

## Manuscript Details

<b>Manuscript number</b>	PROOCE_2018_81_R2
<b>Title</b>	Macrofaunal assemblages in canyon and adjacent slope of the NW and Central Mediterranean systems
<b>Article type</b>	Full Length Article

### Abstract

Macrofaunal assemblages were studied along bathymetric transects in six canyons and four adjacent open slopes of the Mediterranean Sea. The different areas investigated were located approximately along a longitudinal gradient at similar latitudes. Three regions were investigated: the Catalan (from 334 to 1887 m depth), the Ligurian (from 222 to 2005 m depth) and the South Adriatic margins (from 196 to 908 m depth). The analysis of the meso-scale distribution of assemblage structure and biomass showed significant differences among regions, which resulted in high values of  $\delta$ -diversity. Clear differences in trophic composition were also observed, and a decreasing pattern in the individual body size of macrofaunal organisms moving Eastward. These patterns were apparently linked to changes in food supply, whereas macrofaunal abundance and number of taxa showed a decrease pattern with increasing water depth. When the assemblage structure was compared between canyons and adjacent open slope, a very high  $\beta$ -diversity was observed, indicating that the bottom topography exerted a strong effect on the assemblage characteristics.

<b>Keywords</b>	Macrofauna; biomass; biodiversity; deep-sea; canyons; Mediterranean Sea
<b>Manuscript category</b>	Biological Oceanography
<b>Corresponding Author</b>	Anabella Covazzi Harriague
<b>Order of Authors</b>	Anabella Covazzi Harriague, Roberto Danovaro, Cristina Misic
<b>Suggested reviewers</b>	Georges Stora, Francisco Sardà, Maite Louzao

## Submission Files Included in this PDF

### File Name [File Type]

Covazzi Harriague et al response to reviewers second round.docx [Response to Reviewers]

Covazzi Harriague et al highlights.docx [Highlights]

Covazzi Harriague et al text revised second round.doc [Manuscript File]

Figure 1.tif [Figure]

Figure 2.tif [Figure]

Figure 3.tif [Figure]

Figure 4.tif [Figure]

Figure 5.tif [Figure]

Figure 6.tif [Figure]

Figure 7.tif [Figure]

Covazzi Harriague et al Tables revised second round.docx [Table]

Covazzi Harriague et al Supporting material second round.docx [Table]

To view all the submission files, including those not included in the PDF, click on the manuscript title on your EVISE Homepage, then click 'Download zip file'.

Point by point response to Reviewers. In black the reviewer's observations and suggestions, in red our response.

### **-Reviewer 1**

Line 130: .....(Gage, 2003). this gradient is CHANGE TO "This" **Done**

LINES 143-145: CHANGE TO: a very high number of canyons whose complexity reflect the features .... **Done**

LINE 155 "to tidal currents" I WOULD SPECIFY that tides re negligible in the Mediterranean **Done**

LINE 196: You should, quote along with Mamouridis et al., 2011 also:

TECCHIO S., RAMÍREZ-LLODRA E., AGUZZI J., FLEXAS M.M., COMPANY J.B., SARDÀ F. 2013. Seasonal fluctuations of deep megabenthos: Finding evidence of standing stock accumulation in a flux-rich continental slope. Progress in Oceanography 118: 188-198

**Done**

LINE 220: i) SHOULD BE ii) **Done**

LINES 235-262 THESE INFO SHOULD BE CONDENSED INTO Table 1 adding new columns for N replicates and for R/V name.

