 2                                        | 9 June 2014                   |
| 6. Copeland | 11 May 2014 & 23 May 2014      | 54.69             | 5.52              | 405 (3.5)                                       | 993                                           | 683        | 141 (48)                 | 22                                       | 15 June 2014, 29 June 2014    |
| 7. Strangford | 14 May 2014                  | 54.40             | 5.61              | 406 (2)                                         | 1004                                          | 190        | 106 (39)                 | 13                                       | 11 June 2014                  |
| Total      | –                               | –                 | –                 | –                                               | –                                             | –          | –                        | –                                        | –                             |

*Wave fetch is calculated based on the exposure of a coastline depending on its topography (Burrows et al. 2008). A low (or short) wave fetch value reflects a more sheltered intertidal habitat, with short distances to the nearest land mass, whereas high (or long) wave fetch values reflect an exposed coastline, with greater distances to the nearest land mass. Rocky shorelines with low wave fetch support a greater abundance and diversity of potential intertidal prey species (Burrows 2012). bColony size is the count of apparently occupied nests (AON) in 2012 or 2013 using standard monitoring techniques (Walsh et al. 1995). |

Statistical analyses

We performed all statistical analyses in R, Version 2.12.0 (R Development Core Team 2014). For each egg trait (eggshell colour, maculation, shape and size) we used the mean value of all eggs measured in a clutch as the unit of analysis, as the exact order of laying was unknown. To determine whether the egg traits were related to one another, we performed linear mixed effect models using lme4 with one egg trait as the response variable and the other egg traits as the explanatory variables, so that all combinations were tested. To account for the data being representative of a population, these models included colony as a random effect and included only clutches where all four egg traits were measured (n = 128 clutches). As our visits were not standardized to the phenology of each colony, we performed Spearman rank correlation tests between egg sampling dates and each egg trait, and chick down sampling dates and resource assimilation.

Resource assimilation of a colony was characterized by the first principal component of the stable isotope ratios δ13C and δ15N from chick down feathers collected in that colony (isotope-PC1) using the R package prcomp. Isotope-PC1 explained 84.6% of the variation in isotope ratios, with an eigenvalue of 7.13 and factor loadings for δ13C = 0.90 and δ15N = 0.44. Hence, isotope-PC1 reflected where on the marine–terrestrial gradient birds from a colony foraged, with low isotope-PC1 values when a terrestrial diet was consumed and high values for marine diets, and also reflected trophic level, which increased from terrestrial to marine diets (Fig. S5).

To explore the effect of variation in assimilated resources (isotope-PC1) prior to egg-laying on...
among-colony differences in egg traits we used linear mixed effect models. We ran separate models for each egg trait, with the mean egg trait of each clutch as the response variable. Mean isotope-PC1, colony size and clutch size were included as explanatory variables. We included colony size, as the study colonies varied in size from 25 and 830 apparently occupied nests (Table 1). Larger colonies can exert increased levels of competition and deplete local resources (Furness & Birkhead 1984, Birt et al. 1987, Lewis et al. 2001), which could affect egg traits. We included clutch size as the number of eggs in each clutch varied between one and three, which might also influence the egg trait measurements (see above). The variable isotope-PC1 only varied among colonies, so the power of our mixed model to find an effect of assimilated diet on egg traits is reduced, and hence we applied a conservative test. Colony identity was again included as a random effect to account for the non-independence of clutches from the same colony. We tested the main effects (mean isotope-PC1, colony size and clutch size) and all two-way interactions between them using an information theoretic approach (Burnham & Anderson 2002) to determine the best-fitting model across all competing models. Akaike’s information criterion (AIC_c), corrected for small sample sizes, and Akaike weights (\( w_i \)) were calculated for all models in the R package MuMIn (Barton 2012) and compared across candidate models. To account for model uncertainty, we retained all competing models with \( \Delta AIC_c \leq 2 \) compared with the model with the lowest AIC_c (Burnham & Anderson 2002). We calculated the weighted average of each supported explanatory variable across all retained models to obtain parameter estimates and 95% confidence intervals. The fact that the 95% confidence intervals did not overlap with zero supports that variable is important.

We checked all diagnostic plots to ensure model assumptions were met; colony size had to be log-transformed. We visually inspected all relationships for linearity and, where plots suggested a non-linear relationship, we also tested the fit of a second-order polynomial function. The significance threshold was set at a two-tailed \( P < 0.05 \).

**RESULTS**

All four egg traits (eggshell colour, maculation, shape and size) varied independently of each other with no significant relationships among any of the traits across colonies (linear mixed effect models, all \( P > 0.10, n = 128 \)). We found no seasonal change in mean egg traits (\( P > 0.62 \)) or assimilated diet (\( r_s = 0.003, n = 132, P = 0.98 \)) among colonies. Female Herring Gulls’ assimilated diet during the pre-laying period, reflected in chick’s down feather isotope_PC1, varied significantly across the seven colonies (\( F_{6,126} = 15.73, \ P < 0.001, R^2 = 0.40 \)). Variation in assimilated diet among colonies during egg formation was important in explaining variation in Herring Gull egg size and maculation, with 95% confidence intervals that did not overlap zero (Tables 2 and 3). In colonies with the most terrestrial, low trophic level diets (low values of \( ^{13}C \) and \( ^{15}N \) contribute to a negative isotope_PC1 value representing grain, terrestrial invertebrates and some human refuge; Islay and Portpatrick) or the most marine, high trophic level diets (high values of \( ^{13}C \) and \( ^{15}N \) contribute to a positive isotope_PC1 representing intertidal invertebrates, fish and occasional mammal and bird prey; Oronsay and Pladda) eggs were larger than in the remaining colonies which had an intermediate diet (Table 3, Fig. 2a). Furthermore, egg size declined with increasing colony size (Table 3, Fig. S6a). In colonies with a more marine, higher trophic level diet, eggs were also more maculated (Table 3, Fig. 2b). We found no relationship between isotope_PC1 and eggshell colour or egg shape; however, eggs in larger colonies were more pointed (Table 3, Fig. S6b). There was no support for clutch size explaining any of the observed variation in egg traits.

