

Feeding grounds for waders in the Bay of the Mont Saint-Michel (France): the Lanice conchilega reef serves as an oasis in the tidal flats

Bart de Smet, Laurent Godet, Jérôme Fournier, Nicolas Desroy, Mikaël Jaffré, Magda Vincx, Marijn Rabaut

▶ To cite this version:

Bart de Smet, Laurent Godet, Jérôme Fournier, Nicolas Desroy, Mikaël Jaffré, et al.. Feeding grounds for waders in the Bay of the Mont Saint-Michel (France): the Lanice conchilega reef serves as an oasis in the tidal flats. Marine Biology, 2013, 160, pp.751-761. 10.1007/s00227-012-2130-3 . hal-00804122

HAL Id: hal-00804122 https://hal.science/hal-00804122

Submitted on 5 Feb 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés. Marine Biology

April 2013, Volume 160, Issue 4, pp 751-761 <u>http://dx.doi.org/10.1007/s00227-012-2130-3</u> © Springer-Verlag Berlin Heidelberg 2012

The original publication is available at http://www.springerlink.com

Feeding grounds for waders in the Bay of the Mont Saint-Michel (France): the *Lanice conchilega* reef serves as an oasis in the tidal flats

Bart De Smet^{1, *}, Laurent Godet², Jérôme Fournier^{3, 4}, Nicolas Desroy⁵, Mikaël Jaffré⁶, Magda Vincx¹, Marijn Rabaut¹

- ² CNRS, UMR 6554 LETG-Nantes Géolittomer, Université de Nantes, B.P. 81227, 44312, Nantes Cedex 3, France
- ³ CNRS, UMR 7208 BOREA, Muséum National d'Histoire Naturelle, 7 Rue Cuvier, CP 32, 75231, Paris Cedex 05, France

⁴ Station Marine de Dinard, USM 404 Muséum National d'Histoire Naturelle, 38 Rue du Port Blanc, 35800, Dinard, France

⁵ IFREMER Laboratoire Environnement et Ressources FBN, CRESCO, 38 Rue du Port Blanc, 35800, Dinard, France

⁶ Université de Lille 1, UMR 8187 LOG, Station Marine de Wimereux, 28 Avenue Foch, 62930, Wimereux, France

*: Corresponding author : Bart De Smet, tel.: +32 (0)9 264 85 34 ; Fax: +32 (0)9 264 85 98 ; email address : <u>badsmet.desmet@ugent.be</u>

Abstract:

The tube-building polychaete *Lanice conchilega* can form dense populations, often called reefs, which promote benthic community change and constitute feeding grounds for secondary consumers. The aim of this study was to quantify the role of the *L. conchilega* reef of the Bay of the Mont Saint-Michel (BMSM) for feeding waders, by combining macrobenthos data, bird counts and bird diet information. Wader densities in the reef were on average 46.6 times higher than in non-reef areas. According to faecal analyses, waders in the reef mainly selected the accompanying fauna and especially crustaceans. The attractiveness of the reef to feeding birds may be largely explained by the high abundance, richness and biomass of macrobenthic species in the reef compared with the rest of the BMSM.

¹ Department of Biology, Marine Biology Section, Ghent University, Krijgslaan 281/S8, 9000, Ghent, Belgium

37 Introduction

38 Lanice conchilega is a widespread tubicolous polychaete that can form dense aggregations, considered biogenic reefs (Rabaut et al. 2009; Callaway et al. 2010). Above particular density thresholds, the 39 40 structures of the tubes as well as the biological activity of the "engineer species" (Callaway 2006; Godet et al. 2008), generate specific sedimentological "structures" (Carey 1987; Feral 1989) and 41 42 enhance the species diversity and abundance of the associated benthic macrofauna by stabilizing the 43 sediments (Zühlke 2001; Callaway 2006; Rabaut et al. 2007; Van Hoey et al. 2008). These reefs are 44 also important feeding grounds for flatfishes, particularly the juveniles of *Pleuronectes platessa* (Rijnsdorp and Vingerhoed 2001; Rabaut et al. 2010). 45

Lanice conchilega can be an important item in the diet of several waders (Goss-Custard and Jones 46 1976; Yates et al. 1993). Godet et al. (2009) showed that ovstercatchers may significantly select L. 47 conchilega reefs to feed in and that their spatial distribution can change greatly when these reefs 48 disappear. But to our knowledge, the study of Petersen and Exo (1999) in the German part of the 49 50 Wadden Sea is, so far, the only extensive study investigating the role of large L. conchilega dominated 51 tidal flats for waders and gulls. In comparison to the surrounding area, they found higher densities of 52 four bird species feeding in these flats, though the relative biomass consumption on these flats was 53 similar to other regions of the Wadden Sea. Furthermore, the study revealed that gulls tend to feed on L. conchilega specimens, while waders rather select the accompanying benthic macrofauna. 54

The area surveyed by Petersen and Exo (1999) was special because of the spatial dominance of *L. conchilega* sand flats (60% of the 6 680 ha of the study site), and because the remaining area was covered by two other habitat-creating species: *Arenicola marina* (20%) and *Mytilus edulis* (5 to 10%). Despite of the dominance of *L. conchilega*, the tubeworm aggregates did not generate the specific sedimentological structures, typically mounds and depressions, previously described for other *L. conchilega* reefs (e.g. Carey 1987).

The aim of this current study was to quantify the role of *L. conchilega* reefs as potential "oases" for feeding waders; do the reefs constitute a localized and discrete habitat for birds among large and

homogeneous tidal flats. Therefore, one of the largest intertidal L. conchilega reefs in Europe, located 63 in the Bay of the Mont Saint-Michel (BMSM) (France), was selected. For the first time, a combination 64 65 of macrobenthos data, bird counts and bird diet information was used to stress the importance of a Lanice conchilega reef. Following hypotheses have been tested: i) the benthic macrofaunal 66 composition of the reef clearly differs from the macrofaunal composition of the whole bay; ii) the 67 abundance and composition of waders on the reef is significantly different from the wader abundances 68 and composition at the scale of the whole BMSM. Additionally, the diet of waders feeding on the reef 69 70 was determined by means of a faecal analysis.

