

Notman, Gill ORCID: <https://orcid.org/0000-0002-7531-2709> , McGill, Rona A.R., Hawkins, Stephen J. and Burrows, Michael T. (2016) Macroalgae contribute to the diet of *Patella vulgata* from contrasting conditions of latitude and wave exposure in the UK. *Marine Ecology Progress Series*, 549 . pp. 113-123.

Downloaded from: <https://insight.cumbria.ac.uk/id/eprint/2125/>

***Usage of any items from the University of Cumbria's institutional repository 'Insight' must conform to the following fair usage guidelines.***

Any item and its associated metadata held in the University of Cumbria's institutional repository Insight (unless stated otherwise on the metadata record) may be copied, displayed or performed, and stored in line with the JISC fair dealing guidelines (available [here](#)) for educational and not-for-profit activities

**provided that**

- the authors, title and full bibliographic details of the item are cited clearly when any part of the work is referred to verbally or in the written form
  - a hyperlink/URL to the original Insight record of that item is included in any citations of the work
- the content is not changed in any way
- all files required for usage of the item are kept together with the main item file.

**You may not**

- sell any part of an item
- refer to any part of an item without citation
- amend any item or contextualise it in a way that will impugn the creator's reputation
- remove or alter the copyright statement on an item.

The full policy can be found [here](#).

Alternatively contact the University of Cumbria Repository Editor by emailing [insight@cumbria.ac.uk](mailto:insight@cumbria.ac.uk).

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15

Macroalgae contribute to the diet of *Patella vulgata* from contrasting conditions of latitude  
and wave exposure in the UK

Consumption of macroalgae by *Patella vulgata*

Gillian M. Notman\*<sup>1,4,5</sup>, Rona A. R. McGill<sup>2</sup>, Stephen J. Hawkins<sup>3,5</sup>  
and Michael T. Burrows<sup>4</sup>

<sup>1</sup>Department of Science, Natural Resources and Outdoor Studies, University of Cumbria,  
Ambleside, UK, LA22 1BB (gill.notman@cumbria.ac.uk); <sup>2</sup>NERC Life Sciences Mass  
Spectrometry Facility, Scottish Universities Environmental Research Centre, East Kilbride,  
UK, G75 0QF; <sup>3</sup>Ocean and Earth Science, National Oceanography Centre Southampton,  
University of Southampton, UK, SO14 3ZH; <sup>4</sup> Scottish Association for Marine Science,  
Scottish Marine Institute, Oban, UK, PA37 1QA. <sup>5</sup> Marine Biological Association of the  
UK, The Laboratory, Citadel Hill, Plymouth, UK, PL1 2PB.

16 **Abstract**

17 Analysis of gut contents and stable isotope composition of intertidal limpets (*Patella vulgata*)  
18 showed a major contribution of macroalgae to their diet, along with microalgae and  
19 invertebrates. Specimens were collected in areas with limited access to attached macroalgae,  
20 suggesting a major dietary component of drift algae. Gut contents of 480 animals from two  
21 moderately wave exposed and two sheltered rocky shores in each of two regions: western  
22 Scotland (55-56°N) and southwest England (50°N), were analysed in two years (n = 30 per  
23 site per year). The abundance of microalgae, macroalgae and invertebrates within the guts  
24 was quantified using categorical abundance scales. Gut content composition was compared  
25 among regions and wave exposure conditions; showing that the diet of *P. vulgata* changes  
26 with both wave exposure and latitude. Microalgae were most abundant in limpet gut contents  
27 in animals from southwest sites, whilst leathery/corticated macroalgae were more prevalent  
28 and abundant in limpets from sheltered and northern sites. *P. vulgata* appears to have a more  
29 flexible diet than previously appreciated and these keystone grazers consume not only  
30 microalgae, but also large quantities of macroalgae and small invertebrates. To date, limpet  
31 grazing studies have focussed on their role in controlling recruitment of macroalgae by  
32 feeding on microscopic propagules and germlings. Consumption of adult algae suggests *P.*  
33 *vulgata* may also directly control the biomass of attached macroalgae on the shore, whilst  
34 consumption of drift algae indicates the species may play important roles in coupling subtidal  
35 and intertidal production.

36

## 37 **Introduction**

38           The intertidal limpet, *Patella vulgata*, is the most abundant grazer on rocky shores in  
39 the north east Atlantic and plays an important role in structuring intertidal communities  
40 throughout its range (Hawkins & Hartnoll 1983, Hawkins et al. 1992). Classic limpet  
41 removal experiments (Jones 1948, Lodge 1948, Burrows & Lodge 1950, Southward 1964),  
42 observations following the mass limpet kills in the aftermath of the Torrey Canyon oil spill  
43 clean-up (Southward & Southward 1978, Hawkins & Southward 1992) and limpet exclusion  
44 experiments (Hawkins 1981, Hartnoll & Hawkins 1985, Jenkins et al. 2005, Coleman et al.  
45 2006, Jonnson et al. 2006) have all shown that removal or exclusion of limpets produces lush  
46 growth of canopy forming furoid macroalgae on wave exposed and moderately exposed  
47 rocky shores. These indirect studies have led to the conclusion that *P. vulgata* is a keystone  
48 species in the intertidal, preventing macroalgal growth through microphagous grazing of  
49 epilithic biofilm and associated macroalgal propagules (spores and germlings) (Hill &  
50 Hawkins 1991, Jenkins & Hartnoll 2001, Jenkins et al. 2005, Coleman et al. 2006, Jonnson et  
51 al. 2006, Moore et al. 2007). This view is supported by observational studies of limpet  
52 foraging behaviour (Hartnoll & Wright 1977, Little et al. 1988, Della Santina et al. 1994) and  
53 several studies which directly examined limpet gut contents (Hawkins et al. 1989, Hill &  
54 Hawkins 1991, Little et al. 1990).

55           More recent work suggests that *P. vulgata* may play an even more significant role in  
56 structuring rocky shore ecosystems through direct grazing of mature macroalgae as well as  
57 biofilm and propagules. On many more sheltered shores *P. vulgata* aggregate around  
58 established stands of *Ascophyllum nodosum* and *Fucus vesiculosus* and ‘bite marks’ are  
59 visually apparent on the algal fronds. The fronds are often much shorter than might be  
60 expected, consistent with the ends being directly grazed away by the nearby limpets (Davies  
61 et al. 2007, Davies et al. 2008, personal observation). Lorenzen (2007) also recorded *P.*

62 *vulgata* feeding on stranded macroalgae during periods of emersion, and the results of stable  
63 isotope analyses from several sources provide strong evidence to suggest that macroalgae  
64 may be a more significant source of organic carbon to *P. vulgata* than previously appreciated  
65 (Campbell 2004, Riera et al. 2009, Schaal et al. 2010, Notman 2011).

66 The limited number of studies which have examined gut contents in *P. vulgata* to date  
67 probably give an incomplete view of the dietary range of *P. vulgata*. Sample sizes for foregut  
68 analysis were small and acid digestion methods may have provided biased information on  
69 only a subset of the diet rather than representing whole gut contents (Hawkins et al. 1989,  
70 Hill & Hawkins 1991). This has led to an overemphasis on grazing of the microalgal biofilm  
71 by the species, and a potential underestimation of their capacity for feeding on adult or  
72 detrital macroalgae. Quantification of whole diet composition without such bias is lacking,  
73 and is crucial for a re-evaluation of the role of the species as a keystone grazer across its  
74 range.

75 The overall aim of this study was to compare the diets of limpets between contrasting  
76 latitudinal and wave exposure conditions. Specifically we quantified the relative importance  
77 of microalgal biofilms and other food sources across these locations. We focussed on the diet  
78 of limpets collected from bare rock microhabitats where access to other types of foods was  
79 restricted. Gut contents of adult *P. vulgata* (30-50 mm in length) were examined from wave  
80 exposed and wave sheltered sites in western Scotland and southwest England over two years.  
81 Analysis of carbon ( $\delta^{13}\text{C}$ ) and nitrogen ( $\delta^{15}\text{N}$ ) stable isotope ratios from a subset of these  
82 animals, together with their potential foods, was used to provide additional information on  
83 limpet diets integrated over longer periods of time.