**DISCUSSION**

Colonies of Herring Gulls, generalist foragers inhabiting landscapes with a range of marine and terrestrial foraging opportunities, differed in their assimilated diet during egg formation. These differences explained a significant part of the among-colony variation in egg size and maculation, but not in eggshell colour or egg shape. Based on previous work in the study region, this variation would be expected to be related to local availability of foraging habitats (O’Hanlon et al. 2017). Our results thus indicate that egg traits of Herring Gulls are sensitive to variation in landscape structure reflecting relative local availabilities of marine and terrestrial resources.

In our study region, breeding Herring Gulls during incubation and chick-rearing generally
consumed the most abundant resources in the local landscape, which varied among colonies (O’Hanlon et al. 2017). Female Herring Gulls also probably consumed resources gathered locally for egg formation as they depend largely on exogenous resources for egg formation (income breeders, Kilpi et al. 1996). Although Herring Gulls can make regional movements in the non-breeding season (Coulson & Butterfield 1985, Camphuysen et al. 2011), they return to their breeding area 1–2 months prior to the start of egg-laying (N. J. O’Hanlon unpubl. data). Some stored lipid reserves can be used for egg formation (Hario et al. 1991) but it is not known when they are accumulated and whether they are differentially allocated among the eggs of a clutch. Furthermore, down feathers are thought to reflect the protein more than the lipid stores deposited into the egg (Hobson 2006). Consequently, the isotope values of chick down feathers are thought to provide an accurate reflection of the diet of females prior to egg-laying.

At the colony level, Herring Gulls foraging more on either marine or terrestrial resources laid

### Table 2. Rank of general linear multivariate regression models explaining variation in Herring Gull egg traits by isotope_PC1 (proxy for assimilated diet), colony size and clutch size using Akaike’s information criterion corrected for small sample size (AICc): k is the number of estimated parameters included, wi is the Akaike weight, and ΔAICc is the AICc difference. Supported models (ΔAICc < 2) are shown in bold and were used in the model averaging to obtain parameter estimates (Table 3).

| Variables included within model | k | AICc | ΔAICc | wi |
|--------------------------------|---|------|-------|----|
| (A) Eggshell colour            |   |      |       |    |
| Intercept only                 | 1 | −913.3 | 0.00  | 0.320 |
| Clutch Size                    | 2 | −912.6 | 0.70  | 0.225 |
| Colony Size                    | 2 | −911.3 | 1.98  | 0.119 |
| Isotope_PC1                    | 2 | −911.2 | 2.06  | 0.114 |
| Clutch Size, Isotope_PC1       | 3 | −910.5 | 2.78  | 0.080 |
| Clutch Size, Colony Size       | 3 | −910.4 | 2.86  | 0.077 |
| Colony Size, Isotope_PC1       | 3 | −909.1 | 4.12  | 0.041 |
| Clutch Size, Colony Size, Isotope_PC1 | 4 | −908.3 | 5.03  | 0.026 |
| (B) Maculation                 |   |      |       |    |
| Colony Size, Isotope_PC1       | 3 | −601.3 | 0.00  | 0.487 |
| Isotope_PC1                    | 2 | −599.9 | 1.44  | 0.238 |
| Clutch Size, Colony Size, Isotope_PC1 | 4 | −598.4 | 2.92  | 0.113 |
| Clutch Size, Isotope_PC1       | 3 | −598.2 | 3.13  | 0.102 |
| Clutch Size                    | 2 | −595.1 | 6.25  | 0.021 |
| Intercept only                 | 1 | −594.9 | 6.45  | 0.019 |
| Colony Size                    | 2 | −593.5 | 7.78  | 0.010 |
| Clutch Size, Colony Size       | 3 | −593.4 | 7.90  | 0.009 |
| (C) Shape                      |   |      |       |    |
| Colony Size                    | 2 | −254.8 | 0.00  | 0.432 |
| Colony Size, Isotope_PC1       | 3 | −253.6 | 1.24  | 0.232 |
| Clutch Size, Colony Size       | 3 | −253.0 | 1.86  | 0.171 |
| Clutch Size, Colony Size, Isotope_PC1 | 4 | −251.6 | 3.27  | 0.084 |
| Intercept only                 | 1 | −249.9 | 4.93  | 0.037 |
| Clutch Size                    | 2 | −249.2 | 5.59  | 0.026 |
| Isotope_PC1                    | 2 | −247.5 | 7.36  | 0.011 |
| Clutch Size, Isotope_PC1       | 3 | −246.6 | 8.18  | 0.007 |
| (D) Size                       |   |      |       |    |
| Clutch Size, Colony Size, Isotope_PC1² | 4 | 1234.3 | 0.00  | 0.474 |
| Colony Size, Isotope_PC1²      | 3 | 1234.8 | 0.46  | 0.377 |
| Colony Size                    | 2 | 1238.2 | 3.88  | 0.068 |
| Clutch Size, Colony Size       | 3 | 1238.9 | 4.60  | 0.048 |
| Isotope_PC1²                   | 2 | 1241.4 | 7.05  | 0.014 |
| Clutch Size, Isotope_PC1²      | 3 | 1241.6 | 7.23  | 0.013 |
| Intercept only                 | 1 | 1243.8 | 9.49  | 0.004 |
| Clutch Size                    | 2 | 1244.6 | 10.25 | 0.003 |

The superscript 2 refers to the quadratic term in the model as Isotope_PC1 was not linear so was fit to a second order polynomial. Isotope_PC1 without the subscript in the linear term.

