



HAL
open science

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

<http://dx.doi.org/10.1007/s00227-012-2130-3>

© Springer-Verlag Berlin Heidelberg 2012

Archimer
<http://archimer.ifremer.fr>

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

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

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 : badsmet.desmet@ugent.be

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.

37 **Introduction**

38 *Lanice conchilega* is a widespread tubicolous polychaete that can form dense aggregations, considered
39 biogenic reefs (Rabaut et al. 2009; Callaway et al. 2010). Above particular density thresholds, the
40 structures of the tubes as well as the biological activity of the “engineer species” (Callaway 2006;
41 Godet et al. 2008), generate specific sedimentological “structures” (Carey 1987; Feral 1989) and
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*
45 (Rijnsdorp and Vingerhoed 2001; Rabaut et al. 2010).

46 *Lanice conchilega* can be an important item in the diet of several waders (Goss-Custard and Jones
47 1976; Yates et al. 1993). Godet et al. (2009) showed that oystercatchers may significantly select *L.*
48 *conchilega* reefs to feed in and that their spatial distribution can change greatly when these reefs
49 disappear. But to our knowledge, the study of Petersen and Exo (1999) in the German part of the
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
54 *L. conchilega* specimens, while waders rather select the accompanying benthic macrofauna.

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

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

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

71 **Material and Methods**

72 Study area

73 The *Lanice conchilega* reef is located in the central region of the BMSM (48°40'45" N-01°41'25" W,
74 south-eastern part of the Normand-Breton Gulf, France) (Figure 1) and in the lower section of the tidal
75 flats. In 2008, the reef covered 105 ha; *i.e.* 0.42% of the sand flats of the BMSM (Godet et al. 2011)
76 (Figure 1). The BMSM is subjected to an extreme megatidal regime (tidal range up to 15.5 m during
77 spring tides), resulting in large tidal flats - covering 25 000 ha - and mainly dominated by a *Macoma*
78 *balthica* community characterized by low macrobenthic abundances and diversity (Retière 1979;
79 Thorin et al. 2001). The BMSM is an internationally important migration stopover and wintering site
80 for birds (Le Drean-Quenec'hdu et al. 1995), designated as a RAMSAR site and classified as a Special
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.
83 1998). At this site, Dunlin *Calidris alpina*, Red Knot *Calidris canutus*, Oystercatcher *Haematopus*
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
86 2000).

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

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

102 To assess the total number of waders in the whole BMSM, all waders of the site were monitored 5
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
105 associations, Bretagne-Vivante/SEPNB and GONm, was followed (Beaufils et al. 2009). Waders were
106 counted in their high-tide roosts by people equipped with telescopes and binoculars. Because of the
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
110 systematically reported with the exact time and flight direction.

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
113 month, except for November and December), but only while spring tide fully exposed the reef (*i.e.*
114 corresponding to a low tide level of less than 2.5 m above extreme low water spring tide). Birds were
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
118 point, located on a sandbank just outside the study area, ensured visibility of the entire site as well as a
119 minimal bird disturbance.

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

121 In 2010, faeces of five wader species were collected within the reef: Oystercatcher *H. ostralegus*,
122 Dunlin *C. alpina*, Curlew *Numenius arquata*, Grey Plover *P. squatarola* and Bar-tailed Godwit *L.*
123 *lapponica*. We selected these species because they: i) are abundant in the reef; ii) feed regularly on the
124 reef; iii) are species for which faeces are easy to collect because they feed in dense and virtually
125 monospecific groups. Faeces of Curlew and Dunlin were collected on the 27th of April 2010, while
126 faeces of Oystercatcher, Bar-tailed Godwit and Grey Plover were collected on the 13th of August 2010,
127 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
129 confident that a dropping came from the target species. After 20 to 30 min (in order to be sure that the
130 collected faeces resulted from a feeding activity on the reef), the entire droppings were scraped off the
131 surface and preserved in 70% ethanol. In the laboratory, the ethanol was removed by pouring the
132 dropping onto a 20- μ m sieve. For analysis, the sample was transferred to a 100-ml jar, containing a
133 mix of 80% distilled water and 20% hydrogen peroxide (H_2O_2), for a period of at least 24 h. The
134 samples were shaken regularly and, after sedimentation, the supernatant was poured through a 20- μ m
135 sieve. The supernatant remaining on the sieve was transferred to a 25-cm³ petri dish in order to
136 observe and identify lighter animal parts (e.g. polychaete chaetae) with an inverted microscope. A

137 stereo-microscope was used to screen the entire settled sand fraction for hard remnants of bivalves,
138 polychaete jaws, etc. Prior to investigation, we selected for each of the five bird species 15 faecal
139 samples (except for Curlew: 11 samples).