71 Material and Methods

72 Study area

The Lanice conchilega reef is located in the central region of the BMSM (48°40'45" N-01°41'25" W, 73 south-eastern part of the Normand-Breton Gulf, France) (Figure 1) and in the lower section of the tidal 74 flats. In 2008, the reef covered 105 ha; *i.e.* 0.42% of the sand flats of the BMSM (Godet et al. 2011) 75 (Figure 1). The BMSM is subjected to an extreme megatidal regime (tidal range up to 15.5 m during 76 77 spring tides), resulting in large tidal flats - covering 25 000 ha - and mainly dominated by a Macoma balthica community characterized by low macrobenthic abundances and diversity (Retière 1979; 78 79 Thorin et al. 2001). The BMSM is an internationally important migration stopover and wintering site for birds (Le Drean-Quenec'hdu et al. 1995), designated as a RAMSAR site and classified as a Special 80 81 Protection Area (SPA) and a Special Area of Conservation (SAC). More than 50 000 waders over-82 winter in the BMSM; 12% of the French wintering abundances of waders (Le Drean-Quenec'hdu et al. 1998). At this site, Dunlin Calidris alpina, Red Knot Calidris canutus, Oystercatcher Haematopus 83 84 ostralegus, Grey Plover Pluvialis squatarola, Black-tailed Godwit Limosa limosa, and Bar-tailed 85 Godwit Limosa lapponica reach international abundance levels in winter (Deceuninck and Mahéo 2000). 86

87 Macrobenthos sampling and treatment at the scale of the *L. conchilega* reef

Benthic macrofauna was sampled from the 10th to the 12th of January 2009 within 1 ha squares of a 88 regular grid (consisting of 150 squares). The L. conchilega density within each square was estimated 89 by counting aboveground tubes on pictures of 3 randomly selected $1/4 \text{ m}^2$ quadrats. Samples were only 90 taken in one out of every two squares of each row of the grid, though every square with L. conchilega 91 densities ≥ 200 ind. m⁻² was sampled as well (*i.e.* 80 sampled squares in total) (Figure 2). The number 92 93 of tubes is highly correlated with the number of individuals in the sediment (e.g. Van Hoey et al. 2006). At every selected square, one macrofaunal core was collected (1/40 m², 30 cm deep). Benthic 94 samples were sieved in the field through a 1 mm circular mesh size and the retained biological 95 material was immediately preserved in a 4.5% buffered formalin solution. In the laboratory, samples 96 97 were sorted and macrobenthos was identified to the lowest possible taxonomic level. Total biomass was estimated by determining the dry mass of all individuals per species (60 °C for 48 h). The ash-free 98 99 dry mass (AFDM) was calculated as the difference between the dry mass and the ash mass (500 °C for 100 3 h).

101 Wader counts at the scale of the BMSM

To assess the total number of waders in the whole BMSM, all waders of the site were monitored 5 102 103 times (January '09, March '09, May '09, September '09 and January '10) by 10 to 30 people. A 104 standardized protocol to monitor water birds of the BMSM, developed by two ornithological associations, Bretagne-Vivante/SEPNB and GONm, was followed (Beaufils et al. 2009). Waders were 105 counted in their high-tide roosts by people equipped with telescopes and binoculars. Because of the 106 107 extent of the BMSM, it was divided in sectors assigned to one or two observers. To avoid double 108 counts, surveys in the different sectors were performed on the same day and during the same time 109 interval (20 to 30 min). Additionally, any bird group seen flying from one sector to another was systematically reported with the exact time and flight direction. 110

111 Wader counts at the scale of the reef

112 On the reef, birds were surveyed 21 times from February 2009 to January 2010 (at least once each month, except for November and December), but only while spring tide fully exposed the reef (i.e. 113 corresponding to a low tide level of less than 2.5 m above extreme low water spring tide). Birds were 114 115 counted by two persons equipped with a pair of binoculars and a telescope (magnification respectively 116 10X and 20-60X). All individuals were counted (in case of a few tens of individuals) or estimated in 117 tens of individuals (if several hundreds or thousands of individuals were present). The observation point, located on a sandbank just outside the study area, ensured visibility of the entire site as well as a 118 minimal bird disturbance. 119

120 Estimating the diet of waders feeding on the reef: faeces sampling and treatment

In 2010, faeces of five wader species were collected within the reef: Oystercatcher *H. ostralegus*, Dunlin *C. alpina*, Curlew *Numenius arquata*, Grey Plover *P. squatarola* and Bar-tailed Godwit *L. lapponica*. We selected these species because they: i) are abundant in the reef; ii) feed regularly on the reef; iii) are species for which faeces are easy to collect because they feed in dense and virtually monospecific groups. Faeces of Curlew and Dunlin were collected on the 27th of April 2010, while faeces of Oystercatcher, Bar-tailed Godwit and Grey Plover were collected on the 13th of August 2010, the 9th of September 2010 and the 7th of October 2010 respectively.

128 Prior to collection, a large monospecific flock of birds feeding on the reef was observed in order to be confident that a dropping came from the target species. After 20 to 30 min (in order to be sure that the 129 collected faeces resulted from a feeding activity on the reef), the entire droppings were scraped off the 130 surface and preserved in 70% ethanol. In the laboratory, the ethanol was removed by pouring the 131 132 dropping onto a 20-µm sieve. For analysis, the sample was transferred to a 100-ml jar, containing a mix of 80% distilled water and 20% hydrogen peroxide (H₂O₂), for a period of at least 24 h. The 133 samples were shaken regularly and, after sedimentation, the supernatant was poured through a 20-µm 134 sieve. The supernatant remaining on the sieve was transferred to a 25-cm³ petri dish in order to 135 136 observe and identify lighter animal parts (e.g. polychaete chaetae) with an inverted microscope. A stereo-microscope was used to screen the entire settled sand fraction for hard remnants of bivalves,
polychaete jaws, etc. Prior to investigation, we selected for each of the five bird species 15 faecal
samples (except for Curlew: 11 samples).

Although all different items in a faecal sample were quantified, identification to genus or species level was not always possible. Therefore, each unique unidentified animal part was assigned to a morphotype, leading up to the creation of a catalogue consisting of pictures. Finally, for several analyses morphotypes were pooled into taxonomic groups, as mentioned hereafter.