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on average larger eggs than those with an intermediate diet. A positive relationship between egg size and food quantity and quality has been observed in several gull species (Hiom et al. 1991, Bolton et al. 1992, Blight 2011, Svagelj et al. 2015, Bennett et al. 2017). The observed association of assimilated diet with egg size may reflect differences in resource quality and availability (Sibly & McCleery 1983, Pierotti & Annett 1991). Foraging on marine resources at a higher trophic level, especially fish, is generally thought to provide more energy and protein than foraging on lower trophic and terrestrial items (Hiom et al. 1991, Pierotti & Annett 1991, Ronconi et al. 2014, Blight et al. 2015). For example, among-colony differences in egg size in Balearic Shearwaters Puffinus mauretanicus over a relatively small spatial scale were attributed to differences in trophic level (Louzao et al. 2008). In our study, Herring Gulls at colonies where they consumed more marine items (Oronsay, Strangford) laid the largest eggs, although the Gulls only rarely consumed fish (O’Hanlon et al. 2017). More marine items were consumed in colonies along sheltered coastlines (O’Hanlon et al. 2017) harbouring a higher abundance and diversity of marine invertebrates (Burrows 2012). In contrast, terrestrial and low trophic level resources have been considered a lower quality diet for gulls (Murphy et al. 1984, Pierotti & Annett 1991, Pons 1992, Belant et al. 1993, Duhem et al. 2008, Weiser & Powell 2010, Hobson et al. 2015, O’Hanlon et al. 2017), although some resources, for example refuse from landfills, can have a high energy and protein content (Sibly & McCleery 1983, Pierotti & Annett 1987, 1991, Pons & Migot 1995). In our study region, grain was the main terrestrial food (O’Hanlon et al. 2017), and its source (farmland) was the dominant terrestrial habitat (86.2–99.7% of all potential terrestrial foraging habitat), close to all study colonies. However, female Gulls in colonies with a mainly terrestrial diet (Islay, Portpatrick) also laid large eggs, possibly because terrestrial resources are thought to be more predictably available compared with intertidal resources that may only be available at low tide (Van Donk et al. 2017), which may compensate for small differences in resource quality (Pierotti & Annett 1991, Steigerwald et al. 2015). Future

| Parameter | Effect | se | Lower | Upper | Relative importancea |
|-----------|--------|----|-------|-------|-----------------------|
| (A) Eggshell colour | | | | | |
| Intercept | 1.113 | | | | |
| Clutch Size | 0.001 | 0.002 | -0.002 | 0.008 | 0.408 |
| Colony Size | 0.0003 | 0.001 | -0.005 | 0.007 | 0.263 |
| (B) Maculation | | | | | |
| Intercept | 0.068 | | | | |
| Isotope_PC1 | -0.004 | 0.0012 | -0.007 | -0.002 | 0.940 |
| Colony Size | -0.003 | 0.0029 | -0.009 | 0.001 | 0.619 |
| (C) Shape | | | | | |
| Intercept | 0.3233 | | | | |
| Colony Size | 0.025 | 0.010 | 0.006 | 0.043 | 0.919 |
| Isotope_PC1 | 0.001 | 0.003 | -0.005 | 0.013 | 0.334 |
| Clutch Size | 0.001 | 0.007 | -0.021 | 0.035 | 0.288 |
| (D) Size | | | | | |
| Intercept | 84.388 | | | | |
| Colony Size | -2.068 | 0.533 | -3.119 | -1.018 | 0.967 |
| Isotope_PC1 | 16.115 | 6.301 | 3.684 | 28.539 | |
| Isotope_PC12 | 20.864 | 6.548 | 7.946 | 33.782 | 0.878 |
| Clutch Size | -0.594 | 0.723 | -2.367 | 0.232 | 0.538 |

aRelative importance refers to the sum of all model weights in which a given parameter occurs. The superscript 2 refers to the quadratic term in the model as Isotope_PC1 was not linear so was fit to a second order polynomial. Isotope_PC1 without the subscript in the linear term.
work should test whether terrestrial resources are relatively more predictable and abundant than intertidal food resources in our region. Intermediate diets probably represent a mixed diet of terrestrial and marine food consumed by an individual. As specialized behavioural, physiological or morphological adaptations are often required to obtain or digest particular food items, these individuals may be less efficient (Bolnick et al. 2003). If Herring Gulls show such specializations, this could result in smaller eggs laid by individuals consuming a mixed diet. Thus, it may be advantageous to Herring Gulls to select a specific resource, either because they exploit the most abundant or profitable resource or because they become efficient in using that resource.