140 Although all different items in a faecal sample were quantified, identification to genus or species level
141 was not always possible. Therefore, each unique unidentified animal part was assigned to a
142 morphotype, leading up to the creation of a catalogue consisting of pictures. Finally, for several
143 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*

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

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

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

163 flats), counting n individuals of a species at high tide in the whole BMSM results in a predicted
164 abundance of $n/238$ individuals of this species on the reef. Only counts which were performed at
165 similar time periods over the entire BMSM and the reef were selected for the analysis; *i.e.* four
166 observation dates (March '09, May '09, September '09, and January '10). The number of days
167 between a count across the BMSM and on the reef varied between 0 and 12 days. G -tests for
168 goodness-of-fit were conducted in order to compare predicted and actual abundances of waders on the
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
173 investigated. Therefore, bird faeces data were transformed into presence/absence data, followed by
174 lumping morphotypes into taxonomic groups. Due to diagnostic features of some morphotypes they
175 could be linked to a certain species. Nonetheless, in most of the cases it was not possible to assign a
176 morphotype to a species but only to an order, class, or even phylum. For every wader species, the
177 frequency of occurrence for each taxon i ($FO_i\%$) was calculated: $FO_i\% = (\text{the number of faecal}$
178 $\text{samples of the wader species } s \text{ where taxon } i \text{ is present} / \text{the total number of faecal samples of the}$
179 $\text{wader species } s) * 100$. Second, to determine which taxa are preferentially found in faecal samples of
180 particular wader species, the proportion of different taxonomic groups ($N_i\%$) per faecal sample was
181 calculated: $N_i\% = (\text{Number of items in taxonomic group } i / \text{total number of items in the faecal sample})$
182 $* 100$. To test whether relative abundances of higher taxonomic groups and the most abundant lower
183 taxonomic groups differed significantly among bird species, a generalized linear model (GLM) was
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
186 overdispersion became apparent in the model output, the model was rerun, taking the overdispersion
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
189 response are not linearly related to each other, the relationship was specified by a log link function.

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

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

209 Across the entire BMSM, 22 wader species were counted, representing a mean density of 1.03 ± 0.58
210 birds ha⁻¹, $N = 5$. The five most abundant species were: Dunlin (42%), Red Knot (18%), Oystercatcher
211 (13%), Grey Plover (9%) and Curlew (7%). On the reef, 15 wader species were counted, representing
212 a mean density of 51.38 ± 19.11 birds ha⁻¹, $N = 5$. Dunlin (39%) was the most abundant species on the
213 reef followed by Red Knot (20%), Grey Plover (14%), Oystercatcher (13%), Bar-tailed Godwit (8%)
214 and Curlew (5%).

215 The one-way global ANOSIM test failed to detect a significant difference in the wader community in
216 the two habitats ($p > 0.05$; $R = 0.108$). Unlike bird composition, bird counts revealed that all wader
217 species together exhibited a high proportion of individuals in the reef in relation to the whole BMSM
218 (Table 1). The frequency of waders in the reef was on average 46.6 times higher than expected
219 assuming a random distribution of waders over the entire BMSM. The same was observed for the five
220 wader species selected for the faecal analysis. Focussing on each of the 5 wader species separately
221 revealed frequencies ranging from 31.1 times (Curlew) up to 112.3 times (Bar-tailed Godwit) higher
222 than expected. Consequently, the observed numbers of these 5 species in the reef were significantly
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
227 much among different bird species (Table 2). Crustaceans were present in all faecal samples of all
228 birds except for Bar-tailed Godwit (FO% = 93.3%). Both polychaetes and bivalves were present in all
229 bird species but polychaetes in a much higher percentage of the faeces (ranging from 72.7% in Curlew
230 to 93.3% in Dunlin) than bivalves (ranging from 13.3% in Oystercatcher and Grey Plover to 45.5% in
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
233 faeces - were encountered in all bird species and reached the highest FO% in Curlew (respectively
234 81.8%, 45.5% and 90.9%). The three most frequently occurring bivalve species among all bird species
235 were *Abra alba*, *C. edule* and *M. balthica*, although they never exceeded a FO% of 6.7%, 36.4%, and
236 20% respectively.