**The info on the R/V name and sampling period was added in the Table 1. The number of replicates was always 3 and it was reported in the text.**

LINE 286: Split the paragraphs (a new argument is starting). **Done**

LINE 308-309: this is a single phrase and should be joined into the previous paragraph (although it is a different argument). **Done**

LINE 430 sp SHOULD BE CHANGED TO sp. **Done**

LINE 505 "Notwithstanding" SHOULD BE CHANGED for "Despite" **Done**

LINE 517: CHANGE TO (21%) **Done**

LINE 594: You should also quote:

Coll M., Piroddi C., Steenbeek J., Kaschner K., Ben RaisLasram F., Aguzzi J., et al. 2010. Biodiversity of the Mediterranean Sea: status, patterns and threats. PLoS ONE (5): e11842

**Done**

LINES 535-637: Quotations need to be added in chronologic mode). **Done**

LINES 681-683: You declare that "... we observed a lack of relationships between the macrofaunal parameters and grain size". A DESCRIPTION OF THESE RESULTS SHOULD BE GIVEN IN THE RESULT SECTION (Table S3 should be reported there). **Done**

LINE 808: eliminate one indent (too large paragraph space). **Done**

Table 1: ADD SEPARATION IN Catalanmargin, Ligurianmargin **Done**

Also, in this table you code the sampling areas (add the head name in that column) as e.g. Southern Open Slope (SOS) or Cap de Creus Canyon (CCC).....but then in Tables 3 and 4, those codes are forgotten. Please, choose if eliminate the codes from Table 1 or change names of areas in Table 3 and 4 for those codes for consistency. **Done**

In table 3. All percentage values should be with numerals e.g. 3rd line form above of values as

33 0 0 33 0 33 is equals top 99!

**We add the decimal values**

In Table S1: ALL "sp" should be "sp." **Done**

In Table S2. Add space in "...diversity),between" **Done**

Also here...Canyon and Open Slope should be written with the first capital letters as in the Tables before (and please, DO THE SAME FOR Table S1). **Done**

Also:

MODIFY "Table S3.Result" for "Table S3. Result" **Done**

NOT commas for decimals but DOTS as in all other numbers. **Done**

## **-Reviewer 2**

Dear authors.

I recommend minor revisions. These concerns mostly to the discussion which still needs improvement for a better understanding of it. The authors begin the discussion by explaining the differences in community structure between the three investigated regions, then follow by examining patterns observed in the canyon and open slope within each region. Next, the authors use food availability to explain longitudinal patterns in biomass, individual size and trophic composition and then the same variable to explain the variability in trophic composition between canyon and open slope. Food availability could be used in the canyon vs open slope topic and elaborate the discussion in this part that sometimes is vague. In fact, the reason why there were no differences between canyon and open slope in the Ligurian margin, is at the end of the discussion, while in my opinion it is missing in the part where is discussed the local patterns.

Also, the canyon and open slope results discussion has some confusion and inconsistencies. For example, the authors compare changes in the assemblage structure between canyon and open slope that were not significant with significant differences found in community structure with other canyons. Also, the authors give examples of species that can explain the differences in dissimilarity between canyon and open slope but it is not clear how the presence of one of these species, the ophiurid *Amphiura filiformis* in both canyon and open slope explain the dissimilarity. The next part is somehow confusing because the authors indicate studies (Gage et al., 1995; Vetter and Dayton, 1999; Curdia et al., 2004) that are in accordance with the present results in relation to the lower number of taxa found inside some canyons but next indicate discrepancy of these findings with previous studies. It is not clear which results and in which canyons are the authors referring to establish the comparison.

I indicate these and other minor comments throughout the discussion in the document. In addition, some comments in the remaining sections, also indicated in the document.

**We reorganized the Discussion following carefully all the reviewer suggestions and we took into account all the reviewer comments indicated in the document.**

### Introduction

Missing reference in the sentence:

“The analyses conducted on macrofaunal assemblages along longitudinal gradients at large spatial-scales (from the Gulf of Lyon to the Ligurian Sea and Southern Adriatic) in the Mediterranean Sea support the important role of these variables and the importance of these systems in promoting deep-sea biodiversity. “

**This sentence was deleted**

### Material and methods:

Please indicate the dissimilarity measure used in the MDS (bray Curtis measure). **Done**

Please indicate how diversity was estimated (number of taxa). **Done**

### Results:

In the subsection *Macrofaunal assemblage structure* please change “more significant” to “significant”. **Done**

In the subsection *Macrofaunal biomass* please add the results of the regression analysis of biomass according to the longitude.