Several studies have investigated the association between egg pigmentation (eggshell colour and maculation) and female condition or environmental conditions experienced during egg formation, but the relationship is not clear (Moreno & Osorno 2003, Osorno et al. 2004, Siefferman et al. 2006, Krist & Grim 2007, Martinez-de la Puente et al. 2007, Morales et al. 2008, Soler et al. 2008, Sanz & García-Navas 2009, López de Hierro & De Neve 2010, De Coster et al. 2013, Duval et al. 2013, Hargitai et al. 2016, 2018). Different studies have measured different components of eggshell pigmentation using different approaches, and it is likely that each measurement provides different information (Badás et al. 2017). Background eggshell colour did not vary with consumed diet in our study. This is consistent with experimental food restriction in Japanese Quails Coturnix coturnix japonica not affecting eggshell colour, although it did increase the amount of pigment in the shell (Duval et al. 2013). Background colour of Herring Gull eggs has been shown to vary with the egg’s contamination load (Hanley & Doucet 2012). The results from our analysis of eggshell colour cannot support any differences in food limitation between colonies or potential systematically different contamination loads associated with different diets. We also measured maculation (size and intensity of spots) with less maculated eggs (fewer and lighter spots) laid in colonies with a low trophic level, terrestrial diet. Previous studies found that less maculated eggs were associated with poorer foraging and female condition (Christians 2002, Reynolds et al. 2009, Duval et al. 2014), whereas
Martinez-de la Puente et al. (2007) and Hargitai et al. (2016) found the opposite association. The relationships between diet, female condition and foraging conditions in Herring Gulls need further study before a clear conclusion can be drawn. Including a measure of the distribution of spots, which also can be associated with female condition (Sanz & García-Navas 2009), may provide further information that we missed by only looking at size and intensity of spots.

Interestingly, female Herring Gulls in colonies consuming the most terrestrial diet, largely composed of grain (Islay, Portpatrick) and in colonies consuming the most marine diet (mostly intertidal invertebrates) laid similarly large eggs, but differed in maculation. As maculation possibly reflects differences in egg or parental quality as discussed above, the quality of Herring Gull eggs may vary with assimilated diet, independent of egg size. However, the existing information on the association between maculation and environmental and fitness-related traits is still unclear. Therefore, to elucidate further the observed maculation relationship, and test whether eggs formed on one diet are of higher quality than those formed on the other diet would require a study of hatching and fledgling success. Larger eggs can more probably result in surviving offspring (reviewed by Krist 2011), and egg pigmentation can play an important role in the successful development of the embryo (Maurer et al. 2011, Lahti & Ardia 2016). Unfortunately, due to the logistical constraints of studying seven colonies, we do not have measures of success for individual eggs or nests. Future studies should consider relationships between egg traits, diet and breeding success among nests within a colony. These logistical constraints also meant that we were unable to standardize our colony visits to specific times of breeding. However, we did not sample eggs or down feathers consistently early or late, and egg traits and diet were not related to sampling date. Therefore, differences in breeding phenology related to parental quality are unlikely to have influenced the relationships we found among egg traits and Herring Gull diet.

We found that egg size and shape, but not pigmentation, were associated with colony size. In larger colonies, Herring Gulls laid smaller eggs (also observed by Coulson et al. 1982, Bennett et al. 2017) and more pointed eggs. Individuals nesting in larger colonies may experience increased competition for limited resources (Ashmole 1963, Lewis et al. 2001, Wakefield et al. 2013), resulting in smaller and more pointed eggs (Houston et al. 1983, Kilpi et al. 1996, Nager et al. 2000). If the larger colonies are expanding, there may be more young, inexperienced birds that lay smaller eggs (Coulson et al. 1982), whereas in declining colonies larger eggs may be attributed to reduced resource depletion or disproportionately more older individuals breeding (Bennett et al. 2017). In this study region, colony size was unrelated to population trends (O’Hanlon & Nager 2018), which makes it unlikely that the colonies differed in age structure; instead, female Gulls in larger colonies are likely to have experienced greater resource competition.

CONCLUSION

At the colony level, Herring Gull egg traits measured by digital photography in the field, which was an efficient tool to measure multiple egg traits with minimal disturbance to the colony, were related to differences in resources consumed when eggs were formed. As Herring Gulls, at least in our study region, used the resources from habitats that were most readily available to them (O’Hanlon et al. 2017), differences in resource use during egg formation among colonies with different composition of habitats in their landscape also affected egg traits. Anthropogenic changes in local landscapes that alter the resources available to generalist birds such as Herring Gulls may therefore affect reproduction from the earliest stage in the breeding cycle.

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AUTHOR CONTRIBUTIONS

R.G.N. and N.J.O’H. originally formulated the idea and developed the methodology. N.J.O’H., J.A.O.M. and S.A. conducted fieldwork and
extracted data from the images. R.A.R.M. carried out the stable isotope analysis. N.J.O'H. analysed the data. N.J.O'H. and R.G.N. wrote the manuscript, with all authors providing editorial advice.

REFERENCES

Alverson, D.L. 1992. Commercial fisheries and the Steller Sea Lion (Eumetopias jubatus): the conflict arena. Rev. Aquat. Sci. 6: 203–256.

Amar, A. & Redpath, S.M. 2005. Habitat use by Hen Harriers Circus cyaneus on Orkney: implications of land-use change for this declining population. Ibis 147: 37–47.

Ardendt, W.J. 2004. A quarter century of variation in color and allometric characteristics of eggs from a rain forest population of the Pearly-eyed Thrasher (Margarops fuscatus). Caribb. J. Sci. 40: 204–217.

Ashmole, N.P. 1963. The regulation of numbers of tropical oceanic birds. Ibis 103: 458–473.

Avilés, J.M., Stokke, B.G., Moksnes, A., Roskaft, E. & Møller, A.P. 2006. Environmental conditions influence egg color of Reed Warblers Acrocephalus scirpaceus and their parasite, the Common Cuckoo Cuculus canorus. Behav. Ecol. Sociobiol. 61: 475–485.

Badás, E.P., Martinez, J., Rivero-de Aguilar, J., Stevens, M., van der Velde, M., Komdeur, J. & Merino, S. 2017. Eggshell pigmentation in the Blue Tit: male quality matters. Behav. Ecol. Sociobiol. 71: 57.