84

## 85 **Materials and Methods**

### 86 **Study sites & sample collection**

87 *Patella vulgata* were collected from mid-shore bare rock microhabitats at four sites in  
88 southwest England, near Plymouth, Devon, and four sites in western Scotland, around Oban  
89 and the Kintyre peninsula. Two moderately wave-exposed (300 – 500 km fetch) and two  
90 sheltered (< 50 km fetch) sites were selected in each region, with the ranges of wave  
91 exposure set using calculated values for wave fetch in accordance with Burrows *et al.* (2008).  
92 Adult limpets were collected from areas of bare rock in spring 2005 and 2006. These habitats  
93 seldom comprised entirely clean rock, but also often contained small areas of macroalgae and  
94 barnacles. To investigate the potential influence of these alternate food sources on diet  
95 composition, we visually estimated the percentage cover of bare rock, barnacle and attached  
96 macroalgal cover in the mid-intertidal using twenty five 0.5 × 0.5 m quadrats at the eight sites  
97 in each year. Limpets were immediately chilled upon collection to minimise digestion of gut  
98 contents, and frozen (-20°C) on return to the laboratory. Thirty animals of similar size (mean  
99 length 42 ± 4 mm) were selected from each site per year for gut contents analysis. A subset  
100 of twelve limpets were randomly selected for stable isotope analysis from the different  
101 locations and wave exposure conditions, and samples of potential foods (biofilm and  
102 macroalgae) were collected at each site in each year. Macroalgal species included  
103 *Ascophyllum nodosum*, *Fucus vesiculosus*, *F. serratus*, *F. spiralis*, *Himanthalia elongata*, and  
104 detrital *L. digitata*.

### 105 **Dissection of gut contents**

106 The digestive organs of *P. vulgata* are long and complex, extending from the mouth  
107 and buccal mass, through the crop and stomach to the long intestine embedded in the visceral  
108 mass (Fretter & Graham, 1962). Limpets were dissected ventrally, whilst partially frozen,  
109 with shells attached. The muscular foot and internal organs were completely removed to

110 reveal the crop and anterior portion of the stomach. Only the anterior sections of the  
111 alimentary canal were excised as the posterior sections are difficult to separate from the  
112 digestive gland and visceral mass.

### 113 **Examination of gut contents**

114 Dissected gut contents were initially examined at  $\times 4$  magnification, and ingested algae  
115 and invertebrates identified and quantified using categorical abundance scales (Table 1). It  
116 was not possible to identify ingested macroalgae to species level. These were therefore  
117 classified according to Steneck and Watling's (1982) functional group scheme (Table 1), with  
118 categories 4 and 5 (corticated and leathery macroalgae) combined due to the difficulties in  
119 objectively and consistently discriminating between these groups in gastropod guts (Raffaelli,  
120 1985, personal observation). The abundance of microalgae (Group 1 including diatoms and  
121 cyanobacteria) was assessed at  $\times 20$  magnification using a compound microscope. A 1 cm<sup>2</sup>  
122 subsample of material was covered with a glass coverslip and five fields of view were  
123 randomly selected for closer examination.

### 124 **Stable isotope analysis**

125 Tissue from the muscular foot of *P. vulgata* was chosen for isotope analysis because  
126 muscle tissues integrate consumer diets over long periods of time (Hobson & Clark 1992)  
127 and, due to their low fat content, do not require lipid extraction (Pinnegar & Polunin 1999).  
128 Dissected foot tissues from 48 of the limpets used in gut contents analysis were rinsed in  
129 deionised water and freeze dried to constant weight for stable isotope analysis.

130 Biofilm was obtained from three 200 cm<sup>2</sup> rock samples (free from macroalgae and  
131 encrusting organisms) from each site in each year. The rocks were soaked in 0.7  $\mu\text{m}$  filtered  
132 seawater for 15 minutes in the laboratory and the rehydrated biofilm removed by brushing  
133 with an electric toothbrush. Biofilm was filtered onto glass fibre filter paper (Whatman  
134 GF/F), frozen at -20°C and freeze dried to constant weight. Samples of corticated/leathery

135 macroalgae were rinsed in deionised water and epibiont-free fronds were frozen at -20°C  
136 before freeze drying. The stable isotope values of barnacles were not measured due to the  
137 difficulties inherent in analysing inorganic carbon (Soreide *et al* 2006).

138 Samples were ground into homogenous powders, weighed into tin capsules (0.7 mg  
139 limpet tissue, 1.5 mg macroalgal tissue and two 9 mm discs of biofilm/filter paper),  
140 randomised and loaded into an automatic carousel for simultaneous analysis of carbon and  
141 nitrogen isotopes using continuous-flow isotope ratio mass spectrometry (CF-IRMS)  
142 (Costech model ECS 4010 elemental analyser coupled with a ThermoFinnegan Delta Plus XP  
143 mass spectrometer). Two laboratory standards were analysed every 10 samples allowing  
144 instrument drift to be corrected if required.

#### 145 **Data analysis**

146 Abundance of algae and invertebrates ingested by limpets was compared between the  
147 two regions, between the two classes of wave exposure and among the sites (nested within  
148 region and exposure), using ordinal logistic regression (OLR, Minitab version 15.1.20,  
149 MINITAB Inc.) on ranked categorical abundance data to determine influences on the  
150 probability of obtaining particular abundance categories. Best OLR models were selected by  
151 comparisons of deviance likelihood ratios following step-wise exclusion of factors from a  
152 saturated model (Notman 2011).

153 Multivariate analyses were used to examine patterns of gut content composition  
154 between the sampling regions and wave exposure conditions. The Gower similarity  
155 coefficient ( $S_{15}$ ) was used to obtain a matrix of similarities for examination of ordinal  
156 categories of limpet gut contents (Gower, 1971; Podani, 1999; Legendre & Legendre, 1998).  
157 Normalisation was not necessary as similar abundance scales were used for each variable.  
158 Non-metric multidimensional scaling (MDS, PRIMER 6, PRIMER-E Ltd.) was used to  
159 produce a two-dimensional ordination of gut contents data from the similarity matrix (Clarke,



160 1993). Similarity indices were used to determine the effects of region and wave exposure on  
161 composition of limpet gut contents (2-way ANOSIM), with the contribution of each taxon to  
162 differences among levels of each factor assessed using similarity percentage analysis  
163 (SIMPER, based on the Euclidian distance measure of association).

164 Consumption of small invertebrates was assessed using counts of ingested items in a  
165 hierarchical cluster analysis. A matrix of similarities of ingested fauna was calculated for  
166 each site using the Bray-Curtis similarity coefficient following square root transformation of  
167 ingested faunal abundance data (Bray & Curtis, 1957) and analyses of similarity (ANOSIM)  
168 were used to assess whether the observed patterns of ingested invertebrates differed  
169 significantly between sampling regions and wave exposure conditions.

170 Stable carbon and nitrogen isotope ratios of biofilm, macroalgae and *P. vulgata* foot  
171 tissue were compared among the two regions, the two classes of wave exposure and the sites  
172 (nested within region and exposure) using analysis of variance. Site was treated as a random  
173 factor, with all other factors fixed and orthogonal. Stable isotope ratios for macroalgae were  
174 calculated for region and exposure type combinations as the mean and standard deviation  
175 across all species of corticated/leathery macroalgae for each site.

176 Habitat composition was evaluated using Mann-Whitney U-tests to compare percentage  
177 cover of bare rock at sampling sites among regions and Wilcoxon signed ranks tests to  
178 evaluate bare rock cover among levels of wave exposure within each region (with Bonferroni  
179 corrections for multiple comparisons) (SPSS version 22.0, IBM Corp).

180

## 181 **Results**

### 182 **Study Sites**

183           The percentage cover of bare rock was similar between moderately exposed sites in  
184 both regions at 92 % (Mann-Whitney U-test,  $p = 0.75$ ). Although bare microhabitats were  
185 deliberately chosen at all sites, the percentage bare rock was lower on sheltered shores,  
186 around 88 % in the north and 84 % in the south, and differed significantly between regions  
187 (Mann-Whitney U-test,  $p < 0.01$ ). Percentage macroalgal and barnacle cover was highest at  
188 southern sheltered shores at around 3 and 13 % respectively ( $p < 0.01$  for all comparisons).