**For the relationship between individual biomass and longitude we have highlighted a general trend. A statistical analysis such as regression was not performed because longitude data were considered not a quantitative parameter as, for instance, depth is, but something more similar to a nominal variable. The individual biomass data in each area (Catalan, Ligurian and South Adriatic margins) are referred to very similar longitude values, therefore regression output would be somehow**

influenced by this homogeneity. Maybe the sentence was misleading, suggesting that a regression result should exist. We change the sentence.

Some comments on figures and tables:

Figure 2: The legend should indicate that the two MDS images correspond to two different analysis.

Done

The authors indicate that the available data on mud contribution to the sediment composition were added but is missing.

The data of mud contribution to the sediment composition are presented in Table 1 (the last column)

A table with the ANOSIM results for the trophic composition should also be added.

We added the ANOSIM results for the trophic composition in the table 2

Finally, some typos were assigned throughout the text.

- Macrofauna was studied along bathymetric transects in three areas of the Mediterranean
- Canyons and open slopes were investigated in Catalan, Ligurian and South Adriatic margins
- Assemblages were significantly different among regions, as shown by high  $\delta$ -diversity
- This was likely due to changes in food supply as a consequence of the longitudinal position
- Canyons and adjacent open slope showed a very high  $\beta$ -diversity

1  
2  
3 **Macrofaunal assemblages in canyon and adjacent slope of the**  
4  
5 **NW and Central Mediterranean systems**  
6  
7  
8  
9

10  
11 Anabella Covazzi Harriague<sup>1</sup>, Roberto Danovaro<sup>2,3</sup>, Cristina Misic<sup>1</sup>

12  
13 <sup>1</sup>DisTAV – Università di Genova – Genoa, Italy

14  
15 <sup>2</sup>DiSVA – Università Politecnica delle Marche – Ancona, Italy

16  
17 <sup>3</sup> Stazione Zoologica Anton Dohrn –Naples, Italy  
18  
19  
20  
21  
22  
23

24 Submitted for publication to: Progress in Oceanography

25 Date of submission: 6<sup>th</sup> April 2018

26  
27 Date of submission of the amended version: 30<sup>th</sup> November 2018  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41

42 **Corresponding author:**

43  
44 Dr. Anabella Covazzi Harriague

45  
46 Dipartimento di Scienze della Terra, dell’Ambiente e della Vita - Università di Genova

47  
48 C.so Europa 26 - 16132 Genova, Italy

49  
50 Phone: 0039 01035338224

51  
52 e-mail: anabella7@hotmail.com  
53  
54  
55  
56  
57  
58  
59  
60

61  
62  
63 **Abstract**  
64

65 Macrofaunal assemblages were studied along bathymetric transects in six canyons and four  
66 adjacent open slopes of the Mediterranean Sea. The different areas investigated were located  
67 approximately along a longitudinal gradient at similar latitudes. Three regions were  
68 investigated: the Catalan (from 334 to 1887 m depth), the Ligurian (from 222 to 2005 m  
69 depth) and the South Adriatic margins (from 196 to 908 m depth). The analysis of the meso-  
70 scale distribution of assemblage structure and biomass showed significant differences among  
71 regions, which resulted in high values of  $\delta$ -diversity. Clear differences in trophic composition  
72 were also observed, and a decreasing pattern in the individual body size of macrofaunal  
73 organisms moving Eastward. These patterns were apparently linked to changes in food  
74 supply, whereas macrofaunal abundance and number of taxa showed a decrease pattern with  
75 increasing water depth. When the assemblage structure was compared between canyons and  
76 adjacent open slope, a very high  $\beta$ -diversity was observed, indicating that the bottom  
77 topography exerted a strong effect on the assemblage characteristics.  
78  
79  
80  
81  
82  
83  
84  
85  
86  
87  
88  
89  
90  
91  
92  
93  
94

95 Key words: Macrofauna, biomass, biodiversity, deep-sea, canyons, Mediterranean Sea.  
96  
97  
98  
99  
100  
101  
102  
103  
104  
105  
106  
107  
108  
109  
110  
111  
112  
113  
114  
115  
116  
117  
118  
119  
120