237 Based on the relative abundances of taxonomic groups in each of the 5 bird species a diet composition
238 can be displayed for the 5 bird species examined (Table 3). Relative abundances of all higher
239 taxonomic groups differed significantly ($p < 0.05$) among the 5 wader species (GLM, $p < 0.05$, Table
240 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
242 or Bar-tailed Godwit, while bivalves were eaten more frequently by Dunlin than Curlew. Significant
243 differences in the relative abundances of *L. conchilega* (GLM, $p < 0.0001$) and ‘Other polychaetes’
244 (GLM, $p < 0.0001$) were detected among bird species (Table 5). According to pairwise tests, *L.*
245 *conchilega* was eaten more frequently by Oystercatcher than by Dunlin and Curlew. Additionally,
246 Bar-tailed Godwit ate *L. conchilega* more frequently than Dunlin, but less than Curlew. Within the
247 group of crustaceans, ‘Crabs’ differed significantly among bird species (GLM, $p = 0.005$), although no
248 significant pairwise differences were detected. Lastly, no bivalve species showed a significant
249 difference among waders (GLM, $p > 0.05$).

250 **Discussion**

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

263 However, when abundances are included, the composition of waders clearly differed between the two
264 study sites. The observed frequency of total waders on the *Lanice*-reef was on average 5 799 birds,
265 which is approximately 46.6 times higher than expected assuming a uniform distribution of the birds
266 in the BMSM. Petersen and Exo (1999) observed higher than expected bird abundances in *L.*

267 *conchilega* dominated tidal flats of the Wadden Sea, although densities were only 6 times higher than
268 expected. Similar counting surveys in the Chausey archipelago (France) and preliminary counts in the
269 BMSM already showed the attractiveness of the *L. conchilega* reefs for birds and consequently their
270 potentially important role in the conservation of the avifauna (Godet et al. 2008). The results of the
271 current study prove that within a site of international importance for birds, several wader species are
272 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
275 authors in different study areas (e.g. Zühlke et al. 1998; Zühlke 2001; Callaway 2006; Rabaut et al.
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
281 accommodate a more abundant fauna ($6\,903 \pm 5\,339 \text{ ind.m}^{-2}$) than the average macrobenthic
282 assemblage in the entire BMSM ($1\,568 \pm 299 \text{ ind.m}^{-2}$). As part of a study of Leloup et al. (2008),
283 biomasses of different trophic groups which make up the BMSM were modelled. The biomass of
284 carnivorous and necrophagous macrobenthic fauna in combination with intertidal filter feeders was
285 transformed according to weight-to-weight conversion factors proposed by Ricciardi and Bourget
286 (1998). Comparing the resulting biomass ($2.88 \text{ g AFDM. m}^{-2}$) to the reef biomass ($49.69 \text{ g AFDM.m}^{-2}$)
287 revealed a reef biomass which is more than 17 times higher. Therefore, the reef area can be considered
288 a high productivity area in the bay. Nevertheless, the importance of the whole BMSM for the
289 productivity and functioning of the reef cannot be neglected.

290 Faecal analysis conducted in this study revealed information on the diet composition of waders in the
291 reef. Polychaetes were represented in all bird species and they occurred in a high percentage of the
292 faecal samples, which can probably be attributed to the dominance of polychaetes in the reef benthos.
293 *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
298 and Jago 1993) can lead to a lower accessibility and possibly explains the lower than expected portion
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.*
301 *conchilega* in the reef, might lead to the great contribution of one individual to the nutritional demands
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
306 avoiding the uptake of bivalve shell pieces, as described by Hulscher (1982), and hence the low
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
310 reef benthos were very low (1.77% of the total abundance), particularly for crabs. The observed
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
313 with crustaceans, can have properties which make them particularly perceptible to the avian eye
314 (Goss-Custard 1977).