144 Data and statistical analysis

145 Macrobenthic density, richness, diversity and production in the reef

Macrobenthic densities (D), species richness (S) and species diversity (H', Shannon Index, Shannon 146 147 1948) were calculated. Abundances of macrobenthos in the L. conchilega reef were evaluated by summing the numbers of individuals from the different squares. Subsequently, relative abundances 148 were calculated. The annual macrobenthic production in the reef was estimated using an empirical 149 model based on biomass and abundance data (Brey 1999, 2001). The model takes additional data on 150 151 benthic taxa and environmental variables such as bathymetry and temperature in consideration. Prior to production estimation, benthic biomass in g of AFDM was converted to kJ via conversion factors 152 for aquatic organisms (Brey 2001; Brey et al. 2010). 153

154 Abundances and community composition of waders on the reef versus the entire BMSM

The community composition of waders was analysed with the PRIMER v6 statistical package (Clarke 155 and Warwick 1994). Analysis of similarity (one-way ANOSIM) was used to describe (dis-)similarities 156 in wader communities between the entire bay and the L. conchilega reef. Data were standardized (in 157 order to eliminate the abundance effect) and square root transformed before conducting the analysis. 158 To test whether waders significantly selected the reef at low tide, we compared the abundances of 159 waders present on the reef with predicted wader abundances present on the reef assuming a 160 161 homogeneous distribution of birds across the entire BMSM during low tide. Because the ratio of the area covered by the reef to the area of the tidal flats is 1:238 (105 ha reef among 25 000 ha of tidal 162

flats), counting n individuals of a species at high tide in the whole BMSM results in a predicted 163 abundance of n/238 individuals of this species on the reef. Only counts which were performed at 164 165 similar time periods over the entire BMSM and the reef were selected for the analysis; i.e. four observation dates (March '09, May '09, September '09, and January '10). The number of days 166 between a count across the BMSM and on the reef varied between 0 and 12 days. G-tests for 167 goodness-of-fit were conducted in order to compare predicted and actual abundances of waders on the 168 169 reef, assuming no selective use of the reef habitat. A significant total G-value means that the data do 170 not fit the expected ratio.

171 Analysing bird faeces data

172 First, differences in the frequency of occurrence of benthic taxa among wader species were investigated. Therefore, bird faeces data were transformed into presence/absence data, followed by 173 174 lumping morphotypes into taxonomic groups. Due to diagnostic features of some morphotypes they could be linked to a certain species. Nonetheless, in most of the cases it was not possible to assign a 175 176 morphotype to a species but only to an order, class, or even phylum. For every wader species, the frequency of occurrence for each taxon i (FOi%) was calculated: FOi% = (the number of faecal 177 samples of the wader species s where taxon i is present / the total number of faecal samples of the 178 wader species s) * 100. Second, to determine which taxa are preferentially found in faecal samples of 179 180 particular wader species, the proportion of different taxonomic groups (Ni%) per faecal sample was calculated: Ni% = (Number of items in taxonomic group i / total number of items in the faecal sample) 181 * 100. To test whether relative abundances of higher taxonomic groups and the most abundant lower 182 taxonomic groups differed significantly among bird species, a generalized linear model (GLM) was 183 184 used in the SAS 9.2 software package (Glimmix procedure). Because the response variables are 185 percentage data, the residual error structure was tested against a binomial distribution. When overdispersion became apparent in the model output, the model was rerun, taking the overdispersion 186 187 into account by adding an overdispersion component (random residual) to the variance function. 188 Accordingly, underestimation of the standard errors was avoided. Because the predictor and the mean response are not linearly related to each other, the relationship was specified by a log link function. 189

190 Results

191 Macrobenthic density, richness, diversity, biomass and production in the Lanice conchilega reef

192 In 2009, 13 806 macroinvertebrates belonging to 61 different taxa were sampled on the reef. Excluding L. conchilega itself, the macrobenthic abundance was dominated by the bivalves Macoma 193 194 balthica and Cerastoderma edule and the polychaete Nephtys hombergii. Taking into account L. *conchilega*, the mean density of macrobenthic species was 6 903 \pm 5 339 ind. m⁻², N = 80; mean 195 species richness was 11.8 ± 4.7 species per square and mean species diversity (H') was 2.1 ± 0.5 . At 196 the phylum level, the benthic community within the reef was dominated by annelids (59%), followed 197 198 by molluscs (38%) and arthropods (1.8%). More than 99.9% of the annelids in the reef belonged to the class Polychaeta. Moreover, this taxon was dominated by L. conchilega (69%). The average L. 199 conchilega density was 200 ± 351 ind.m⁻², N = 150, and a maximum density of 1985 ind.m⁻² was 200 reached. Within the phylum of Mollusca, the most abundant species were M. balthica (55%) and C. 201 202 edule (44%). Crustaceans were the most abundant taxon within the phylum of the arthropods (99.6%). 203 The order of the amphipods (71%) dominated the crustaceans within the reef. Cumaceans, isopods and decapods (crabs) constituted respectively 14.7%, 7.8%, and 6.5% of total abundances. The mean 204 benthic biomass in the reef was 49.7 ± 50.4 g of AFDM.m⁻². The annual macrobenthic production in 205 the reef (95% confidence interval) was estimated at 1552.9 (1368.5-1762.3) kJ.m⁻².year⁻¹ or 70.6 g 206 AFDM.m⁻².year⁻¹ (60.2-80.1). 207

208 Wader density and community composition in the BMSM and on the reef

Across the entire BMSM, 22 wader species were counted, representing a mean density of 1.03 ± 0.58 birds ha⁻¹, N = 5. The five most abundant species were: Dunlin (42%), Red Knot (18%), Oystercatcher (13%), Grey Plover (9%) and Curlew (7%). On the reef, 15 wader species were counted, representing a mean density of 51.38 ± 19.11 birds ha⁻¹, N = 5. Dunlin (39%) was the most abundant species on the reef followed by Red Knot (20%), Grey Plover (14%), Oystercatcher (13%), Bar-tailed Godwit (8%) and Curlew (5%).

The one-way global ANOSIM test failed to detect a significant difference in the wader community in 215 the two habitats (p > 0.05; R = 0.108). Unlike bird composition, bird counts revealed that all wader 216 217 species together exhibited a high proportion of individuals in the reef in relation to the whole BMSM (Table 1). The frequency of waders in the reef was on average 46.6 times higher than expected 218 assuming a random distribution of waders over the entire BMSM. The same was observed for the five 219 wader species selected for the faecal analysis. Focussing on each of the 5 wader species separately 220 221 revealed frequencies ranging from 31.1 times (Curlew) up to 112.3 times (Bar-tailed Godwit) higher than expected. Consequently, the observed numbers of these 5 species in the reef were significantly 222 223 higher than their predicted numbers (assuming that the total number of waders counted at high tide 224 have a homogeneous distribution in the BMSM at low tide) (for each species, G-test, p < 0.001).