Barrett, R.T., Nilsen, E.B. & Anker-Nilsen, T. 2012. Long-term decline in egg size of Atlantic Puffins Fratercula arctica is related to changes in forage fish stocks and climate conditions. Mar. Ecol. Prog. Ser. 457: 1–10.

Barton, K. 2012. MuMIn: Multi-Model Inference. R package version 1.13.4.

Belant, J.L., Seaman, T.W., Gabrey, S.W. & Ickes, S.K. 1993. Importance of landfills to nesting Herring Gulls. Condor 95: 817–830.

Bennett, J.L., Jamieson, E.G., Ronconi, R.A. & Wong, S.N.P. 2017. Variability in egg size and population declines of Herring Gulls in relation to fisheries and climate conditions. Avian Conserv. Ecol. 12: 16.

Benton, T.G., Vickery, J.A. & Wilson, J.D. 2003. Farmland biodiversity: Is habitat heterogeneity the key?. Trends Ecol. Evol. 18: 182–188.

Birt, V.L., Birt, T.P., Goulet, D., Cairns, D.K. & Montevecchi, W.A. 1987. Ashmole’s halo: direct evidence for prey depletion by a seabird. Mar. Ecol. Prog. Ser. 40: 205–208.

Bligh, L.K. 2011. Egg production in a coastal seabird, the Glaucous-winged Gull (Larus glaucescens), declines during the last century. PLoS One 6: e22027.

Blight, L.K., Drevier, M.C. & Arcese, P. 2015. A century of change in Glaucous-winged Gull (Larus glaucescens) populations in a dynamic coastal environment. Condor 117: 108–120.

Bolnick, D.J., Svanbäck, R., Fordyce, J.A., Yang, L.H., Davis, J.M., Hulse, C.D. & Forister, M.L. 2003. The ecology of individuals: incidence and implications of individual specialization. Am. Nat. 161: 1–28.

Bolton, M., Houston, D. & Monaghan, P. 1992. Nutritional constraints on egg formation in the lesser black-backed gull: an experimental study. J. Anim. Ecol. 61: 521–532.

Bond, G., Burnside, N.G., Metcalfe, D.J., Scott, D.M. & Blamire, J. 2004. The effects of land-use and landscape structure on Barn Owl (Tyto alba) breeding success in southern England, U.K. Landsc. Ecol. 20: 555–566.

Brulez, K., Miksis, I., Cooney, C.R., Hauber, M.E., Lovell, P.G., Maurer, G., Portugal, S.J., Russell, D., Reynolds, S.J. & Cassey, P. 2016. Eggshell pigment composition covaries with phylogeny but not with life history or with nesting ecology traits of British passerines. Ecol. Evol. 6: 1637–1645.

Burnham, K.P. & Anderson, D.R. 2002. Model Selection and Multimodel Inference. New York, NY: Springer.

Burrows, M.T. 2012. Influences of wave fetch, tidal flow and ocean colour on subtidal rocky communities. Mar. Ecol. Prog. Ser. 445: 193–207.

Burrows, M.T., Harvey, R. & Robb, L. 2008. Wave exposure indices from digital coastlines and the prediction of rocky shore community structure. Mar. Ecol. Prog. Ser. 353: 1–12.

Camphuysen, C.J., Verhulst, H.J.P. & Spans, A.L. 2011. Colony- and age-specific seasonal dispersal of Herring Gulls Larus argentatus breeding in The Netherlands. J. Ornithol. 152: 849–868.

Chace, J.F. & Walsh, J.J. 2006. Urban effects on native avifauna: a review. Landsc. Urban Plan. 74: 46–69.

Christians, J.K. 2002. Avian egg size: variation within species and inflexibility within individuals. Biol. Rev. Camb. Philos. Soc. 77: 1–26.

Coulson, J.C. & Butterfield, J. 1985. Movements of British Herring Gulls. Bird Study 32: 91–103.

Coulson, J.C. & Coulson, B.A. 2008. Lesser Black-backed Gulls Larus fuscus nesting in an inland urban colony: the importance of earthworms (Lumbricidae) in their diet. Bird Study 55: 37–41.

Coulson, J.C., Duncan, N. & Thomas, C. 1982. Changes in the breeding biology of the Herring Gull (Larus argentatus) induced by reduction in the size and density of the colony. J. Anim. Ecol. 51: 739–756.

Davoren, G.K. & Montevecchi, W.A. 2003. Signals from seabirds indicate changing biology of capelin stocks. Mar. Ecol. Prog. Ser. 258: 253–261.

De Coster, G., De Neve, L. & Lens, L. 2013. Intra-clutch variation in avian eggshell pigmentation covaries with female quality. J. Ornithol. 154: 1057–1065.

Duhem, C., Roche, P., Vidal, E. & Tatoni, T. 2008. Effects of anthropogenic food resources on Yellow-legged Gull colony size on Mediterranean islands. Popul. Ecol. 50: 91–100.

Duval, C., Cassey, P., Miksis, I., Reynolds, S.J. & Spence, K.A. 2013. Condition-dependent strategies of eggshell pigmentation: an experimental study of Japanese Quail (Coturnix coturnix japonica). J. Exp. Biol. 216: 700–708.

Duval, C., Zimmer, C., Miksis, I., Cassey, P. & Spence, K.A. 2014. Early life stress shapes female reproductive strategy through eggshell pigmentation in Japanese Quail. Gen. Comp. Endocrinol. 208: 146–153.
Furness, R.W. & Birkehead, T.R. 1984. Seabird colony distributions suggest competition for food supplies during the breeding season. *Nature* 311: 655–656.