315 Based on relative abundances of taxonomic groups, bird species that largely dominate the wader
316 community on the reef exhibited significant differences in their dietary composition. Differences in
317 polychaetes, and especially *L. conchilega*, contributed highly to differences in the wader diets.
318 Differences in diets due to bivalves were rather low and in all probability largely overshadowed by an
319 inability to trace back bivalve shells in faeces. For crustaceans, differences in the relative abundance
320 between waders were the result of significant differences in crabs, though no significant pairwise tests

321 were noted. Nevertheless, it can be visually seen that crabs reached much higher abundances in the
322 faeces of Curlew compared to other waders (Table 3), which is in accordance with studies revealing
323 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
329 specifically on *L. conchilega*. This result confirms the investigations of Petersen and Exo (1999),
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
334 of waders feeding in the reef. However, faecal analysis is hampered by a high proportion of
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.
339 Nevertheless, to get a more complete view on a wader's diet, the use of complementary approaches
340 (e.g. visual observations) is recommended (Scheiffarth 2001). In this study morphotypes were
341 counted, which cannot be compared with specimens. Since morphotypes are mostly parts of animals
342 and can belong to one or several specimens of the same species or taxonomic group, caution regarding
343 the interpretation of the results is recommended. Faecal samples of the wader species were collected
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

348 2001). As neither sexual nor seasonal variations in the diet composition of waders were taken into
349 account in our study, comparing wader diets is restricted. Additionally, future studies should try to
350 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
352 fundamental to gain knowledge on the feeding ecology, but is also essential for predicting the effects
353 of a possible loss of the reef habitat in the future. Overall, the *L. conchilega* reef of the BMSM can be
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
356 that birds are attracted by the reef. According to faecal analyses, waders in the *L. conchilega* reef tend
357 to feed on the associated fauna, and especially crustaceans, rather than specifically on *L. conchilega*
358 itself. Within the BMSM, which is characterized by low species diversity and low macrofaunal
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 Beaufiles (Bretagne-Vivante/SEPNB, *La société pour l'Etude et la Protection de la*
364 *Nature en Bretagne*, and GONm, *Groupe Ornithologique Normand*), Sébastien Provost (GONm) and
365 Régis Morel (Bretagne-Vivante/SEPNB) for coordinating and providing the bird counts at the scale of
366 the BMSM for 2009. Both Bretagne-Vivante/SEPNB and the GONm are acknowledged for funding
367 the counts at the scale of the BMSM. Thanks to Thomas Brey for providing and helping with the
368 empirical model to estimate benthic production and to Andrea Moore for help with the faecal analysis.
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
371 University, Belgium. The second author acknowledges the '*Centre National de la Recherche*
372 *Scientifique*' for his Post-Doc Grant.