225 Diet of waders feeding on the reef

226 In general, the frequency of occurrence (FO%) of higher taxonomic groups in the faeces did not differ much among different bird species (Table 2). Crustaceans were present in all faecal samples of all 227 birds except for Bar-tailed Godwit (FO% = 93.3%). Both polychaetes and bivalves were present in all 228 bird species but polychaetes in a much higher percentage of the faeces (ranging from 72.7% in Curlew 229 to 93.3% in Dunlin) than bivalves (ranging from 13.3% in Oystercatcher and Grey Plover to 45.5% in 230 231 Curlew). Lanice conchilega was eaten by all birds though it never exceeded a FO% of 63.6%. Other 232 polychaetes were scarce. Crabs, amphipods and ostracods - the most abundant crustacean groups in the faces - were encountered in all bird species and reached the highest FO% in Curlew (respectively 233 81.8%, 45.5% and 90.9%). The three most frequently occurring bivalve species among all bird species 234 235 were Abra alba, C. edule and M. balthica, although they never exceeded a FO% of 6.7%, 36.4%, and 20% respectively. 236

Based on the relative abundances of taxonomic groups in each of the 5 bird species a diet composition can be displayed for the 5 bird species examined (Table 3). Relative abundances of all higher taxonomic groups differed significantly (p < 0.05) among the 5 wader species (GLM, p < 0.05, Table 4). Globally, polychaetes differed significantly among species of birds, though no significant pairwise 241 differences were detected. Crustaceans were more frequently eaten by Curlew than by Oystercatcher or Bar-tailed Godwit, while bivalves were eaten more frequently by Dunlin than Curlew. Significant 242 243 differences in the relative abundances of L. conchilega (GLM, p < 0.0001) and 'Other polychaetes' (GLM, p < 0.0001) were detected among bird species (Table 5). According to pairwise tests, L. 244 conchilega was eaten more frequently by Oystercatcher than by Dunlin and Curlew. Additionally, 245 Bar-tailed Godwit ate L. conchilega more frequently than Dunlin, but less than Curlew. Within the 246 247 group of crustaceans, 'Crabs' differed significantly among bird species (GLM, p = 0.005), although no significant pairwise differences were detected. Lastly, no bivalve species showed a significant 248 249 difference among waders (GLM, p > 0.05).

250 Discussion

Our study showed that the Lanice conchilega reef of the BMSM is remarkable because of the wader 251 density which easily exceeds the expected frequency. In the entire BMSM a total of 22 wader species 252 was observed and counted during the study period, while only 15 species were observed in the L. 253 conchilega reef. Since the entire BMSM is almost 240 times bigger than the reef area, the observed 254 difference in species richness is not surprising keeping in mind the species-area relationship (Connor 255 256 and McCoy 1979). Consequently, species that were rarely counted at the scale of the entire BMSM are 257 not likely to be observed at the scale of the reef. Despite this difference, both areas were mainly dominated by the same species: Dunlin, Red Knot, Grey Plover, Oystercatcher and Curlew. Eybert et 258 259 al. (2003) already demonstrated that 96% of the wintering shorebird community in the entire BMSM was represented by 7 species: Dunlin, Oystercatcher, Red Knot, Curlew, Grey Plover, Bar-tailed 260 261 Godwit and Black-tailed Godwit. Overall, the composition of waders can be considered the same in the entire BMSM and the reef. 262

However, when abundances are included, the composition of waders clearly differed between the two study sites. The observed frequency of total waders on the *Lanice*-reef was on average 5 799 birds, which is approximately 46.6 times higher than expected assuming a uniform distribution of the birds in the BMSM. Petersen and Exo (1999) observed higher than expected bird abundances in *L*. *conchilega* dominated tidal flats of the Wadden Sea, although densities were only 6 times higher than
 expected. Similar counting surveys in the Chausey archipelago (France) and preliminary counts in the
 BMSM already showed the attractiveness of the *L. conchilega* reefs for birds and consequently their
 potentially important role in the conservation of the avifauna (Godet et al. 2008). The results of the
 current study prove that within a site of international importance for birds, several wader species are
 able to select preferentially habitats generated by *L. conchilega*.

273 In general, the attractiveness of L. conchilega reefs can be attributed to the good food supply, i.e. the 274 high diversity, abundance and biomass of associated macrobenthic invertebrates, as proven by several authors in different study areas (e.g. Zühlke et al. 1998; Zühlke 2001; Callaway 2006; Rabaut et al. 275 276 2007; Van Hoey et al. 2008). The situation in the *L. conchilega* reef of the BMSM seems to be alike. 277 In our study, the macrofaunal density is 4.4 times higher than in a study of Trigui (2009), which 278 investigated the general characteristics of the benthic macrofauna of the entire intertidal zone of the 279 BMSM in 2003. Trigui's survey is, with 176 sampled stations, the most extensive benthic survey ever 280 done in the BMSM. Comparison with current research reflects well the fact that the reef does accommodate a more abundant fauna (6 903 \pm 5 339 ind.m⁻²) than the average macrobenthic 281 assemblage in the entire BMSM (1 568 \pm 299 ind.m⁻²). As part of a study of Leloup et al. (2008), 282 biomasses of different trophic groups which make up the BMSM were modelled. The biomass of 283 284 carnivorous and necrophagous macrobenthic fauna in combination with intertidal filter feeders was transformed according to weight-to-weight conversion factors proposed by Ricciardi and Bourget 285 (1998). Comparing the resulting biomass (2.88 g AFDM. m⁻²) to the reef biomass (49.69 g AFDM.m⁻²) 286 revealed a reef biomass which is more than 17 times higher. Therefore, the reef area can be considered 287 a high productivity area in the bay. Nevertheless, the importance of the whole BMSM for the 288 productivity and functioning of the reef cannot be neglected. 289