Garthe, S., Schwemmer, P., Paiva, V.H., Corman, A., Fock, H.O., Voigt, C.C. & Adler, S. 2016. Terrestrial and marine foraging strategies of an opportunistic seabird species breeding in the Wadden Sea. *PLoS One* 11: e0159630.

Gómez, J. & Liñán-Cembrano, G. 2016. SpotEgg: an image-processing tool for automated analysis of colouration and spotiness. *J. Avian Biol.* 48: 502–512.

Gorski, A., Nowakowski, J.J. & Banbura, J. 2015. Variation in egg-size traits of the European Roller (Coracias garrulus) in eastern Poland. *Ornis Fenn.* 92: 213–220.

Gosler, A.G., Barnett, P.R. & Reynolds, S.J. 2000. Inheritance and variation in eggshell patterning in the Great Tit Parus major. *Proc. R. Soc. Lond. B* 267: 2469–2473.

Gosler, A.G., Higham, J.P. & Reynolds, S.J. 2005. Why are birds’ eggs speckled? *Ecol. Lett.* 8: 1105–1113.

Götmark, F. 1984. Food and foraging in five European Larus gulls in the breeding season: a comparative review. *Ornis Fenn.* 61: 9–18.

Grémillet, D., Picchegru, L., Kuntz, G., Woakes, A.G., Wilkinson, S., Crawford, R.J.M. & Ryan, P.G. 2008. A junk-food hypothesis for gannets feeding on fishery waste. *Proc. R. Soc. Lond. B* 275: 1149–1156.

Gyimesi, A., Boudewijn, T.J., Buijs, R.J., Shamoun-Hobson, K.A. & Wassenaar, L.I. 2000. Using stable isotopes to quantitatively track endogenous and exogenous nutrient allocations to eggs of birds that travel to breed. *Ardea* 94: 359–369.

Hobson, K.A. & Wassenaar, L.I. 1999. Stable isotope ecology: an introduction. *Oecologia* 120: 312–313.

Hobson, K.A., Blight, L.K. & Arcese, P. 2015. Human-induced long-term shifts in gull diet from marine to terrestrial sources in North America’s coastal Pacific: more evidence from more isotopes (15N, 13C). *Environ. Sci. Technol.* 49: 10834–10840.

Houston, D.C., Jones, P.J. & Sibly, R.M. 1983. The effect of female body condition on egg laying in Lesser Black-backed gulls Larus fuscus. *J. Zool. Soc.* 200: 509–520.

Hoyt, D.F. 1976. The effect of shape on the surface-volume relationships of birds’ eggs. *Condor* 78: 343–349.

Hunt, G.L. 1972. Influence of food distribution and human disturbance on the reproductive success of Herring Gulls. *Ecology* 53: 1051–1061.

Kadlec, J.A., William, H., Drury, W.J.H. & Onion, D.K. 1969. Growth and mortality of Herring Gull chicks. *Bird-Banding* 40: 222–233.

Kilpi, M., Hillstrom, L. & Lindstrom, K. 1996. Egg-size variation and reproductive success in the Herring Gull Larus argentatus: adaptive or constrained size of the last egg? *Ibis* 138: 212–217.

Klaassen, M., Baarspul, T., Dekkers, T. & van Tienen, P. 2004. The relationship between carbon stable isotope ratios of, hatching down and egg yolk in Black-headed Gulls. *J. Field Ornithol.* 75: 196–199.

Kouwenberg, A., Hipfner, J.M., Kay, D.W.M.C. & Storey, A.E. 2013. Corticosterone and stable isotopes in feathers predict egg size in Atlantic Puffins Fratercula arctica. *Ibis* 155: 413–418.

Krist, M. 2011. Egg size and offspring quality: a meta-analysis in birds. *Biol. Rev. Camb. Philos. Soc.* 86: 692–716.

Krist, M. & Grim, T. 2007. Are blue eggs a sexually selected signal of female Collared Flycatchers? A cross-fostering experiment. *Behav. Ecol. Sociobiol.* 61: 863–876.

Kurki, S., Nikula, A., Helle, P. & Lindon, H. 2000. Landscape fragmentation and forest composition effects on grous breeding success in boreal forests. *Ecology* 81: 1985–1997.

Lahti, D.C. & Ardia, D.R. 2016. Shedding light on bird egg color: pigment as parasol and the dark car effect. *Am. Nat.* 187: 547–563.

Lévesque, S., Sherratt, T.N., Hamer, K.C. & Wanless, S. 2001. Evidence of intra-specific competition for food in a pelagic seabird. *Nature* 412: 816–819.

Lindstrom, A., Visser, G.H. & Daan, S. 1993. The energetic cost of feather synthesis to basal metabolic rate is proportional. *Physiol. Zool.* 66: 490–510.

López de Hierro, M.D.G. & De Neve, L. 2010. Pigment limitation and female reproductive characteristics influence eggshell thickness and ground colour variation in the House Sparrow (Passer domesticus). *J. Ornithol.* 151: 833–840.

Louzao, M., Igual, J.M., Genovart, M., Forero, M.G., Hobson, K.A. & Oro, D. 2008. Spatial variation in egg size of a top predator: interplay of body size and environmental factors? *Acta Oecol.* 34: 186–193.

Martinez-de la Puente, J., Merino, S., Moreno, J., Moreno, J., Tomas, G., Morales, J., Lobato, E., Garcia-Fraile, S. & Martinez, J. 2007. Are eggshell spottiness and colour indicators of health and condition in Blue Tits Cyanistes caeruleus? *J. Avian Biol.* 38: 377–384.