### 189 **Main constituents of limpet gut contents**

190           Most of the 480 animals examined had quantifiable gut contents (82 %). It was not  
191 possible to identify the species of algae ingested, but microalgae (Group 1) and  
192 corticated/leathery algae (Groups 4/5) were present in almost all of these samples: 90 % and  
193 95 % of gut contents respectively. Filamentous (Group 2) algae were found in around 36 %  
194 of gut samples, and foliose (Group 3), articulated calcareous (Group 6) and crustose coralline  
195 (Group 7) algae were only present in around 10 % of the material analysed (Fig. 1).  
196 Invertebrates were present in 91 of the 392 limpet gut contents (23 % overall).

### 197 **Changes in diet with latitude**

198           There were significant differences in limpet diets between the two sampling regions.  
199 Microalgae were almost five times more likely to be recorded in high abundance categories  
200 (common, abundant and superabundant) in gut contents of southern limpets than northern  
201 limpets (ordinal logistic regression region effect, log odds ratio 1.514, SE 0.367,  $p < 0.001$ ,  
202 Table 2, Fig. 2). In contrast, corticated/leathery macroalgae were around twice as likely to be  
203 recorded in higher abundance categories in northern limpets (Fig. 3). The odds ratio of 0.55  
204 given in Table 2 indicates that high abundance categories of corticated/leathery algae were  
205 around half as likely in the south than in the north ( $p < 0.01$ ). Abundance of other types of

206 algae (filamentous, foliose, articulated calcareous or crustose coralline) was not influenced by  
207 latitude ( $p > 0.05$  for all comparisons).

### 208 **Effect of wave exposure on diet**

209 Microalgae were more prevalent in guts of limpets from exposed shores than sheltered  
210 shores in the south (Fig. 2) but not in the north, and no overall trend in the abundance of  
211 microalgae with wave exposure was seen. In the south, higher abundance categories for  
212 microalgae were more likely at the exposed shores at Andurn and Picklecombe (sites SE1 and  
213 SE2) than at the sheltered sites Jennycliff or Cawsand (SS1 or SS2;  $p < 0.01$ , Fig. 2, Table 2).  
214 There were no significant differences in microalgal abundance among the sites in the north,  
215 but there was some evidence that the probability of higher microalgal abundances was greater  
216 in samples from the moderately exposed Easdale (NE1) than for those from the nearby,  
217 sheltered Ellenabeich (NS1; odds ratio = 1.95,  $p = 0.079$ , Table 2).

218 Corticated/leathery algae were almost twice as likely to be at least common (Table 1) in  
219 the gut contents of limpets from sheltered sites than in those from exposed sites (odds ratio =  
220 1.96,  $p < 0.001$ , Table 2 and Fig. 3). Abundance categories of filamentous, foliose,  
221 articulated calcareous or crustose coralline algae were not significantly affected by wave  
222 exposure ( $p > 0.05$  for all comparisons).

### 223 **Ingestion of invertebrates**

224 Small invertebrates were present in 23 % of quantifiable limpet gut contents and were  
225 found at every site (9 to 42 % of limpets examined). Barnacles were the most frequently  
226 ingested taxon, found in 15 % of limpet guts. The small snail *Skeneopsis planorbis* was  
227 ingested by 6 % of limpets, and acarid mites were found in 3% of guts. Other taxa included  
228 bivalves (present in 4 individuals), foraminifera (3), ostracods (3), *Littorina mariae* (2),  
229 copepods (1), hydroids (1), *Melaraphe neritoides* (1), and *Patella vulgata* itself (1). Few  
230 limpets had ingested more than one type of invertebrate (only 4 %). Hierarchical cluster

231 analysis of ingested invertebrates and analyses of similarity indicated that there were no  
232 significant effects of sampling region or wave exposure on the faunal composition of limpet  
233 gut contents (ANOSIM region effect  $r = -0.25$ ,  $p = 0.78$ ; ANOSIM wave exposure effect  $r = -$   
234  $0.25$ ,  $p = 1.0$ ).

### 235 **Patterns of gut content composition in *P. vulgata***

236 Multidimensional ordination of the abundance of the functional groups of algae and  
237 ingested fauna (Fig. 4) also showed that the composition of the diet of *P. vulgata* varied with  
238 region and wave exposure (ANOSIM region effect  $r = 0.017$ ,  $p = 0.025$ ; ANOSIM wave  
239 exposure effect  $r = 0.03$ ,  $p = 0.001$ ). Although no clear, discrete groupings of diet types are  
240 immediately apparent from the ordination plot, there is a general pattern for individuals from  
241 exposed sites in the south to be positioned towards the bottom right of the figure, and  
242 individuals from sheltered sites in both regions to be positioned towards the top left. Samples  
243 from northern exposed sites show less pronounced separation than other samples and overlap  
244 considerably with other data. The MDS plot (Fig. 4) places individual limpets with high  
245 abundance categories for microalgae (Fig. 4c) towards the right of the figure, and shows them  
246 to be mainly from exposed and southern sites. Individuals with high abundance categories  
247 for corticated/leathery algae (Fig. 4f) are positioned towards the top left of the MDS plot and  
248 tend to be from sheltered sites in the north. Limpets which had ingested invertebrate fauna  
249 are positioned towards the bottom of the ordination (Fig. 4b), and show no clear patterns  
250 according to wave exposure conditions or sampling region. The few individuals with high  
251 abundance categories for filamentous, foliose, articulated calcareous and encrusting coralline  
252 algae (Groups 2, 3, 6 and 7; Figs. 4d, e, g and h respectively) appear as outliers in the MDS  
253 plot and show no significant patterns according to the factors examined.

254 SIMPER analyses indicate that microalgae, corticated/leathery and filamentous algae  
255 contribute most to differences in diet composition by region and wave exposure (Table 3).

256 Average abundance categories for microalgae and filamentous algae were higher for samples  
257 from southern sites and average abundance categories for corticated/leathery algae were  
258 higher for limpet gut contents from northern and sheltered sites. Higher abundance  
259 categories of microalgae (Group 1 algae) were found in samples from wave exposed sites  
260 than in those from sheltered areas.

## 261 **Stable isotopes**

262 Analysis of variance of the  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  values in *P. vulgata* foot tissues by region,  
263 wave exposure and site (nested within region and exposure) showed a significant effect of  
264 region on carbon isotope composition and significant effects of site on both carbon and  
265 nitrogen isotope ratios (Table 4, Fig. 5). The average  $\delta^{13}\text{C}$  values of animals from the south  
266 were around 2 ‰ enriched in  $^{13}\text{C}$  compared to animals from the north ( $p < 0.01$ ). Variability  
267 in  $\delta^{15}\text{N}$  values of limpet tissues was largely due to differences between sites ( $p < 0.01$ ) and,  
268 although nitrogen isotope values in limpets from exposed sites in the north were slightly  
269 higher than those of sheltered sites in this region, this trend was not significant ( $p > 0.05$ ).  
270 Average  $\delta^{13}\text{C}$  values for corticated/leathery macroalgae differed significantly between the  
271 sampling regions (-16.4 ‰ in the north and -17.2 to -18.5 ‰ in the south;  $p < 0.01$ ; Fig. 5),  
272 and average  $\delta^{15}\text{N}$  values were lower in the north (5.3 to 6.2 ‰) compared to similar algae  
273 from the south (7.4 to 7.6 ‰). No significant wave exposure effects were seen. The average  
274  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  values of epilithic biofilm were -20.7 ‰ and 6.4 ‰ respectively and no  
275 significant differences among region and wave exposure conditions were shown (Fig. 5).