Faecal analysis conducted in this study revealed information on the diet composition of waders in the reef. Polychaetes were represented in all bird species and they occurred in a high percentage of the faecal samples, which can probably be attributed to the dominance of polychaetes in the reef benthos. *Lanice conchilega* was the most abundant species and polychaete in the reef, which was partially 294 reflected in the faecal samples since L. conchilega was the only identifiable polychaete species present 295 in all bird species. Nevertheless, based on the relative abundances on the reef it could be expected that 296 L. conchilega counted as a larger part of a bird's diet. The fact that the tube-building polychaete is 297 large (up to 30cm; Hartmann-Schröder 1996), sturdy and buried quite deeply in the sediment (Jones and Jago 1993) can lead to a lower accessibility and possibly explains the lower than expected portion 298 299 of L. conchilega in the diet of waders. Additionally, the fact that the energy content of polychaetes in 300 temperate waters is high (Dauvin and Joncourt 1989), in combination with the high biomass of L. conchilega in the reef, might lead to the great contribution of one individual to the nutritional demands 301 302 of birds. The bivalves Macoma balthica and Cerastoderma edule were, next to L. conchilega, the most 303 abundant macrobenthic species in the reef. However, the general occurrence and portion of these two 304 species in the diet of the investigated bird species was low. Even in Oystercatchers, which are 305 specialized bivalve feeders (e.g. Hulscher 1982), bivalves were underrepresented. A feeding strategy avoiding the uptake of bivalve shell pieces, as described by Hulscher (1982), and hence the low 306 307 detection success of shell remains in the faeces of Oystercatchers, is the most plausible explanation. 308 Crustaceans, and in particular crabs and ostracods, were the most frequently occurring and abundant 309 taxonomic group in the faeces of all bird species. However, relative abundances of crustaceans in the reef benthos were very low (1.77% of the total abundance), particularly for crabs. The observed 310 311 ubiquity of crabs (and crustaceans in general) in the faecal samples implies a selective feeding 312 behaviour of waders for this benthic group, which may be due to the fact that the stimuli associated with crustaceans, can have properties which make them particularly perceptible to the avian eye 313 314 (Goss-Custard 1977).

Based on relative abundances of taxonomic groups, bird species that largely dominate the wader community on the reef exhibited significant differences in their dietary composition. Differences in polychaetes, and especially *L. conchilega*, contributed highly to differences in the wader diets. Differences in diets due to bivalves were rather low and in all probability largely overshadowed by an inability to trace back bivalve shells in faeces. For crustaceans, differences in the relative abundance between waders were the result of significant differences in crabs, though no significant pairwise tests were noted. Nevertheless, it can be visually seen that crabs reached much higher abundances in the
faeces of Curlew compared to other waders (Table 3), which is in accordance with studies revealing
that crabs are a major food source of Curlew (e.g. Goss-Custard and Jones 1976).

324 Considering the frequency of occurrence of taxonomic groups in the diet, waders foraging within the 325 reef can be seen as opportunistic feeders, while focusing on the relative abundance gives the 326 impression of a more selective feeding strategy. Based on current wader diet analysis, the actual 327 strategy in the reef probably lies somewhere in between these two extremes. In general, waders 328 feeding on the reef tend to feed on the associated fauna, and especially crustaceans, rather than specifically on L. conchilega. This result confirms the investigations of Petersen and Exo (1999), 329 330 providing evidence that L. conchilega, within L. conchilega dominated tidal flats, was a less important 331 food source for birds (mainly waders) than the accompanying macrobenthos. However, L. conchilega 332 was of overriding importance for a few species, especially for gulls (Petersen and Exo 1999).

333 As experienced in this current study, investigating bird faeces can give us better insights into the diet of waders feeding in the reef. However, faecal analysis is hampered by a high proportion of 334 335 unidentified prey items and is unlikely to reveal all prey taken by the predator (Barrett et al. 2007). 336 Despite some shortcomings, faecal analysis is a valuable and easily applicable technique for which 337 samples can be obtained all-year round without causing harm to the birds. Moreover, this approach 338 can reveal the presence of prey species which could not be noticed by means of visual observations. Nevertheless, to get a more complete view on a wader's diet, the use of complementary approaches 339 (e.g. visual observations) is recommended (Scheiffarth 2001). In this study morphotypes were 340 341 counted, which cannot be compared with specimens. Since morphotypes are mostly parts of animals and can belong to one or several specimens of the same species or taxonomic group, caution regarding 342 the interpretation of the results is recommended. Faecal samples of the wader species were collected 343 344 on different dates ranging from the end of April 2010 until the beginning of October 2010. It is known 345 that the diet composition of waders, as well as the nutritive value of some macrobenthic organisms, 346 can change seasonally (e.g. Scheiffarth 2001; Braeckman et al. 2012). Furthermore, sexual differences 347 in the diet composition have been demonstrated for several birds (e.g. Bar-tailed Godwit; Scheiffarth

2001). As neither sexual nor seasonal variations in the diet composition of waders were taken into
account in our study, comparing wader diets is restricted. Additionally, future studies should try to
compare diet composition of waders both inside and outside the *L. conchilega* reef.

351 Evaluating the importance of *Lanice conchilega* reefs as trophic resources for waders is not merely fundamental to gain knowledge on the feeding ecology, but is also essential for predicting the effects 352 of a possible loss of the reef habitat in the future. Overall, the L. conchilega reef of the BMSM can be 353 354 considered an oasis within the tidal flats composed of a similar composition of waders but with much 355 higher bird densities compared to the non-reef areas of the BMSM. This result clearly demonstrates that birds are attracted by the reef. According to faecal analyses, waders in the L. conchilega reef tend 356 357 to feed on the associated fauna, and especially crustaceans, rather than specifically on L. conchilega itself. Within the BMSM, which is characterized by low species diversity and low macrofaunal 358 359 abundances, the L. conchilega reef constitutes a rich feeding area. For future studies, the use of 360 complementary techniques to study a wader's diet is recommended, as well as the inclusion of 361 information on the alimentary regime of waders in the BMSM.

362 Acknowledgements

363 We thank Matthieu Beaufils (Bretagne-Vivante/SEPNB, La société pour l'Etude et la Protection de la Nature en Bretagne, and GONm, Groupe Ornithologique Normand), Sébastien Provost (GONm) and 364 Régis Morel (Bretagne-Vivante/SEPNB) for coordinating and providing the bird counts at the scale of 365 366 the BMSM for 2009. Both Bretagne-Vivante/SEPNB and the GONm are acknowledged for funding the counts at the scale of the BMSM. Thanks to Thomas Brey for providing and helping with the 367 empirical model to estimate benthic production and to Andrea Moore for help with the faecal analysis. 368 369 We also thank the anonymous reviewers who improved earlier versions of the manuscript. The first 370 author acknowledges a PhD scholarship provided by the Special Research Fund (BOF), Ghent University, Belgium. The second author acknowledges the 'Centre National de la Recherche 371 Scientifique' for his Post-Doc Grant. 372