373 **References**

- 374 Barrett RT, Camphuysen K, Anker-Nilssen T, Chardine JW, Furness RW, Garthe S, Hüppop O,
375 Leopold MF, Montevecchi WA, Veit RR (2007) Diet studies of seabirds: a review and
376 recommendations. *Ices J Mar Sci* 64 (9):1675-1691
- 377 Beaufils M, Morel R, Provost S (2009) Suivi 2009 de la fréquentation de la Baie du Mont Saint-
378 Michel par les oiseaux d'eau. Rapport SEPNEB-GONm
- 379 Braeckman U, Provoost P, Sabbe K, Soetaert K, Middelburg JJ, Vincx M, Vanaverbeke J (2012)
380 Temporal dynamics in the diet of two marine polychaetes as inferred from fatty acid
381 biomarkers. *J Sea Res* 68 (0):6-19. doi:10.1016/j.seares.2011.11.003
- 382 Brey T (1999) A collection of empirical relations for use in ecological modelling. *NAGA The*
383 *ICLARM Quarterly* 22 (3):24-28
- 384 Brey T (2001) Population dynamics in benthic invertebrates. A virtual handbook. [http://www.thomas-](http://www.thomas-brey.de/science/virtualhandbook)
385 [brey.de/science/virtualhandbook](http://www.thomas-brey.de/science/virtualhandbook). Accessed 1st February 2012.
- 386 Brey T, Muller-Wiegmann C, Zittier ZMC, Hagen W (2010) Body composition in aquatic organisms -
387 A global data bank of relationships between mass, elemental composition and energy content.
388 *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
- 391 Callaway R, Desroy N, Dubois SF, Fournier J, Frost M, Godet L, Hendrick VJ, Rabaut M (2010)
392 Ephemeral Bio-engineers or Reef-building Polychaetes: How Stable are Aggregations of the
393 Tube Worm *Lanice conchilega* (Pallas, 1766)? *Integrative and Comparative Biology* 50
394 (2):237-250. doi:10.1093/icb/icq060
- 395 Carey DA (1987) Sedimentological effects and paleoecological implications of the tube-building
396 polychaete *Lanice conchilega* Pallas. *Sedimentology* 34 (1):49-66. doi:10.1111/j.1365-
397 3091.1987.tb00559.x
- 398 Clarke KR, Warwick RM (1994) Change in marine communities: an approach to statistical analysis
399 and interpretation. Primer-E Plymouth Marine Laboratory, UK
- 400 Connor EF, McCoy ED (1979) The Statistics and Biology of the Species-Area Relationship. *Am Nat*
401 113 (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
- 406 Eybert M-C, Geslin T, Questiau S, Feunteun E (2003) Shorebird community variations indicative of a
407 general perturbation in the Mont-Saint-Michel (France). *CR Biologies* 326:140-147
- 408 Feral P (1989) Influence des populations de *Lanice conchilega* (Pallas)(Annelida, Polychaeta) sur la
409 sédimentation sableuse intertidale de deux plages bas-normandes (France). *Bull Soc Geol Fr*
410 8:1193-1200