121  
122  
123 **1. Introduction**  
124

125 The Mediterranean Sea is characterised by a clear gradient in primary productivity with  
126 lowest values in the eastern basin (Danovaro et al., 1999). Since food supply to deep-sea  
127 benthic fauna depends on the productivity of the euphotic zone (Gage, 2003). This gradient is  
128 typically reflected by a decreasing abundance, biomass and species richness of the deep  
129 benthic assemblages moving eastward (Tselepidis et al., 2000; Danovaro et al., 2008; Tecchio  
130 et al., 2011). One of the reasons for the extreme oligotrophy of the eastern Mediterranean  
131 basin is phosphorous depletion (Salihoglu et al., 1990), while in the western region the fluvial,  
132 atmospheric and upwelling inputs contribute to a higher productivity (Bas, 2009). Another  
133 main feature of the Mediterranean continental margins is the presence of a very high number  
134 of canyons, whose complexity reflect the features of the fluvial basins of the continent (Bas,  
135 2009).

136 Submarine canyons contribute to the formation of a number of different habitats that show  
137 spatial as well as temporal variability, and are often characterised by peculiar benthic  
138 assemblages (Curdia et al., 2004). The dynamic characteristics of these environments, subject  
139 to tidal currents albeit tides are negligible in Mediterranean, sediment gravity flows and  
140 turbidity changes (Canals et al., 2006; de Stigter et al., 2007), increase the physical forcing,  
141 habitat heterogeneity and the complexity of the morphological features of the seafloor (Thistle  
142 and Levin, 1998). Canyons are areas with enhanced transfer of materials from the continental  
143 shelf to bathyal depths (Epping et al., 2002; Martin et al., 2006; Sanchez-Vidal et al., 2008;  
144 Thomsen et al., 2017), and provide an important food supply to the deep-sea benthos (Epping  
145 et al., 2002; Gunton et al., 2015). The canyons act also as collectors of land-derived organic  
146 material, which contribute to shape and sustain benthic assemblages (de Stigter et al., 2007).  
147 For instance, massive inputs of organic materials, including macrophyte detritus (Rowe et al.,  
148 1982; Vetter and Dayton, 1999), are transported from the shelf to the deep sea during periodic  
149

181  
182  
183 flushing events (Canals et al., 2006). These processes contribute to make the Mediterranean  
184  
185 canyons hotspots of benthic production and ecosystem functioning (Vetter and Dayton, 1999;  
186  
187 Puseddu et al 2010; Fernández-Arcaya et al., 2017) and are responsible for their generally  
188  
189 high macrofaunal abundance and biomass (Vetter and Dayton, 1999; Duineveld et al. 2001;  
190  
191 Gunton et al. 2015). For this reason, studies conducted in Mediterranean Sea have, in many  
192  
193 cases, highlighted the presence of differences between canyons and the adjacent open slopes  
194  
195 (Mamouridis et al., 2011; Tecchio et al., 2013). Such differences include also the presence of  
196  
197 some taxa that are exclusively or preferentially associated to canyon sediments (Bianchelli et  
198  
199 al., 2010) and are the result of the combined action of habitat heterogeneity and food  
200  
201 availability.  
202  
203

204  
205 In the present study, macrofaunal assemblages were investigated within the frame of several  
206  
207 cruises conducted in different regions at depths comprised approximately from 200 to 2000 m  
208  
209 depth. The following hypotheses were tested: i) structural and functional features of the  
210  
211 macrofaunal assemblages among different continental margins change along a longitudinal  
212  
213 gradient; ii) macrofaunal variables change between canyons and adjacent open slopes in the  
214  
215 same area.  
216  
217

## 218 219 **2. Material and Methods**

### 220 221 *Sampling strategy*

222  
223 Samples were collected in the Western and Central Mediterranean Sea during several  
224  
225 multidisciplinary oceanographic cruises. Three regions were studied from 196 to 2005 m  
226  
227 depth. The Catalan margin was sampled during the HERMES cruises in the Gulf of Lyon (11  
228  
229 sampling stations), the Ligurian margin was sampled during the BioLig (Ligurian Canyons)  
230  
231 cruise (12 sampling stations) and the South Adriatic margin was sampled during the  
232  
233  
234  
235  
236  
237  
238  
239  
240