373 References

- Barrett RT, Camphuysen K, Anker-Nilssen T, Chardine JW, Furness RW, Garthe S, Hüppop O,
 Leopold MF, Montevecchi WA, Veit RR (2007) Diet studies of seabirds: a review and
 recommendations. Ices J Mar Sci 64 (9):1675-1691
- Beaufils M, Morel R, Provost S (2009) Suivi 2009 de la fréquentation de la Baie du Mont Saint Michel par les oiseaux d'eau. Rapport SEPNB-GONm
- Braeckman U, Provoost P, Sabbe K, Soetaert K, Middelburg JJ, Vincx M, Vanaverbeke J (2012)
 Temporal dynamics in the diet of two marine polychaetes as inferred from fatty acid
 biomarkers. J Sea Res 68 (0):6-19. doi:10.1016/j.seares.2011.11.003
- Brey T (1999) A collection of empirical relations for use in ecological modelling. NAGA The
 ICLARM Quarterly 22 (3):24-28
- Brey T (2001) Population dynamics in benthic invertebrates. A virtual handbook. <u>http://www.thomas-brey.de/science/virtualhandbook</u>. Accesses 1st February 2012.
- Brey T, Muller-Wiegmann C, Zittier ZMC, Hagen W (2010) Body composition in aquatic organisms A global data bank of relationships between mass, elemental composition and energy content.
 J Sea Res 64 (3):334-340. doi:10.1016/j.seares.2010.05.002
- 389 Callaway R (2006) Tube worms promote community change. Mar Ecol-Prog Ser 308:49-60.
 390 doi:10.3354/meps308049
- Callaway R, Desroy N, Dubois SF, Fournier J, Frost M, Godet L, Hendrick VJ, Rabaut M (2010)
 Ephemeral Bio-engineers or Reef-building Polychaetes: How Stable are Aggregations of the
 Tube Worm *Lanice conchilega* (Pallas, 1766)? Integrative and Comparative Biology 50
 (2):237-250. doi:10.1093/icb/icq060
- Carey DA (1987) Sedimentological effects and paleoecological implications of the tube-building
 polychaete *Lanice conchilega* Pallas. Sedimentology 34 (1):49-66. doi:10.1111/j.1365 3091.1987.tb00559.x
- Clarke KR, Warwick RM (1994) Change in marine communities: an approach to statistical analysis
 and interpretation. Primer-E Plymouth Marine Laboratory, UK
- Connor EF, McCoy ED (1979) The Statistics and Biology of the Species-Area Relationship. Am Nat 113 (6):791-833
- 402 Dauvin JC, Joncourt M (1989) Energy values of marine benthic invertebrates from the western english
 403 channel. Journal of the Marine Biological Association of the United Kingdom 69 (3):589-595
- 404 Deceuninck B, Mahéo R (2000) Synthèse des dénombrements et analyse des tendances des limicoles
 405 hivernant en France 1978-1999. Rapport DNP:82
- Eybert M-C, Geslin T, Questiau S, Feunteun E (2003) Shorebird community variations indicative of a
 general perturbation in the Mont-Saint-Michel (France). CR Biologies 326:140-147
- Feral P (1989) Influence des populations de *Lanice conchilega* (Pallas)(Annelida, Polychaeta) sur la sedimentation sableuse intertidale de deux plages bas-normandes (France). Bull Soc Geol Fr
 8:1193-1200

- Godet L, Toupoint N, Olivier F, Fournier J, Retière C (2008) Considering the functional value of
 common marine species as a conservation stake: The case of sandmason worm *Lanice conchilega* (Pallas 1766) (Annelida, Polychaeta) beds. Ambio 37 (5):347-355.
 doi:10.1579/07-a-317.1
- Godet L, Toupoint N, Fournier J, Le Mao P, Retiere C, Olivier F (2009) Clam farmers and 415 Oystercatchers: Effects of the degradation of Lanice conchilega beds by shellfish farming on 416 417 the spatial distribution of shorebirds. Mar Pollut Bull 58 (4):589-595. doi:10.1016/j.marpolbul.2008.11.001 418
- Godet L, Fournier J, Jaffre M, Desroy N (2011) Influence of stability and fragmentation of a wormreef on benthic macrofauna. Est Coast Shelf Sci 92 (3):472-479.
 doi:10.1016/j.ecss.2011.02.003
- 422 Goss-Custard JD (1977) The energetics of prey selection by redshank, *Tringa totanus* (L.), in relation
 423 to prey density. J Anim Ecol 46 (1):1-19
- 424 Goss-Custard JD, Jones RE (1976) The diets of redshank and curlew. Bird Study 23:233-270
- Hartmann-Schröder G (1996) Annelida, Borstenwürmer, Polychaeta. In: The fauna of Germany and adjacent seas with their characteristics and ecology, second revised ed., vol. 58. Gustav Fisher, Jena, Germany, 648 p.
- Hulscher JB (1982) The oystercatcher *Haematopus ostralegus* as a predator of the bivalve *Macoma balthica* in the dutch Wadden Sea. Ardea 70 (2):89-152
- Jones SE, Jago CF (1993) In situ assessment of modification of sediment properties by burrowing
 invertebrates. Mar Biol 115:133-142
- Le Drean-Quenec'hdu S, Mahéo R, Boret P (1995) The Mont Saint-Michel bay: a site of international
 importance for wintering Palearctic waders. Wader Study Group Bull 77:50-54
- Le Drean-Quenec'hdu S, Mahéo R, Boret P (1998) Les limicoles. Penn ar bed 169: La Baie du Mont
 Saint-Michel:1-10
- Leloup FA, Desroy N, Le Mao P, Pauly D, Le Pape O (2008) Interactions between a natural food web,
 shellfish farming and exotic species: The case of the Bay of Mont Saint Michel (France). Est
 Coast Shelf Sci 76 (1):111-120. doi:10.1016/j.ecss.2007.06.011
- Petersen B, Exo KM (1999) Predation of waders and gulls on *Lanice conchilega* tidal flats in the
 Wadden Sea. Mar Ecol Prog Ser 178:229-240. doi:10.3354/meps178229
- Rabaut M, Guilini K, Van Hoey G, Vincx M, Degraer S (2007) A bio-engineered soft-bottom
 environment: The impact of *Lanice conchilega* on the benthic species-specific densities and
 community structure. Est Coast Shelf Sci 75 (4):525-536. doi:10.1016/j.ecss.2007.05.041
- Rabaut M, Vincx M, Degraer S (2009) Do *Lanice conchilega* (sandmason) aggregations classify as reefs? Quantifying habitat modifying effects. Helgoland Mar Res 63 (1):37-46. doi:10.1007/s10152-008-0137-4
- Rabaut M, Van de Moortel L, Vincx M, Degraer S (2010) Biogenic reefs as structuring factor in
 Pleuronectes platessa (Plaice) nursery. J Sea Res 64 (1-2):102-106.
 doi:10.1016/j.seares.2009.10.009
- 450 Retière C (1979) Contribution à la connaissance des peuplements benthiques du Golfe Normano 451 Breton. Dissertation, Université de Rennes