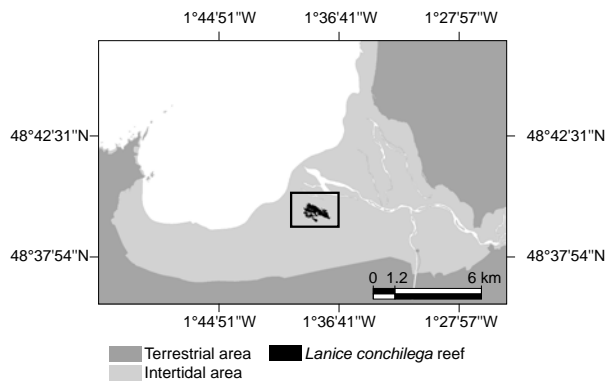
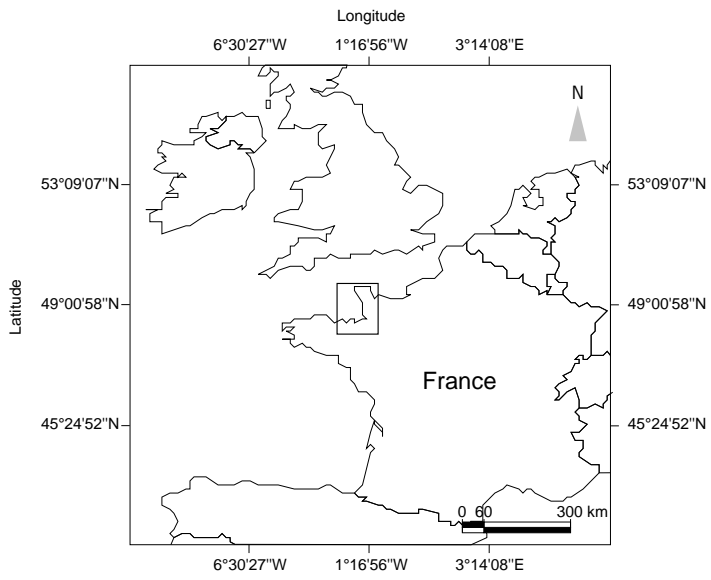
- 411 Godet L, Toupoint N, Olivier F, Fournier J, Retière C (2008) Considering the functional value of
412 common marine species as a conservation stake: The case of sandmason worm *Lanice*
413 *conchilega* (Pallas 1766) (Annelida, Polychaeta) beds. *Ambio* 37 (5):347-355.
414 doi:10.1579/07-a-317.1
- 415 Godet L, Toupoint N, Fournier J, Le Mao P, Retiere C, Olivier F (2009) Clam farmers and
416 Oystercatchers: Effects of the degradation of *Lanice conchilega* beds by shellfish farming on
417 the spatial distribution of shorebirds. *Mar Pollut Bull* 58 (4):589-595.
418 doi:10.1016/j.marpolbul.2008.11.001
- 419 Godet L, Fournier J, Jaffre M, Desroy N (2011) Influence of stability and fragmentation of a worm-
420 reef on benthic macrofauna. *Est Coast Shelf Sci* 92 (3):472-479.
421 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
- 425 Hartmann-Schröder G (1996) Annelida, Borstenwürmer, Polychaeta. In: The fauna of Germany and
426 adjacent seas with their characteristics and ecology, second revised ed., vol. 58. Gustav Fisher,
427 Jena, Germany, 648 p.
- 428 Hulscher JB (1982) The oystercatcher *Haematopus ostralegus* as a predator of the bivalve *Macoma*
429 *balthica* in the dutch Wadden Sea. *Ardea* 70 (2):89-152
- 430 Jones SE, Jago CF (1993) In situ assessment of modification of sediment properties by burrowing
431 invertebrates. *Mar Biol* 115:133-142
- 432 Le Drean-Quenec'hdu S, Mahéo R, Boret P (1995) The Mont Saint-Michel bay: a site of international
433 importance for wintering Palearctic waders. *Wader Study Group Bull* 77:50-54
- 434 Le Drean-Quenec'hdu S, Mahéo R, Boret P (1998) Les limicoles. *Penn ar bed* 169: La Baie du Mont
435 Saint-Michel:1-10
- 436 Leloup FA, Desroy N, Le Mao P, Pauly D, Le Pape O (2008) Interactions between a natural food web,
437 shellfish farming and exotic species: The case of the Bay of Mont Saint Michel (France). *Est*
438 *Coast Shelf Sci* 76 (1):111-120. doi:10.1016/j.ecss.2007.06.011
- 439 Petersen B, Exo KM (1999) Predation of waders and gulls on *Lanice conchilega* tidal flats in the
440 Wadden Sea. *Mar Ecol Prog Ser* 178:229-240. doi:10.3354/meps178229
- 441 Rabaut M, Guilini K, Van Hoey G, Vincx M, Degraer S (2007) A bio-engineered soft-bottom