276

277

278

279

## 280 **Discussion**

281 Our work shows that the intertidal limpet *Patella vulgata* consumes not only  
282 microalgae, but also large quantities of macroalgae and some small invertebrates. Intertidal  
283 consumers are thought to exploit the most common food sources available in their immediate  
284 surroundings (Raffaelli 1985; Steinarsdottir et al. 1995) and *P. vulgata* is a homing species  
285 which tends to forage within 1 m of its 'home scar' (Hartnoll & Wright 1977, Della Santina  
286 et al. 1995). Despite being collected from bare rock habitats, 95 % of the limpets examined  
287 in this study had macroalgae in their guts, and the stable isotope evidence indicates that  
288 assimilated carbon was from macroalgal sources. It is clear that these animals are exploiting  
289 a macroalgal food resource that is not immediately apparent on the shore at low tide. *Patella*  
290 *vulgata* has been observed feeding on drift algae stranded on the shore when exposed at low  
291 water, particularly on dark, damp days and following storms (Lorenzen 2007, GMN, SJH  
292 personal observations). South African limpets (*Patella argenvillei* and *P. granatina*) have  
293 been documented capturing and feeding on drifting kelp directly from the water column  
294 (Bustamante et al. 1995). Our findings suggest that *P. vulgata* may also be able to exploit  
295 drift algae from the water column when the tide is in.

296 The relative abundance of ingested algal material in limpet guts is affected by both  
297 latitude and wave exposure; microalgae are more common in the diet of limpets from  
298 southern and moderately wave exposed sites, whereas corticated/leathery macroalgae are  
299 more frequently encountered in animals from northern and sheltered coastlines. Stable  
300 isotope analyses show that foot tissues from limpets from all sites are enriched in  $^{13}\text{C}$  relative  
301 to biofilm, also indicating that the organic carbon assimilated over time is largely derived  
302 from macroalgal sources rather than microalgae.

303 *Patella vulgata* plays a major role in structuring intertidal communities along  
304 gradients of wave exposure and latitude in the north-east Atlantic by ingestion of biofilm and

305 macroalgal propagules (Hawkins & Hartnoll 1983, Hawkins et al. 1992) and previous gut  
306 contents analyses have supported this (Hawkins et al. 1989, Hill & Hawkins 1991). Ingestion  
307 of invertebrates was reported in these studies at similar levels to those recorded here, but acid  
308 digestion probably removed much of the macroalgal tissues from the analyses and the  
309 presence of any residual macroalgal tissue (*Fucus* spp.) within the gut was interpreted as  
310 consumption of juvenile algae, rather than sections of mature thalli (Hill & Hawkins 1991).  
311 It is difficult dissect the limpet alimentary tract, and even harder to identify fragments of  
312 algae in limpet gut contents (Raffaelli 1985, C. Maggs personal communication). Yet our  
313 examination of gut contents, combined with stable carbon and nitrogen isotope analysis,  
314 provides very strong evidence that *P. vulgata* routinely consumes mature macroalgae as well  
315 as biofilm and macroalgal propagules. Use of live and detrital macroalgal foods may explain  
316 the lack of clear relationships between biofilm standing stock, limpet density and grazing  
317 activity which have been observed on a number of British shores (Jenkins et al. 2001,  
318 Thompson et al. 2004).

319         Some of the ingested macroalgae in this study may be from *P. vulgata* directly  
320 consuming adult furoid plants. In sheltered sea lochs and bays, where intertidal communities  
321 are dominated by leathery, furoid macroalgae, *P. vulgata* may be directly feeding on the  
322 fronds. On more exposed areas, headlands and breakwaters, where such algae are rare and  
323 communities are characterised by barnacles and mussels (Lewis, 1964; Ballantine, 1961),  
324 drift algae may enhance populations of these grazers and so maintain high grazing intensity  
325 and prevent further escapes of furoids (Moore et al 2007). The greater availability of drift  
326 algae in the north may explain the greater incidence of macroalgae in the diet of *P. vulgata*,  
327 given than macroalgal cover increases with latitude. Furoids dominate most shores in  
328 Norway whereas in northern Spain the algae are confined to only the most sheltered sites  
329 (Ballantine 1961, Hawkins & Hartnoll 1983, Hawkins et al. 1992, Coleman et al. 2006).

330 By feeding on allochthonous drift algae, *P. vulgata* may be coupling subtidal and  
331 intertidal production through feeding. Thus their ecological role may be much more  
332 extensive than simply that of a microphagous grazer of rocky substrata. Such grazers may  
333 therefore play crucial roles in facilitating nutrient flow in coastal environments and allowing  
334 horizontal transport of resources between exposed and sheltered areas (Polis et al 1997).

335 Consumption of corticated/leathery algae is reported in many of the early accounts of  
336 limpet biology (see Steneck and Watling (1982) for a comprehensive review), and is also  
337 described in several more recent investigations (Davies et al., 2008; Davies et al., 2007;  
338 Lorenzen, 2007). Lorenzen (2007) provides photographic evidence of rasping marks made  
339 by *P. vulgata* on the thalli of *A. nodosum* from shores in Brittany and a video showing a low  
340 shore individual feeding on attached *Laminaria digitata* was recorded by G. Notman in  
341 Argyll in 2007 (<http://www.youtube.com/watch?v=79RvGRUdnwE>). Stable isotope  
342 evidence from this and other studies also indicates that fucoid macroalgae are a significant  
343 source of nutrition to *P. vulgata*; the limpets are not only ingesting macroalgal foods, but they  
344 are also assimilating them into their tissues (Campbell, 2004; Notman et al. in preparation;  
345 Riera et al., 2009; Schaal et al., 2010).

346 Despite limiting our investigation to relatively large limpets collected from open rock  
347 habitats, and acknowledging that the diet of these molluscs may change seasonally and  
348 ontogenetically, as well as in response to food availability, it is clear that the diet and the  
349 ecological role of *P. vulgata* is not yet fully understood. *Patella vulgata* is considered to be a  
350 keystone species on temperate rocky shores because its grazing activities prevent  
351 establishment of mature algae by consumption of macroalgal propagules and germlings in the  
352 biofilm. The assumption was that biofilm foods provided the main source of energy to these  
353 animals. Our work suggests that the species may also play important roles in modifying  
354 macroalgal cover, especially on more sheltered coastlines where fucoid algae commonly



355 occur. Moreover, previous work has interpreted the aggregation of *P. vulgata* under stands of  
356 mature algae as being primarily a sheltering response, mitigating desiccation stress and  
357 reducing predation (Hartnoll & Hawkins 1985, Coleman et al. 1999, Moore et al. 2007). This  
358 work indicates that limpets associated with patches of attached macroalgae are likely to be  
359 feeding on them too. The species may also be of great ecological importance in terms of  
360 coupling sub- and intertidal production by ingesting allochthonous drift algae across a range  
361 of wave exposure conditions and latitudes. This would help to explain the high biomass of  
362 grazers which occurs on apparently bare shores across the north east Atlantic, which is  
363 unlikely to be supported by epilithic microalgal production alone.