- 452 Ricciardi A, Bourget E (1998) Weight-to-weight conversion factors for marine benthic
 453 macroinvertebrates. Mar Ecol Prog Ser 163:245-251. doi:10.3354/meps163245
- Rijnsdorp AD, Vingerhoed B (2001) Feeding of plaice *Pleuronectes platessa* L. and sole *Solea solea*(L.) in relation to the effects of bottom trawling. J Sea Res 45 (3-4):219-229. doi:10.1016/s1385-1101(01)00047-8
- 457 Scheiffarth G (2001) The diet of Bar-tailed Godwits Limosa lapponica in the Wadden Sea: Combining
 458 visual observations and faeces analyses. Ardea 89 (3):481-494
- 459 Shannon CE (1948) A mathematical theory of communication. Bell System Technical Journal 27:379 460 423
- 461 Thorin S, Radureau A, Feunteun E, Lefeuvre JC (2001) Preliminary results on a high east-west
 462 gradient in the macrozoobenthic community structure of the macrotidal Mont Saint-Michel
 463 bay. Cont Shelf Res 21 (18-19):2167-2183. doi:10.1016/s0278-4343(01)00050-4
- 464 Trigui RJ (2009) Influence des facteurs environnementaux et anthropiques sur la structure et la fonctionnement des peuplements benthiques du Golfe Normano-Breton. Dissertation, Muséum National d'Histoire Naturelle, France
- Van Hoey G, Vincx M, Degraer S (2006) Some recommendations for an accurate estimation of *Lanice conchilega* density based on tube counts. Helgoland Mar Res 60 (4):317-321.
 doi:10.1007/s10152-006-0041-8
- Van Hoey G, Guilini K, Rabaut M, Vincx M, Degraer S (2008) Ecological implications of the
 presence of the tube-building polychaete *Lanice conchilega* on soft-bottom benthic
 ecosystems. Mar Biol 154 (6):1009-1019. doi:10.1007/s00227-008-0992-1
- Yates MG, Goss-Custard JD, McGrorty S, Lakhani KH, Durell S, Clarke RT, Rispin WE, Moy I,
 Yates T, Plant RA, Frost AJ (1993) Sediment characteristics, invertebrate densities and
 shorebird densities on the inner banks of the Wash. J Appl Ecol 30 (4):599-614.
 doi:10.2307/2404240
- Zühlke R (2001) Polychaete tubes create ephemeral community patterns: *Lanice conchilega* (Pallas, 1766) associations studied over six years. J Sea Res 46 (3-4):261-272. doi:10.1016/s1385-1101(01)00091-0
- Zühlke R, Blome D, Van Bernem KH, Dittmann S (1998) Effects of the tube-building polychaete
 Lanice conchilega (Pallas) on benthic macrofauna and nematodes in an intertidal sandflat.
 Senckenb Marit 29:131-138

483 Figure Legends

- 484 Fig. 1 Location of the Bay of the Mont Saint-Michel (BMSM, France) and the *Lanice conchilega* reef
 485 within the bay
- 486 Fig. 2 Macrobenthic sampling design and mean *Lanice conchilega* densities on the *L. conchilega* reef
- 487 in the Bay of the Mont Saint-Michel (BMSM) from 2005 to 2008. Macrobenthic samples were taken
- 488 at alternating squares of each row of the grid (consisting of 150 squares). In addition, every square
- 489 with a mean *L*. *conchilega* density ≥ 200 ind. m⁻² was sampled as well, resulting in 80 sampled squares
- 490 in total





Table 1 Overview of the mean observed and mean expected bird frequencies, and the ratios of these frequencies, for the entire Bay of the Mont Saint-Michel (BMSM) and the *L. conchilega* reef; total G-values, and p-values (G-test) of different groups of birds. Observed frequencies in the BMSM and the *L. conchilega* reef were derived from bird counts in March '09, May '09, September '09, and January '10

		BMSM		L	anice-reef			
	Obs. freq. ± SD	Exp. freq. ± SD	Ratio	Obs. freq. ± SD	Exp. freq. ± SD	Ratio	Total <i>G</i> -value	p-value
Total waders	23928 ± 15976	29602 ± 17513	0.81	5799 ± 1994	124 ± 74	46.64	140976	< 0.0001
5 Species	17651 ± 10761	21830 ± 12882	0.81	$4271{\pm}2444$	92 ± 54	46.58	101915	< 0.0001
Dunlin	9853 ± 9478	12080 ± 11337	0.82	2278 ± 1972	51 ± 48	44.90	53565	< 0.0001
Oystercatcher	3381 ± 1963	3916 ± 2502	0.86	551 ± 614	16 ± 11	33.52	12206	< 0.0001
Grey Plover	2315 ± 884	3157 ± 1476	0.73	856 ± 798	13 ± 6	64.52	23768	< 0.0001
Curlew	1734 ± 1171	1984 ± 1340	0.87	259 ± 274	8 ± 6	31.08	5642	< 0.0001
Bar-tailed Godwit	369 ± 400	694 ± 819	0.53	327 ± 425	3 ± 3	112.34	10697	< 0.0001

Table 2 Frequency of occurrence (FOi%) of higher and the most abundant lower taxonomic groups for all five investigated waders, based on faecal analysis. N = the number of faecal samples investigated. 100% = present in all droppings