241  
242  
243 CANYON BARI 2006 (SETE-06) cruise (11 sampling stations) (see Figure 1, Table 1 for  
244 details).  
245

246  
247 Three replicate samples were collected in each station, using either multi-corer (3 maxicorer,  
248 with inner diameter 9.3 cm for each replicate) and box-corer (29x29 cm). On board, all the  
249 sediment collected was washed and sieved through 500 µm mesh size. After that, the residual  
250 was frozen at -20°C.  
251  
252  
253  
254  
255

### 256 257 *Macrofaunal variables*

258  
259 In laboratory, sediment of all the stations was sorted and organisms were identified down to  
260 species level (Table S1), whenever possible, and counted. Taxonomical identification (Fauvel,  
261 1923; 1927; Parenzan, 1976; Bellan-Santini et al., 1982; 1989; 1993; Pancucci-Papadopoulou  
262 et al., 1999) was checked with the World Register of Marine Species  
263 (<http://www.marinespecies.org>). The number of taxa, intended as a proxy of diversity, was  
264 calculated for each station  
265  
266  
267  
268  
269  
270  
271  
272

273 To determine the biomass, organisms were dried at 60°C for 24 hours and weighed. Data of  
274 number of organisms and weight were normalised to square meter to determine abundance  
275 and biomass applying the following formula:  $\text{abundance/biomass} = \text{organism}$   
276  $\text{number/organism weight of the sample/sampled area in square meter}$ . The individual biomass  
277 of organisms was calculated as the ratio between biomass and abundance. Trophic groups  
278 were determined according to Fauchald and Jumars, 1979; Jangoux and Lawrence, 1982;  
279 Grahame, 1983; Russell-Hunter, 1983 and Gambi and Giangrande, 1985.  
280  
281  
282  
283  
284  
285  
286  
287  
288  
289

### 290 *Statistical analyses*

291  
292 The PRIMER 6 package (Clarke and Warwick 2001) on presence/absence data was used to  
293 perform the multidimensional scaling (MDS, Bray-Curtis similarity index) and the analysis of  
294  
295  
296  
297  
298  
299  
300

301 similarities (ANOSIM), to test the differences in community structure within and between the  
302 sites. Moreover the similarity percentages-species contributions (SIMPER) was performed,  
303 based on a Bray-Curtis similarity matrix, to evaluate the turnover local diversity (hereafter  
304 called  $\beta$ -diversity) and regional turnover diversity (hereafter called  $\delta$ -diversity) (Gray, 2000;  
305 Danovaro et al., 2009). In order to homogenize the data, the taxonomical level family was  
306 used for all analysis. The analysis of similarities (ANOSIM) was performed on the  
307 presence/absence data of the trophic composition in order to verified differences between  
308 sites. The relationship between community parameters and depth and mud contribution to the  
309 sediment composition were verified with the regression analysis.  
310  
311  
312  
313  
314  
315  
316  
317  
318  
319  
320  
321  
322  
323

### 324 **3. Results**

#### 325 *Macrofaunal assemblage structure*

326  
327 The MDS ordination (Figure 2) showed that the stations of the Ligurian margin are grouped  
328 together, while the other two sites are not clearly separated. Notwithstanding, the analysis of  
329 similarities (ANOSIM) showed significant differences in the structure of the communities  
330 between the three regions (Table 2A).  
331  
332

333 The trophic composition of the assemblages was significantly different between the Ligurian  
334 margin and the Catalan and South Adriatic margins (ANOSIM, Table 2B).  
335  
336