Maurer, G., Portugal, S.J., Miksiński, I. & Cassey, P. 2011. Speckles of cryptic black-backed gull eggs show no
机械或流体结构的功能。J. Zool. 285: 194–204.

Millon, A. & Bretagnolle, V. 2008. Predator population dynamics under a cyclic prey regime: numerical responses, demographic parameters and growth rates. Oikos 117: 1500–1510.

Monaghan, P. & Nager, R.G. 1997. Why don’t birds lay more eggs?. Trends Ecol. Evol. 12: 270–274.

Montevecchi, W.A. 1993. Birds as indicators of change in marine prey stocks. In Furness, R.W. & Greenwood, J.J.J. (eds) Birds as Monitors of Environmental Change: 217–266. London: Chapman & Hall.

Morales, J., Velando, A. & Moreno, J. 2008. Pigment allocation to eggs decreases plasma antioxidants in a songbird. Behav. Ecol. Sociobiol. 63: 227–233.

Moreno, J. & Osorno, J.L. 2003. Avian egg colour and sexual selection: does eggshell pigmentation reflect female condition and genetic quality? Ecol. Lett. 6: 803–806.

Murphy, E.C., Day, R.H., Oakley, K.L. & Hoover, A.A. 1984. Dietary changes and poor reproductive performance in Glaucous-winged Gulls. Auk 101: 532–541.

Nager, R.G. 2006. The challenges of making eggs. Ardea 94: 323–346.

Nager, R.G., Monaghan, P. & Houston, D.C. 2000. Within-clutch trade-offs between the number and quality of eggs: Experimental manipulations in gulls. Ecology 81: 1339–1350.

O’Hanlon, N.J. 2016. Spatial variation in herring gull traits and their potential as monitors of the coastal environment. PhD Thesis, University of Glasgow.

O’Hanlon, N.J. & Nager, R.G. 2018. Identifying habitat-driven spatial variation in colony size of Herring Gulls Larus argentatus. Bird Study 65: 306–316.

O’Hanlon, N.J., McGill, R.A.R. & Nager, R.G. 2017. Increased use of marine resources benefits breeding success in a generalist gull species. Mar. Ecol. Prog. Ser. 574: 193–210.

Osorno, L., Morales, J., Merino, S., Toma, G. & Moreno, J. 2004. Egg colouration and male parental effort in the Pied Flycatcher Ficedula hypoleuca. J. Avian Biol. 35: 300–304.

Osterblom, H., Olsson, O., Blenczner, T. & Furness, R.W. 2008. Junk-food in marine ecosystems. Oikos 117: 967–977.

Palma, L., Beja, P., Pais, M. & Cancela Da Fonseca, L. 2006. Why do raptores take domestic prey? The case of Bonell’s Eagles and pigeons. J. Appl. Ecol. 43: 1075–1086.

Penteriani, V., Gallardo, M. & Roche, P. 2002. Landscape structure and food supply affect Eagle Owl (Bubo bubo) density and breeding performance: a case of intra-population heterogeneity. J. Zool. 257: 365–372.

Perrins, C.M. 1996. Eggs, egg formation and the timing of breeding. Ibis 138: 2–15.

Pierotti, R. 1982. Habitat selection and its effect on reproductive output in the Herring Gull in Newfoundland. Ecology 63: 854–868.

Pierotti, R. & Annett, C. 1987. Reproductive consequences of dietary specialization and switching in an ecological generalist. In Kamil, A.C., Krebs, J.R. & Pulliam, H.R. (eds) Foraging Behavior: 417–442. New York, NY: Plenum Press.

Pierotti, R. & Annett, C.A. 1991. Diet choice in the Herring Gull: constraints imposed by reproductive and ecological factors. Ecology 72: 319–328.

Pons, M. 1992. Effects of change in the availability of human refuse on breeding parameters in a Herring Gull Larus argentatus population in Brittany, France. Ardea 80: 143–150.

Pons, J.M. & Migot, P. 1995. Life-history strategy of the Herring Gull: changes in survival and fecundity in a population subjected to various feeding conditions. J. Anim. Ecol. 64: 592–599.

Pyke, G.H., Pulliam, H.R. & Charnov, E.L. 1977. Optimal foraging: a selective review of theory and tests. Q. Rev. Biol. 52: 137–154.

R Development Core Team 2014. R: A Language and Environment for Statistical Computing. Vienna: R Foundation for Statistical Computing.

Reynolds, S.J., Martin, G.R. & Cassey, P. 2009. Is sexual selection blurring the functional significance of eggshell coloration hypotheses? Anim. Behav. 78: 209–215.

Robbins, C.T. 1981. Estimation of the relative protein cost of reproduction in birds. Condor 83: 177–179.

Rock, P. 2005. Urban gulls: problems and solutions. Br. Birds 98: 338–355.

Ronconi, R.A., Steenweg, R.J., Taylor, P.D. & Mallory, M.L. 2014. Gull diets reveal dietary partitioning, influences of isotopic signatures on body condition, and ecosystem changes at a remote colony. Mar. Ecol. Prog. Ser. 514: 247–261.

Sanz, J.J. & Garcia-Navas, V. 2009. Eggshell pigmentation pattern in relation to breeding performance of Blue Tits Cyanistes caeruleus. J. Anim. Ecol. 78: 31–41.

Schneider, C.A., Rasband, W.S. & Elceiir, K.W. 2012. NIH Image to ImageJ: 25 years of image analysis. Nat. Methods 9: 671–675.

Schoener, T.W. 1971. Theory of feeding strategies. Annu. Rev. Ecol. Syst. 2: 369–404.