364

## LITERATURE CITED

- 365 Ballantine WJ (1961) A biologically defined exposure scale for the comparative description  
366 of rocky shores. *Field Studies* 1:1-19
- 367 Bray JR, Curtis JT (1957) An ordination of the upland forest communities of Southern  
368 Wisconsin. *Ecol Monogr* 27:325-349
- 369 Burrows EM, Lodge SM (1950) Note on the interrelationships of *Patella*, *Balanus* and *Fucus*  
370 on a semi-exposed coast. *Rep Mar Biol St, Port Erin* 62:30-34
- 371 Burrows MT, Harvey R, Robb L (2008) Wave exposure indices from digital coastlines and  
372 the prediction of rocky shore community structure. *Mar Ecol Prog Ser* 353:1-12
- 373 Bustamante RH, Branch GM, Eekhout S (1995) Maintenance of an exceptional intertidal  
374 grazer biomass in South Africa - subsidy by subtidal kelps. *Ecology* 76:2314-2329
- 375 Campbell EM (2004) Nitrogen assimilation by limpets, PhD Dissertation, University of  
376 Glasgow, Glasgow
- 377 Clarke KR (1993) Nonparametric multivariate analyses of changes in community structure.  
378 *Aust J Ecol* 18:117-143
- 379 Coleman RA, Goss-Custard JD, Durell SELV, Hawkins SJ (1999). Limpet *Patella* spp.  
380 consumption by oystercatchers *Haematopus ostralegus*: a preference for solitary prey  
381 items. *Mar Ecol Prog Ser* 183: 253-261
- 382 Coleman RA, Underwood AJ, Benedetti-Cecchi L, Aberg P, Arenas F, Arrontes J, Castro J,  
383 Hartnoll RG, Jenkins SR, Paula J, and others (2006) A continental scale evaluation of  
384 the role of limpet grazing on rocky shores. *Oecologia* 147:556-564
- 385 Davies AJ, Johnson MP, Maggs CA (2007) Limpet grazing and loss of *Ascophyllum*  
386 *nodosum* canopies on decadal time scales. *Mar Ecol Prog Ser* 339:131-141
- 387 Davies AJ, Johnson MP, Maggs CA (2008) Subsidy by *Ascophyllum nodosum* increases  
388 growth rate and survivorship of *Patella vulgata*. *Mar Ecol Prog Ser* 366:43-48
- 389 Della Santina P, Naylor E, Chelazzi G (1994) Long-term field actography to assess the timing  
390 of foraging excursions in the limpet *Patella vulgata* L. *J Exp Mar Biol Ecol* 178:193-203
- 391 Della Santina P, Santini G, Chelazzi G (1995) Factors affecting variability of foraging  
392 excursions in a population of the limpet *Patella vulgata* (mollusca, gastropoda). *Mar*  
393 *Biol* 122:265-270
- 394 Fretter V and Graham A (1962) British prosobranch mollusca, their functional anatomy and  
395 ecology. Ray Society, London
- 396 Gower JC (1971) A general coefficient of similarity and some of its properties. *Biometrics*  
397 27:857-871

- 398 Hartnoll RG and Hawkins SJ (1985) Patchiness and fluctuations on moderately exposed  
399 rocky shores. *Ophelia* 24:53-63
- 400 Hartnoll RG, Wright JR (1977) Foraging movements and homing in the limpet *Patella*  
401 *vulgata* L. *Anim Behav* 25: 806-810.
- 402 Hawkins SJ (1981) The influence of patella grazing on the furoid-barnacle mosaic on  
403 moderately exposed rocky shores. *Kieler Meeresforsch* 5:537-543
- 404 Hawkins SJ and Hartnoll RG (1983) Grazing of intertidal algae by marine-invertebrates.  
405 *Oceanogr Mar Biol* 21:195-282
- 406 Hawkins SJ and Southward A (1992) The Torrey Canyon oil spill: Recovery of rocky shore  
407 communities. In: *Restoring the Nation's Marine Environment*:583-619
- 408 Hawkins SJ, Hartnoll RG, Kain JM, Norton TA (1992) Plant-animal interactions on hard  
409 substrata in the north-west Atlantic. In: John DM, Hawkins SJ and Price JH (eds) *Plant-*  
410 *Animal Interactions in the Marine Benthos*. Clarendon Press, Oxford, p. 1-32
- 411 Hawkins SJ, Watson DC, Hill AS, Harding SP, Kyriakides MA, Hutchinson S, Norton TA  
412 (1989) A comparison of feeding mechanisms in microphagous, herbivorous, intertidal,  
413 prosobranchs in relation to resource partitioning. *J Mollus Stud* 55:151-165
- 414 Hill AS and Hawkins SJ (1991) Seasonal and spatial variation of epilithic microalgal  
415 distribution and abundance and its ingestion by *Patella vulgata* on a moderately exposed  
416 rocky shore. *J Mar Biol Assoc UK* 71:403-423
- 417 Hobson KA and Clark RG (1992) Assessing avian diets using stable isotopes .1. turnover of  
418 C-13 in tissues. *Condor* 94:181-188
- 419 Jenkins SR and Hartnoll RG (2001) Food supply, grazing activity and growth rate in the  
420 limpet *patella vulgata* L.: A comparison between exposed and sheltered shores. *J Exp*  
421 *Mar Biol Ecol* 258:123-139
- 422 Jenkins SR, Arenas F, Arrontes J, Bussell J, Castro J, Coleman RA, Hawkins SJ, Kay S,  
423 Martinez B, Oliveros J, and others (2001) European-scale analysis of seasonal variability  
424 in limpet grazing activity and microalgal abundance. *Mar Ecol Prog Ser* 211:193-203
- 425 Jenkins SR, Coleman RA, Della Santina P, Hawkins SJ, Burrows MT, Hartnoll RG (2005)  
426 Regional scale differences in the determinism of grazing effects in the rocky intertidal.  
427 *Mar Ecol Prog Ser* 287:77-86
- 428 Johnson MP, Hanley ME, Frost NJ, Mosley MWJ, Hawkins SJ (2008) The persistent spatial  
429 patchiness of limpet grazing. *J Exp Mar Biol Ecol* 365:136-141
- 430 Jones NS (1948) Observations and experiments on the biology of *Patella vulgata* at Port St.  
431 Mary, Isle of Man. *Trans Proc Liverpool Biol Soc* 56: 60-77

- 432 Jonnson PR, Granhag L, Moschella PS, Aberg P, Hawkins SJ, Thompson RC (2006)  
 433 Interactions between wave action and grazing control the distribution of intertidal  
 434 macroalgae. *Ecology* 87: 1169-1178
- 435 Legendre P and Legendre L (1998) Numerical ecology. In: Developments in environmental  
 436 modelling, 20, Second English. Elsevier Science B. V., Amsterdam
- 437 Lewis JR (1964) The ecology of rocky shores. English Universities Press, London
- 438 Little C, Morrith D, Paterson DM, Stirling P, Williams GA (1990) Preliminary observations  
 439 on factors affecting foraging activity in the limpet *Patella vulgata*. *J Mar Biol Assoc UK*  
 440 70:181-195
- 441 Little C, Williams GA, Morrith D, Perrins JM, Stirling P (1988) Foraging behaviour of  
 442 *Patella vulgata* L. in an Irish sea-lough. *J Exp Mar Biol Ecol* 120:1-21
- 443 Lodge SM (1948) Algal growth in the absence of *Patella* on an experimental strip of  
 444 foreshore, Port St. Mary, Isle of Man. *Trans Proc Liverpool Biol Soc* 56:78-85
- 445 Lorenzen S (2007) The limpet *Patella vulgata* L. at night in air: Effective feeding on  
 446 *Ascophyllum nodosum* monocultures and stranded seaweeds. *J Mollus Stud* 73:267-274
- 447 Moore P, Thompson RC, Hawkins SJ (2007) Effects of grazer identity on the probability of  
 448 escapes by a canopy-forming macroalga. *J Exp Mar Biol Ecol* 344:170-180
- 449 Notman GM (2011) A comparison of the trophic ecology of grazing gastropods on the rocky  
 450 shores of northern and southern Britain, PhD Dissertation, University of Aberdeen,  
 451 Aberdeen
- 452 Peterson BJ and Fry B (1987) Stable isotopes in ecosystem studies. *Annu Rev Ecol Syst*  
 453 18:293-320
- 454 Pinnegar JK and Polunin NVC (1999) Differential fractionation of delta C-13 and delta N-15  
 455 among fish tissues: Implications for the study of trophic interactions. *Funct Ecol* 13:225-  
 456 231
- 457 Podani J (1999) Extending Gower's general coefficient of similarity to ordinal characters.  
 458 *Taxon* 48:331-340
- 459 Polis GA, Anderson WB, Holt RD (1997) Toward an integration of landscape and food web  
 460 ecology: The dynamics of spatially subsidized food webs. *Annu Rev Ecol Syst*:289-316
- 461 Raffaelli D (1985) Functional feeding groups of some intertidal mollusks defined by gut  
 462 contents analysis. *J Mollus Stud* 51:233-239
- 463 Riera P, Escaravage C, Leroux C (2009) Trophic ecology of the rocky shore community  
 464 associated with the *Ascophyllum nodosum* zone (Roscoff, France): A delta C-13 vs delta  
 465 N-15 investigation. *Estuar Coast Shelf S* 81:143-148