442 environment: The impact of *Lanice conchilega* on the benthic species-specific densities and
443 community structure. *Est Coast Shelf Sci* 75 (4):525-536. doi:10.1016/j.ecss.2007.05.041
- 444 Rabaut M, Vincx M, Degraer S (2009) Do *Lanice conchilega* (sandmason) aggregations classify as
445 reefs? Quantifying habitat modifying effects. *Helgoland Mar Res* 63 (1):37-46.
446 doi:10.1007/s10152-008-0137-4
- 447 Rabaut M, Van de Moortel L, Vincx M, Degraer S (2010) Biogenic reefs as structuring factor in
448 *Pleuronectes platessa* (Plaice) nursery. *J Sea Res* 64 (1-2):102-106.
449 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
- 454 Rijnsdorp AD, Vingerhoed B (2001) Feeding of plaice *Pleuronectes platessa* L. and sole *Solea solea*
455 (L.) in relation to the effects of bottom trawling. *J Sea Res* 45 (3-4):219-229.
456 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
465 fonctionnement des peuplements benthiques du Golfe Normano-Breton. Dissertation, Muséum
466 National d'Histoire Naturelle, France
- 467 Van Hoey G, Vincx M, Degraer S (2006) Some recommendations for an accurate estimation of *Lanice*
468 *conchilega* density based on tube counts. *Helgoland Mar Res* 60 (4):317-321.
469 doi:10.1007/s10152-006-0041-8
- 470 Van Hoey G, Guilini K, Rabaut M, Vincx M, Degraer S (2008) Ecological implications of the
471 presence of the tube-building polychaete *Lanice conchilega* on soft-bottom benthic
472 ecosystems. *Mar Biol* 154 (6):1009-1019. doi:10.1007/s00227-008-0992-1
- 473 Yates MG, Goss-Custard JD, McGrorty S, Lakhani KH, Durell S, Clarke RT, Rispin WE, Moy I,
474 Yates T, Plant RA, Frost AJ (1993) Sediment characteristics, invertebrate densities and
475 shorebird densities on the inner banks of the Wash. *J Appl Ecol* 30 (4):599-614.
476 doi:10.2307/2404240
- 477 Zühlke R (2001) Polychaete tubes create ephemeral community patterns: *Lanice conchilega* (Pallas,
478 1766) associations studied over six years. *J Sea Res* 46 (3-4):261-272. doi:10.1016/s1385-
479 1101(01)00091-0
- 480 Zühlke R, Blome D, Van Bernem KH, Dittmann S (1998) Effects of the tube-building polychaete
481 *Lanice conchilega* (Pallas) on benthic macrofauna and nematodes in an intertidal sandflat.
482 *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