Schwemmer, P., Garthe, S. & Mundry, R. 2008. Area utilization of gulls in a coastal farmland landscape: habitat mosaic supports niche segregation of opportunistic species. Landsc. Ecol. 23: 355–367.

Sibly, R.M. & McCleery, R.H. 1983. Increase in weight of Herring Gulls while feeding. J. Anim. Ecol. 52: 35–50.

Siefferman, L., Navara, K.J. & Hill, G.E. 2006. Egg coloration is correlated with female condition in Eastern Bluebirds (Sialia sialis). Behav. Ecol. Sociobiol. 59: 651–656.

Soler, J.J., Navarro, C., Contreras, T.P., Avilés, J.M. & Cuervo, J.J. 2008. Sexually selected egg coloration in Spotless Starlings. Am. Nat. 171: 183–194.

Sorensen, M.C., Hipfner, J.M., Kyser, T.K. & Norris, D.R. 2009. Carry-over effects in a Pacific seabird: stable isotope evidence that pre-breeding diet quality influences reproductive success. J. Anim. Ecol. 78: 460–467.

Steigerwald, E.C., Igual, J.M., Payo-Payo, A. & Tavecchia, G. 2015. Effects of decreased anthropogenic food availability on an opportunistic gull: evidence for a size-mediated response in breeding females. Ibis 157: 439–448.

Stephens, D. & Krebs, J. 1986. Foraging Theory. Princeton, NJ: Princeton University Press.

Stoddard, M.C., Kilner, R.M. & Town, C. 2014. Pattern recognition algorithm reveals how birds evolve individual egg pattern signatures. Nat. Commun. 5: 1–10.

SvagjeI, W.S., Linsner, N., Borboroglu, P.G. & Yorio, P. 2015. Variation in the size of eggs of Kelp Gulls (Larus dominicanus) at two colonies in Patagonia, Argentina. Waterbirds 38: 92–98.
Troscianko, J. 2014. A simple tool for calculating egg shape, volume and surface area from digital images. *Ibis* 156: 874–878.

Van Donk, S., Camphuysen, C.J., Shamoun-Baranes, J. & Van Der Meer, J. 2017. The most common diet results in low reproduction in a generalist seabird. *Ecol. Evol.* 7: 4620–4629.

Vucetich, J.A. & Peterson, R.O. 2004. The influence of prey consumption and demographic stochasticity on population growth rate of Isle Royale wolves *Canis lupus*. *Oikos* 107: 309–320.

Wakefield, E.D., Bodey, T.W., Bearhop, S., Blackburn, J., Colhoun, K., Davies, R., Dwyer, R.G., Green, J.A., Grémillet, D., Jackson, A.L., Jessopp, M.J., Kane, A., Langston, R.H.W., Lescroel, A., Murray, S., Le Nuz, M., Patrick, S.C., Péron, C., Soanes, L.M., Wanless, S., Votier, S.C. & Hamer, K.C. 2013. Space partitioning without territoriality in Gannets. *Science* 341: 68–70.

Walsh, P.M., Halley, D.J., Harris, M.P., Nevo, A., Sim, I.M.W. & Tasker, M.L. 1995. *Seabird Monitoring Handbook for Britain and Ireland*. Peterborough: JNCC/RSPB/ITE/Seabird Group.

Wegmann, M., Vallat-Michel, A. & Richner, H. 2015. An evaluation of different methods for assessing eggshell pigmentation and pigment concentration using Great Tit eggs. *J. Avian Biol.* 46: 597–607.

Weiser, E.L. & Powell, A.N. 2010. Does garbage in the diet improve reproductive output of Glaucous Gulls? *Condor* 112: 530–538.

White, T.C.R. 2008. The role of food, weather and climate in limiting the abundance of animals. *Biol. Rev.* 83: 227–248.

Wikelski, M. & Tarlow, E.M. 2003. Costs of migration in free-flying songbirds. *Nature* 423: 704.

Yohannes, E., Gwinner, H., Lee, R.W. & Schwabl, H. 2016. Stable isotopes predict reproductive performance of European starlings breeding in anthropogenic environments. *Ecosphere* 7: e01566.

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**SUPPORTING INFORMATION**

Additional supporting information may be found online in the Supporting Information section at the end of the article.

**Appendix S1.** Photographic determination of egg size, shape, colour and maculation.

**Figure S1.** Egg holder to photograph Herring Gull eggs in the field with the graph paper background and QPcard+201 colour checker panel and scale. Grey squares P03–P09 highlighted on row 1 and colour square P15 highlighted on row 2.

**Figure S2.** Contrasting egg shapes with (a) low egg shape deviation scores describing oval eggs and (b) high deviation scores describing pointy eggs.

**Figure S3.** Contrasting egg colours with (a) low GB ratios reflecting more blue, less brown, eggs and (b) high GB ratios reflecting brown eggs.

**Figure S4.** Contrasting egg maculation with (a) low similarity scores reflecting highly maculated eggs and (b) high similarity scores reflecting less maculated eggs.

**Figure S5.** Relationship between mean (±sd) δ13C and δ15N values based on stable isotope analysis of Herring Gull down feather samples.

**Figure S6.** Relationship between Herring Gull mean (a) egg volume and (b) egg size (higher deviation scores relate to pointier eggs) with colony size (natural logarithm). Solid lines indicate the trend line with 95% confidence intervals (dashed lines) predicted from model averaged Mixed Effect Models, which also included isotope_PC1 and clutch size (Table 2). Shown are means ±1 se for each colony. Number refers to colony numbers as in Fig. 1.