- 466 Schaal G, Riera P, Leroux C (2010) Trophic ecology in a northern Brittany (Batz Island,  
467 France) kelp (*Laminaria digitata*) forest, as investigated through stable isotopes and  
468 chemical assays. *J Sea Res* 63:24-35
- 469 Soreide JE, Tamelander T, Hop H, Hobson KA, Johansen I (2006) Sample preparation effects  
470 on stable C and N isotope values: A comparison of methods in arctic marine food web  
471 studies. *Mar Ecol Prog Ser* 328:17-28
- 472 Southward A (1964) Limpet grazing and the control of vegetation on rocky shores. In: Crisp  
473 DJ (ed) *Grazing in terrestrial and marine environments*. Blackwell Scientific  
474 Publications, Oxford, p. 296-373
- 475 Southward AJ and Southward EC (1978) Recolonisation of rocky shores in Cornwall after  
476 use of toxic dispersants to clean up the Torrey Canyon spill. *J Fish Res Board Can*  
477 35:682-706
- 478 Steinarsdottir MB, Ingolfsson A, Olafsson E (2009) Trophic relationships on a furoid shore  
479 in south-western Iceland as revealed by stable isotope analyses, laboratory experiments,  
480 field observations and gut analyses. *J Sea Res* 61:206-215
- 481 Steneck RS and Watling L (1982) Feeding capabilities and limitation of herbivorous  
482 mollusks - a functional-group approach. *Mar Biol* 68:299-319
- 483 Thompson RC, Roberts MF, Norton TA, Hawkins SJ (2000) Feast or famine for intertidal  
484 grazing molluscs: A mis-match between seasonal variations in grazing intensity and the  
485 abundance of microbial resources. *Hydrobiologia* 440:357-367

486  
487  
488

Table 1. *Patella vulgata* gut contents. Taxa identified and abundance categories used for gut contents analysis. Algal functional groups follow the classification by Steneck and Watling (1982).

Group	Algal Type /Taxon	Representatives	Measure	S Super- abundant	A Abundant	C Common	F Frequent	O Occasional	R Rare	N None
1	Microalgae	Diatoms, cyanobacteria	No. cells per field of view	> 50	21 to 50	11 to 20	6 to 10	1 to 5	0.5 to 1	< 0.2
2	Filamentous Algae	<i>Cladophora</i> , <i>Ectocarpus</i>	No. of fragments	> 50	31 to 50	21 to 30	11 to 20	6 to 10	1 to 5	None
3	Foliose Algae	<i>Ulva</i> , <i>Porphyra</i>	No. of fragments	> 50	31 to 50	21 to 30	11 to 20	6 to 10	1 to 5	None
4	Corticated Macrophytes	<i>Bryothamnium</i> , <i>Chondria</i> /	No. of fragments	> 50	31 to 50	21 to 30	11 to 20	6 to 10	1 to 5	None
5	Leathery Macrophytes	<i>Laminaria</i> , <i>Fucus</i> , <i>Ascophyllum</i>	No. of fragments	> 50	31 to 50	21 to 30	11 to 20	6 to 10	1 to 5	None
6	Articulated Calcareous Algae	<i>Corallina</i> , <i>Halimeda</i>	No. of fragments	> 50	31 to 50	21 to 30	11 to 20	6 to 10	1 to 5	None
7	Crustose Coralline Algae	Lithothamnion, <i>Lithophyllum</i>	No. of fragments	> 50	31 to 50	21 to 30	11 to 20	6 to 10	1 to 5	None
Fauna	Ingested Invertebrates	Barnacle cyprids, <i>Skeneopsis planorbis</i> , acarinids	No. of individuals	> 50	31 to 50	21 to 30	11 to 20	6 to 10	1 to 5	None

489 Table 2. *Patella vulgata* gut contents. Comparison of abundance of microalgae and  
 490 corticated/leathery macroalgae across regions, sites and wave exposure by ordinal logistic  
 491 regression. Site type abbreviations: NE, northern exposed; NS, northern sheltered; SE,  
 492 southern exposed; SS, southern sheltered. (\*\*\*p < 0.001; \*\*p < 0.01). Results are shown for  
 493 best fit models (see text).  
 494

Region or Site code (region)	Loge odds ratio	SE	z	P	Odds ratio	95 % confidence interval upper lower	
<i>Microalgae: Group 1</i>							
North vs South	1.514	0.367	4.13	< 0.001***	4.54	2.22	9.32
NE1 Easdale							
NE2 Putechan	0.016	0.400	0.04	0.968	1.02	0.46	2.23
NS1 Ellenabeich	0.666	0.379	1.75	0.079	1.95	0.93	4.09
NS2 Luing	0.514	0.369	1.39	0.164	1.67	0.81	3.45
SE1 Andurn							
SE2 Picklecombe	0.429	0.348	1.23	0.217	1.54	0.78	3.04
SS1 Jennycliff	1.150	0.345	3.34	0.001**	3.16	1.61	6.21
SS2 Cawsand	1.654	0.347	4.77	< 0.001***	5.23	2.65	10.32
<i>Corticated/leathery algae: Group 4/5</i>							
North vs South	-0.606	0.184	-3.29	0.001**	0.55	0.38	0.78
Exposed vs Sheltered	0.672	0.184	3.64	< 0.001***	1.96	1.36	2.81

495

496 Table 3. *Patella vulgata* diet composition. Comparison of gut content composition between  
 497 regions and wave exposures by SIMPER analysis using the Euclidian distance measure of  
 498 association. Average abundance categories of algae are shown (categories N to S converted  
 499 to numerical integers 0 to 7) with average squared Euclidian distance (D) and percentage  
 500 contribution (C %) to differences between regions and wave exposures.  
 501

Algal Group	Region				Wave exposure			
	North	South	D	C %	Exposed	Sheltered	D	C %
1 Microalgae	1.90	2.76	4.65	29	2.75	2.04	4.75	29
4/5 Corticated/Leathery	2.64	2.22	4.17	26	2.09	2.69	4.32	26
2 Filamentous	0.608	0.944	3.76	24	0.643	0.928	3.72	23
7 Encrusting coralline	0.08	0.315	1.11	7	0.346	0.087	1.34	8

502

503 Table 4. ANOVA testing for differences in  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  values of *P. vulgata* foot tissue by  
 504 region, wave exposure and site (nested within region and exposure). x denotes that the  
 505 analysis could not perform an exact F test. (\*\* p < 0.01) n = 12 per site.

Source	df	$\delta^{13}\text{C}$			$\delta^{15}\text{N}$		
		MS	F	p	MS	F	p
Region	1	57.602	24.83	0.008** x	7.157	4.28	0.107 x
Exposure	1	1.172	0.51	0.517 x	3.934	2.36	0.200 x
Site (Reg Exp)	4	2.323	5.89	0.001**	1.674	4.47	0.004**
Region*Exposure	1	0.029	0.01	0.917 x	10.649	6.37	0.065 x
Error	44	0.394			0.374		
Levene's Test		1.22, p = 0.315			0.94, p = 0.491		
Bartlett's Test		5.89, p = 0.553			10.52, p = 0.161		
Variance							
Homogeneous		Yes			Yes		