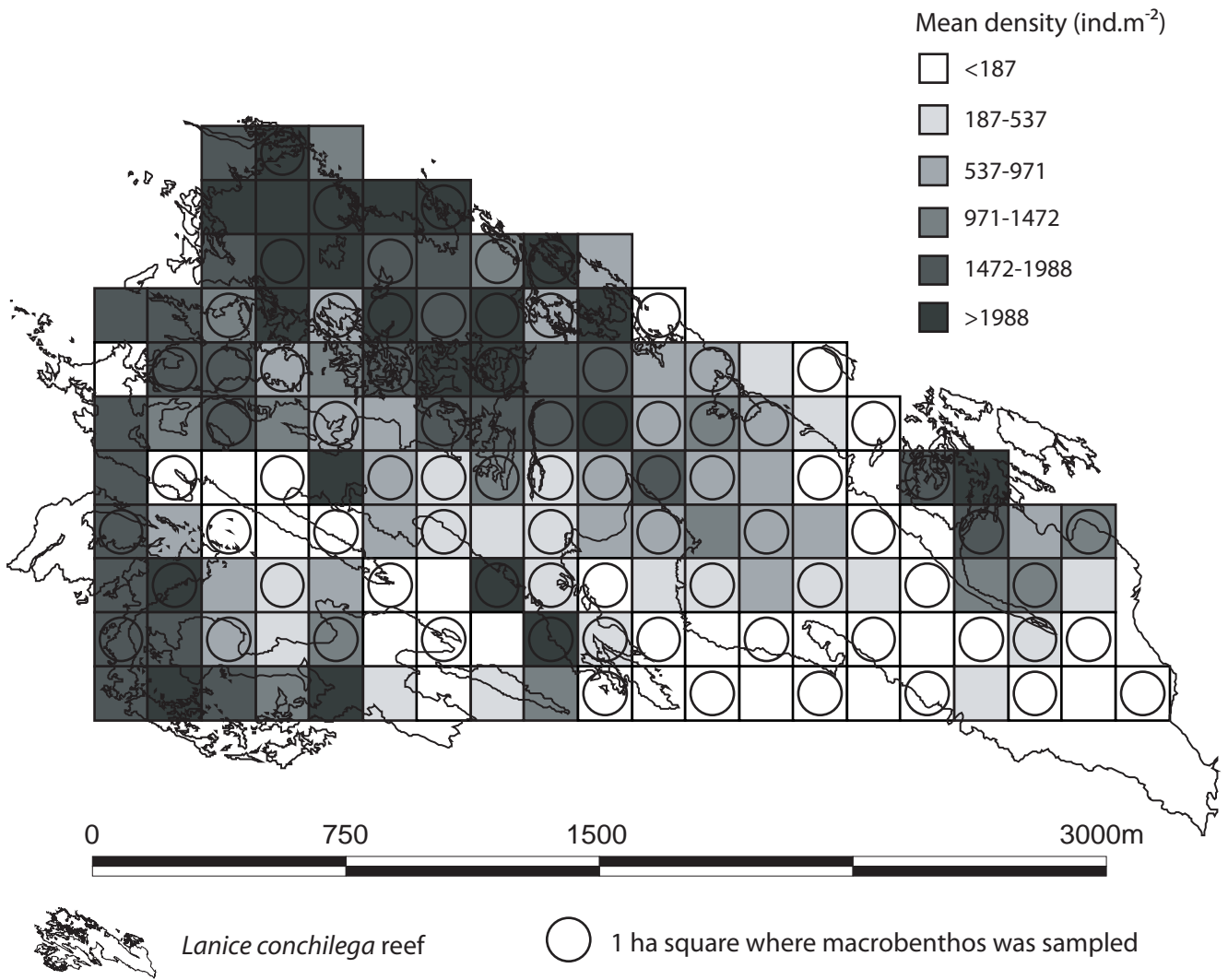


Table 1 Overview of the mean observed and mean expected bird frequencies, and the ratios of these frequencies, for the entire Bay of the Mont Saint-Michel (BMSM) and the *L. conchilega* reef; total G-values, and p-values (G-test) of different groups of birds. Observed frequencies in the BMSM and the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

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

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

		<i>C. alpina</i> (Dunlin) N = 15		<i>H. ostralegus</i> (Oystercatcher) N = 15		<i>N. arquata</i> (Curlew) N = 11		<i>P. squatarola</i> (Grey Plover) N = 15		<i>L. lapponica</i> (Bar-tailed Godwit) N = 15	
Taxon		%	\pm SD	%	\pm SD	%	\pm SD	%	\pm SD	%	\pm SD
Higher taxa	Polychaeta	17.56	17.57	15.15	24.86	18.17	29.61	5.98	9.59	8.93	18.59
	Crustacea	43.31	24.49	26.75	18.16	60.46	35.11	47.26	25.39	18.46	9.65
	Bivalvia	0.23	0.40	0.03	0.10	0.07	0.17	0.01	0.03	0.02	0.05
	Other	38.90	24.57	58.06	26.07	21.30	26.64	46.75	24.63	72.58	21.24
Polychaeta	<i>L. conchilega</i>	6.81	12.19	39.79	32.34	20.10	33.57	18.18	40.45	19.68	36.95
	<i>Nereis</i> sp.	0.48	1.78	0	0	0.02	0.05	0	0	0	0
	<i>E. longa</i>	0	0	0	0	1.01	3.03	0	0	0	0
	Other poly.	92.71	12.03	60.21	32.34	78.88	33.32	81.82	40.45	80.32	36.95
Crustacea	<i>Crab</i> sp.	4.60	5.86	5.12	7.25	32.84	34.00	18.10	18.13	10.06	17.18
	<i>Amphipoda</i> sp.	0.81	1.36	1.30	2.34	6.25	19.06	0.46	0.93	5.69	10.23
	<i>Ostracoda</i> sp.	9.18	15.06	13.36	17.84	15.81	19.30	8.17	22.37	9.15	13.90
	<i>Isopoda</i> sp.	0	0	0	0	0.83	2.74	0	0	0.71	2.67
	<i>Cumacea</i> 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
Bivalvia	<i>A. alba</i>	0	0	50.00	70.71	0	0	50.00	70.71	25.00	50.00
	<i>A. opercularis</i>	0	0	0	0	2.86	6.39	0	0	0	0
	<i>C. edule</i>	0	0	0	0	49.52	47.57	50.00	70.71	0	0
	<i>M. balthica</i>	34.50	44.17	0	0	2.86	6.39	0	0	0	0
	<i>M. bidentata</i>	0	0	0	0	0	0	0	0	25.00	50.00
	<i>Nucula</i> sp.	0	0	50.00	70.71	0	0	0	0	0	0
	<i>S. plana</i>	0	0	0	0	0	0	0	0	25.00	50.00
	<i>S. subtruncata</i>	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 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 - O	D - C	D - G	D - B	O - C	O - G	O - 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 - O	D - C	D - G	D - B	O - C	O - G	O - B	C - G	C - B	G - B
<i>L. conch.</i>	<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