337  $\delta$ -diversity between the three regions was high: Catalan margin vs Ligurian margin 97% (4  
338 common taxa: Cirolanidae, Golfingidae, Nuculidae, Paraonidae), Catalan margin vs South  
339 Adriatic margin 97% (3 common taxa: Canalipalpata, Terebellidae, Xanthidae) and Ligurian  
340 margin vs South Adriatic margin 98% (3 common taxa: Copepoda, Phascolosomatidae,  
341 Spionidae). Each region was characterized by different taxa that determined the regional  
342 variability: in the Ligurian margin the main part of them belonged to polychaetes, while in the  
343  
344  
345  
346  
347  
348  
349  
350  
351  
352  
353  
354  
355  
356  
357  
358  
359  
360

361  
362  
363 Catalan margin the half of the taxa and in the South Adriatic margin only one family belonged  
364  
365 to this class (Figure 3).  
366

367  
368  
369 - Catalan margin  
370

371  
372 The macrofaunal abundance varied between 0 and  $254.6 \pm 28.9$  ind  $\times$  m<sup>-2</sup> (average  $91.2 \pm 24.1$   
373 ind.  $\times$  m<sup>-2</sup>). The highest values were found in the shallowest stations of the NOS and LDC  
374 (Table 3). The average value was higher in the open slope station than in the canyon ones  
375 (Figure 4A). The distribution of abundances was significantly influenced by the depth (Figure  
376 5A). On the whole 30 taxa were identified, 14 in the canyon and 17 in the open slope  
377 (Figure 4B). Canyon and open slope communities showed only 1 common taxa, leading to a very high  
378 local  $\beta$ -diversity (95%; Table S2). Although the community structure did not show a  
379 significant differences between habitats (Table 2A). The number of taxa, instead, did not  
380 show a significant relationship with depth (Figure 5B). The structural parameters did not  
381 show any correlation with the contribution of the mud fraction (Table S3). Assemblages were  
382 dominated by Annelida (50%), followed by Crustacea (19%), Echinodermata (13%),  
383 Sipuncula (10%), Mollusca (5%) and others (Platyhelminthes and Nemertea; 3%). The first  
384 group was the most important in the canyon area, reaching 70% of the abundance, while in  
385 the open slope area 86% of the abundance was composed by Annelida, Crustacea and  
386 Echinodermata (Table 3). The trophic structure of the communities was different in the two  
387 areas: in the assemblages of the canyons the principal contributors to the abundance were  
388 organisms with mixed strategies, deposit-suspension (such as *Onchnesoma steenstrupii*  
389 *steenstrupii* and Terebellidae) and deposit-grazer feeders (such as *Notomastus* sp.), while in the  
390 open slope assemblages suspension feeders (such as *Jassa marmorata* and *Thalassema gigas*),  
391  
392  
393  
394  
395  
396  
397  
398  
399  
400  
401  
402  
403  
404  
405  
406  
407  
408  
409  
410  
411  
412  
413  
414  
415  
416  
417  
418  
419  
420

421  
422  
423 deposit feeders (such as Chaetopteridae and Paraonidae) and predators (such as *Ancistrosyllis*  
424  
425 *cf. groenlandica*) were the most abundant (Figure 6).  
426  
427