506

507

508

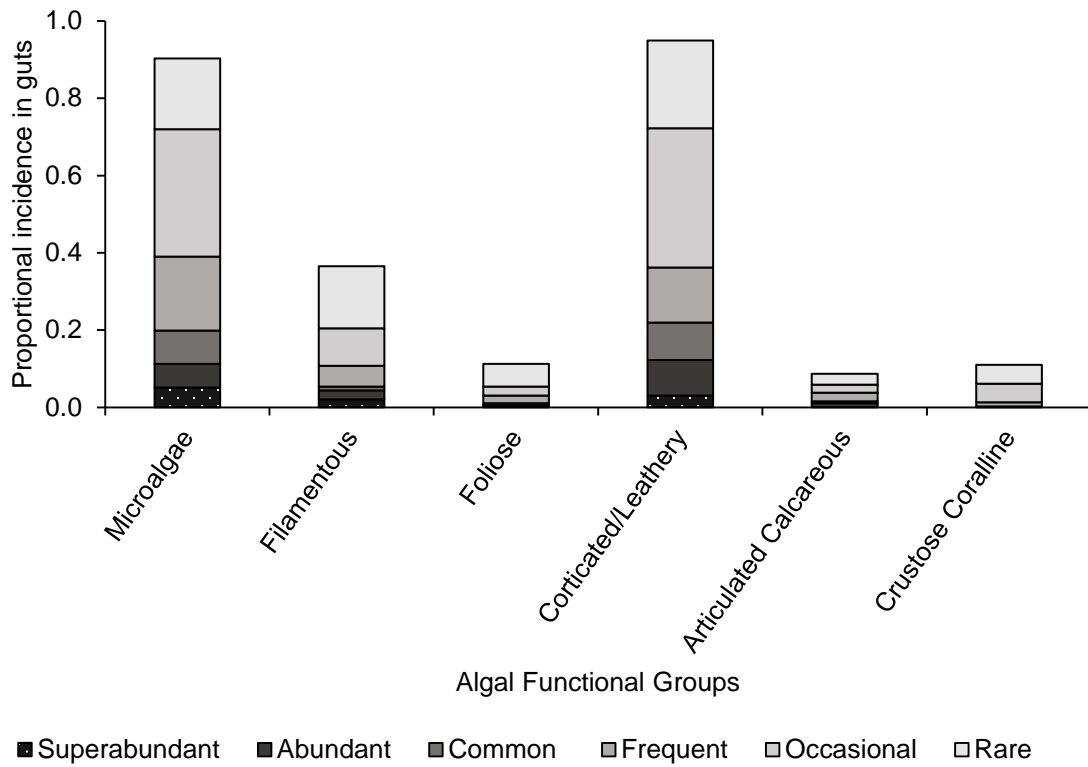
509

510

511

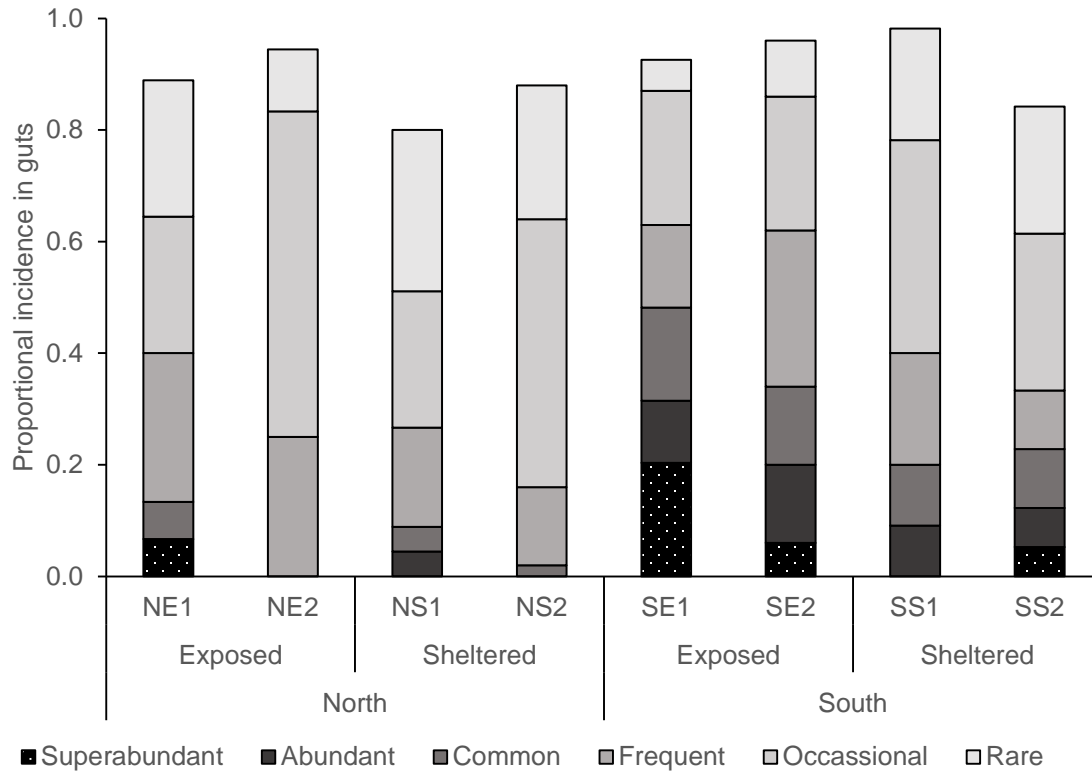
512





514

515 Fig. 1. *Patella vulgata* gut contents (n = 392). Abundance of functional groups of algae in  
 516 gut contents of *P. vulgata* shown by cumulative proportional incidence in abundance  
 517 categories (defined in Table 1).

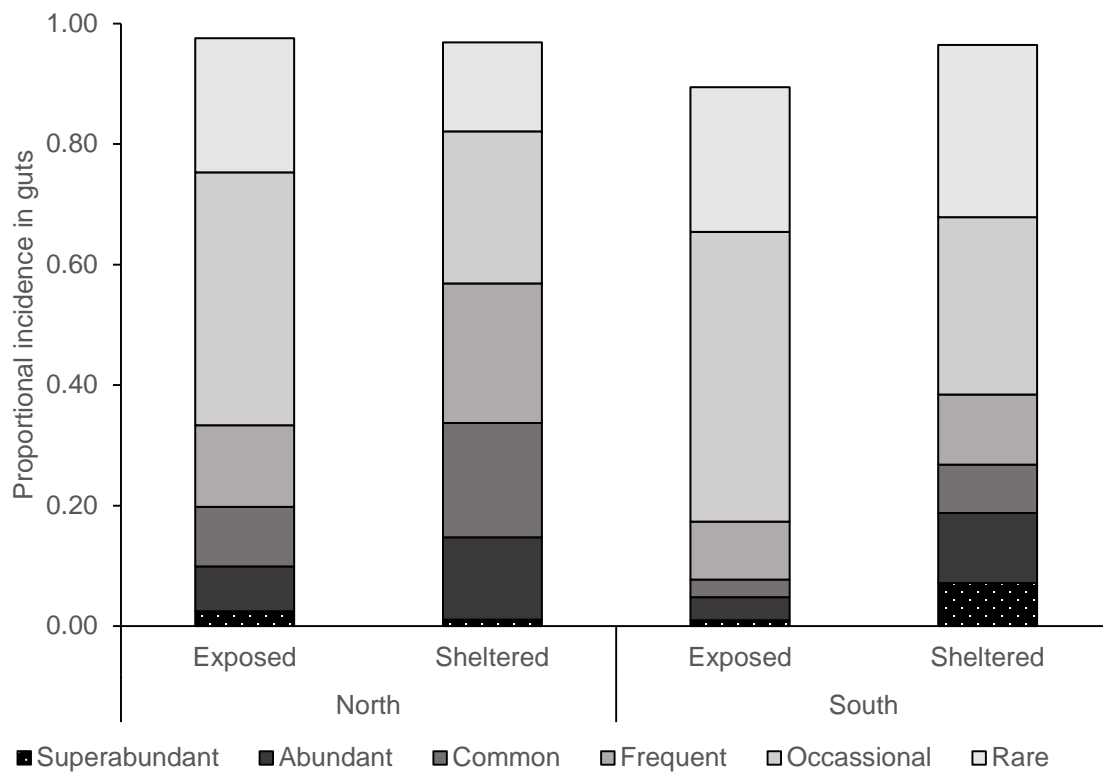


518

519 Fig. 2. *Patella vulgata* gut contents (n = 392). Abundance of microalgae by proportional  
 520 incidence in abundance categories (Table 1) by region, wave exposure and site. Site  
 521 abbreviations are given in Table 2.