	<i>C. alpina</i> (Dunlin)	<i>H. ostralegus</i> (Oystercatcher)	N. arquata (Curlew)	P. squatarola (Grey Plover)	L. lapponica (Bar-tailed Godwit)
Taxon	N = 15	N = 15	N = 11	N = 15	N = 15
Polychaeta	93.3	73.3	72.7	73.3	86.7
Lanice conchilega	40	60	63.6	13.3	33.3
Nereis sp.	6.7	0	9.1	0	0
<i>Eteone longa</i> Other polychaetes	0 93.3	0 73.3	9.1 72.7	0 60	0 73.3
Crustacea	100	100	100	100	93.3
Crab sp.	73.3	53.3	81.8	80	46.7
Amphipoda sp.	33.3	33.3	45.5	26.7	33.3
Ostracoda sp.	80	73.3	90.9	53.3	40
Isopoda sp.	0	0	9.1	0	6.7
<i>Cumacea</i> sp.	0	0	9.1	0	6.7
<i>Copepoda</i> sp.	0	13.3	0	33.3	20
Other crustaceans	100	100	100	100	93.3
Bivalvia	33.3	13.3	45.5	13.3	26.7
Abra alba	0	6.7	0	6.7	6.7
Aequipecten opercularis	0	0	9.1	0	0
Cerastoderma edule	0	0	36.4	6.7	0
Macoma balthica	20	0	9.1	0	0
Mysella bidentata	0	0	0	0	6.7
Nucula sp.	0	6.7	0	0	0
Scrobicularia plana	0	0	0	0	6.7
Spisula subtruncata	0	0	18.2	0	0
Other bivalves	26.7	0	18.2	0	6.7
Other	100	100	100	100	100

		C. al (Du	<i>lpina</i> nlin)	H. ostralegus (Oystercatcher) N = 15		N. ar (Cur	N. arquata (Curlew) N = 11		P. squatarola (Grey Plover)		<i>pponica</i> ed Godwit)
		IN =	= 15	N =	IN = 13		N = 11		N = 15		= 15
	Taxon	%	\pm SD	%	\pm SD	%	\pm SD	%	\pm SD	%	\pm SD
ха	Polychaeta	17.56	17.57	15.15	24.86	18.17	29.61	5.98	9.59	8.93	18.59
sr ta	Crustacea	43.31	24.49	26.75	18.16	60.46	35.11	47.26	25.39	18.46	9.65
ighe	Bivalvia	0.23	0.40	0.03	0.10	0.07	0.17	0.01	0.03	0.02	0.05
Н	Other	38.90	24.57	58.06	26.07	21.30	26.64	46.75	24.63	72.58	21.24
ta	L. conchilega	6.81	12.19	39.79	32.34	20.10	33.57	18.18	40.45	19.68	36.95
hae	Nereis sp.	0.48	1.78	0	0	0.02	0.05	0	0	0	0
olyc	E. longa	0	0	0	0	1.01	3.03	0	0	0	0
P	Other poly.	92.71	12.03	60.21	32.34	78.88	33.32	81.82	40.45	80.32	36.95
	Crab sp.	4.60	5.86	5.12	7.25	32.84	34.00	18.10	18.13	10.06	17.18
	Amphipoda sp.	0.81	1.36	1.30	2.34	6.25	19.06	0.46	0.93	5.69	10.23
cea	Ostracoda sp.	9.18	15.06	13.36	17.84	15.81	19.30	8.17	22.37	9.15	13.90
ıstad	Isopoda sp.	0	0	0	0	0.83	2.74	0	0	0.71	2.67
Cu	Cumacea sp.	0	0	0	0	0.001	0.003	0	0	0.71	2.67
	<i>Copepoda</i> sp.	0	0	0.92	2.90	0	0	0.41	0.78	1.64	3.56
	Other crust.	85.40	19.00	79.30	18.51	44.26	28.28	72.86	25.66	72.04	23.76
	A. alba	0	0	50.00	70.71	0	0	50.00	70.71	25.00	50.00
	A. opercularis	0	0	0	0	2.86	6.39	0	0	0	0
	C. edule	0	0	0	0	49.52	47.57	50.00	70.71	0	0
'ia	M. balthica	34.50	44.17	0	0	2.86	6.39	0	0	0	0
valv	M. bidentata	0	0	0	0	0	0	0	0	25.00	50.00
Bi	Nucula sp.	0	0	50.00	70.71	0	0	0	0	0	0
	S. plana	0	0	0	0	0	0	0	0	25.00	50.00
	S. subtruncata	0	0	0	0	18.10	26.17	0	0	0	0
	Other bivalves	65.50	44.17	0	0	26.67	43.46	0	0	25.00	50.00

Table 3 Relative abundance (\pm SD) of all higher and the most abundant lower taxonomic groups in the diet composition of all five investigated wader species, based on faecal analysis. N = the number of faecal samples investigated

Table 4 Global p-values (GLM) and adjusted p-values of the pairwise tests (Tukey-Kramer) to check differences in the relative abundances of higher taxonomic groups among the five investigated waders. In case of significant differences (p < 0.05) p-values are in bold. D=Dunlin, O=Oystercatcher, C=Curlew, G=Grey Plover, B=Bar-tailed Godwit

	Global	D - 0	D - C	D - G	D - B	0 - C	0 - G	0 - B	C - G	C - B	G - B
Polychaetes	0.0254	0.509	1.000	0.951	0.398	0.197	0.242	1.000	0.894	0.095	0.183
Crustaceans	<0.0001	0.100	0.531	1.000	0.069	0.0010	0.124	1.000	0.354	<0.001	0.087
Bivalves	0.0013	0.307	0.004	0.294	0.182	0.999	1.000	0.994	0.998	0.999	0.992
Other	0.0009	0.833	0.063	0.894	0.898	0.002	1.000	1.000	0.002	0.002	1.000

Table 5 Global p-values (GLM) and adjusted p-values of the pairwise tests (Tukey-Kramer) to check differences in the relative abundances of lower taxonomic groups (*Lanice conchilega*, Other polychaetes, Crabs and Other crustaceans) among the five investigated wader species. In case of significant differences (p < 0.05) p-values are in bold. D=Dunlin, O=Oystercatcher, C=Curlew, G=Grey Plover, B=Bar-tailed Godwit

	Global	D - 0	D - C	D - G	D - B	O - C	0 - G	0 - B	C - G	C - B	G - B
L. conch.	<0.0001	0.039	1.000	0.997	0.023	0.0003	0.675	0.992	0.997	0.0003	0.607
Other poly.	<0.0001	0.036	1.000	0.997	0.022	0.0003	0.672	0.992	0.996	0.0003	0.604
Crab	0.005	1.000	0.070	0.769	1.000	0.570	0.950	1.000	0.135	0.468	0.950
Other crust.	<0.0001	0.830	0.0001	0.704	0.896	0.094	1.000	1.000	0.0002	0.045	1.000