522

523

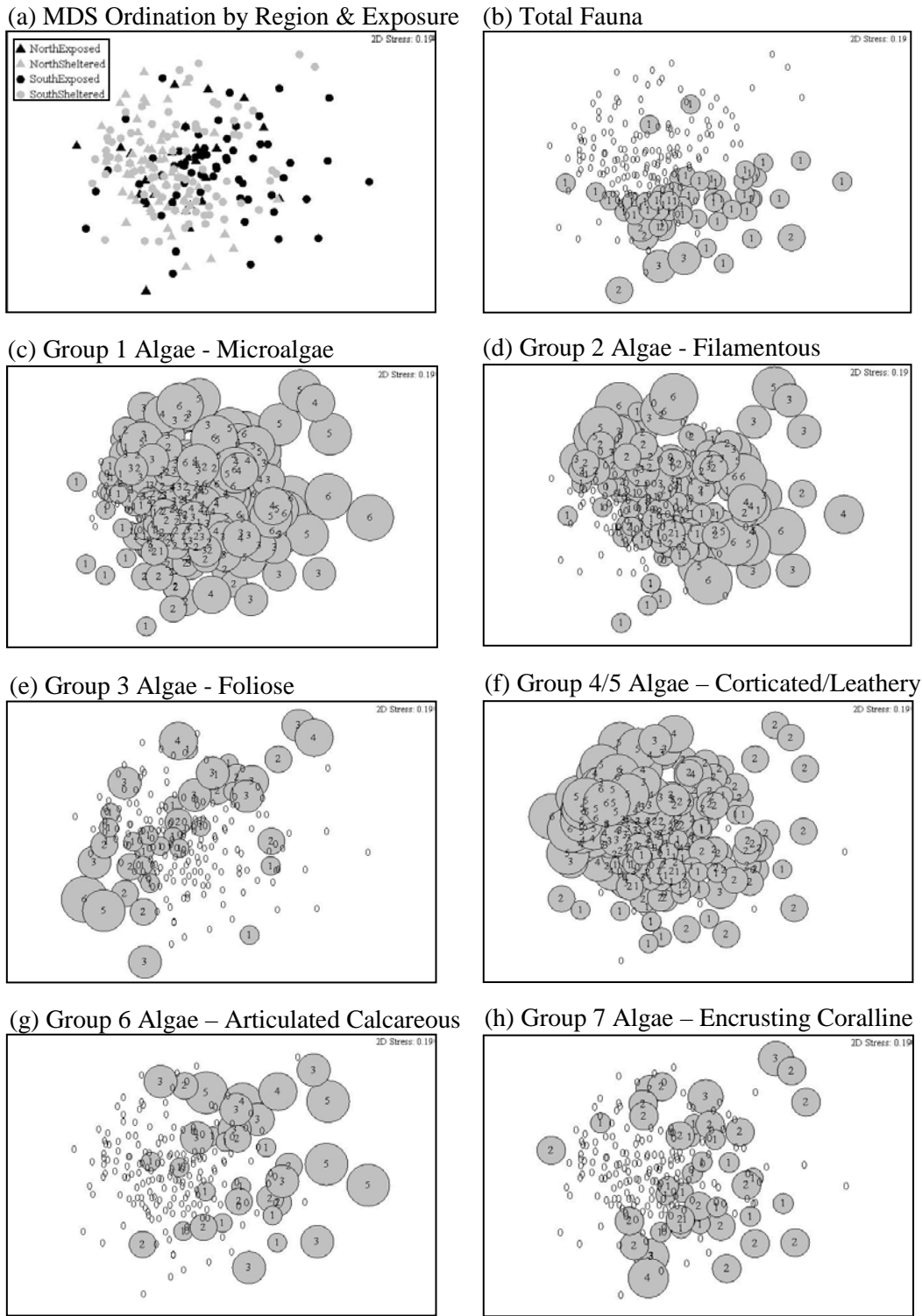


524

525 Fig. 3. *Patella vulgata* gut contents (n = 392). Abundance of corticated/leathery macroalgae  
526 as the proportional incidence in abundance categories (Table 1) by region and wave exposure.  
527

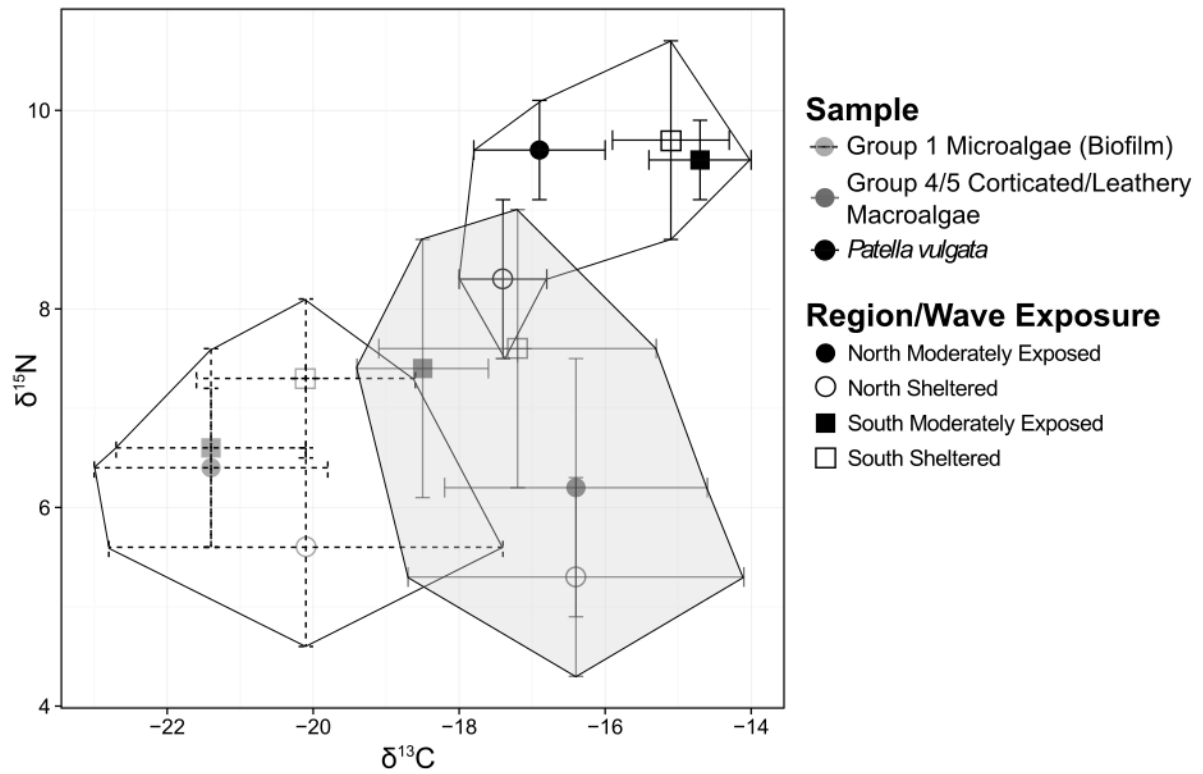
528

529  
 530  
 531  
 532  
 533  
 534  
 535  
 536  
 537  
 538  
 539  
 540  
 541  
 542  
 543  
 544  
 545  
 546  
 547  
 548  
 549  
 550  
 551  
 552  
 553  
 554  
 555  
 556  
 557  
 558  
 559  
 560  
 561  
 562  
 563  
 564  
 565  
 566  
 567  
 568  
 569  
 570



571 ① Rare, ② Occasional, ③ Frequent, ④ Common, ⑤ Abundant and ⑥ Superabundant.  
 572

573 Fig. 4. *P. vulgata* gut contents. MDS ordination of the composition of individual *P. vulgata*  
 574 diets using abundance categories converted to numerical integers using Gower's  $S_{15}$   
 575 similarity matrix ( $n = 392$ ). Subplots show (a) region and wave exposure of collection sites  
 576 (b) the abundance category for total fauna as a varying sized symbol, (c) microalgae, (d)  
 577 filamentous algae, (e) foliose algae, (f) corticated and leathery algae, (g) articulated  
 578 calcareous algae and (h) encrusting coralline algae.



579  
 580  
 581  
 582  
 583  
 584

Fig. 5. Stable isotope ratios of Microalgae, Corticated/Leathery Macroalgae and *P. vulgata* foot tissues. Mean  $\delta^{13}C$  values are given against mean  $\delta^{15}N$  values for tissues from moderately exposed and sheltered sites in the two study regions  $\pm 1$  SD.