428  
429 - Ligurian margin  
430

431  
432 The higher abundances were observed in the shallow stations (see Table 3). The maximum  
433  
434 value was  $396.4 \pm 97.4$  ind.  $\times$  m<sup>-2</sup> in the shallow station of the PC and the minimum  $15.0 \pm 15.0$   
435  
436 ind.  $\times$  m<sup>-2</sup> in the deepest station of the LMOS (average  $141.6 \pm 39.2$  ind.  $\times$  m<sup>-2</sup>). Contrary to the  
437  
438 Catalan margin, the average abundance in the canyon areas was slightly higher than that  
439  
440 observed in the open slope ones (see Figure 4A). In general, a significant decreasing trend of  
441  
442 macrofaunal abundance with depth was observed (see Figure 5A). In the Ligurian margin 43  
443  
444 taxa were found, 23 in the open slope area and 34 in the canyons, and 14 were observed in  
445  
446 both areas. Here the  $\beta$ -diversity was 82%, the lowest of the three sites (Table S2). Despite, the  
447  
448 community structure of the both habitats did not differ significantly (Table 2A). The average  
449  
450 values of the number of taxa were higher in the canyon (see Figure 4B). A significant  
451  
452 decrease of the number of taxa with depth was observed (Figure 5B). The structural  
453  
454 parameters did not show any correlation with the contribution of the mud fraction (Table S3).  
455  
456 Total abundance was composed by Annelida (76%) and Crustacea (15%). In the canyon areas  
457  
458 these groups represented 70% and 15%, respectively, of the total abundance, while in the  
459  
460 open slope assemblages they represented 83% and 12% respectively (see Table 3). From a  
461  
462 trophic point of view, the composition of the assemblages in the canyons and open slope was  
463  
464 quite similar, being dominated by deposit feeders (such as Maldanidae and Onuphidae) (55%  
465  
466 and 60%, respectively), followed by predators (such as Eunicidae and Lumbrineridae) (18%  
467  
468 and 14%, respectively) and deposit-suspension feeders (such as Spionidae and *Urothoe cf.*  
469  
470 *elegans*) (13% and 11%, respectively) (see Figure 6).  
471  
472  
473  
474  
475  
476  
477  
478  
479  
480

481  
482  
483 - South Adriatic margin  
484

485 The higher abundance was observed in the shallowest station of the open slope  
486 (2747.3±1066.7 ind. × m<sup>-2</sup>), while the deepest stations of the two canyons showed the  
487 minimum value (0 ind. × m<sup>-2</sup>, see Table 3). A significant decrease of the abundance with  
488 depth was observed (see Figure 5A). Between the two areas, the open slope displayed a higher  
489 average value than canyon one (see Figure 4A). On the whole, 22 taxa were found in the  
490 South Adriatic margin, 13 in the open slope and 12 in the canyon areas. Despite the similar  
491 number, the average values were different, open slope displayed the highest one (see Figure  
492 4B). Only 3 taxa were common at the two areas and the β-diversity was 88% (Table S2).  
493  
494 Nevertheless, the analysis of similarities didn't show significant differences between the two  
495 habitats (Table 2). The number of taxa showed a significant decrease trend with depth (see  
496 Figure 5B). Abundance did not show any correlation with the contribution of the mud  
497 fraction, while a significant correlation was found for the number of taxa (Table S3).  
498  
499 Communities were composed by Crustacea (28%) followed by Annelida (22%),  
500 Echinodermata (21%), Mollusca (14%), others (Nematoda) (10%) and Sipuncula (5%).  
501  
502 Assemblages on the canyon areas were composed principally by 35% of Mollusca and 35% of  
503 Echinodermata, while in the open slope, the Annelida and Crustacea reached 72% of the total  
504 abundance (see Table 3). Also, the trophic composition was different for the two areas: in the  
505 canyon sites 70% of the abundance was represented by suspension feeders (such as *Amphiura*  
506 *filiformis* and *Clausinella fasciata*), while in the open slope 77% was represented by deposit-  
507 suspension feeders (such as Spionidae), deposit feeders-predators (such as Nematoda) and  
508 predators (such as *Acanthephyra eximia*) (see Figure 6).  
509  
510  
511  
512  
513  
514  
515  
516  
517  
518  
519  
520  
521  
522  
523  
524  
525  
526  
527  
528  
529  
530

531  
532 *Macrofaunal biomass*  
533  
534  
535  
536  
537  
538  
539  
540

541  
542  
543 Values of biomass and the relative contribution of each taxa to the biomass are presented in  
544 Table 4. The decapod *Xantho pilipes* was the responsible for the highest value (13422.9 mg ×  
545 m<sup>-2</sup>) observed in the shallowest station of the SOS in the Catalan margin. The contribution of  
546 the different groups was quite similar for Ligurian and South Adriatic margins, but in the  
547 Catalan margin 93% of the total biomass was due to Crustacea and Echinodermata. The three  
548 sites showed a distribution of the biomass in the canyon and open slope areas similar to the  
549 abundance one (see Figure 4C). In general, macrofaunal biomass did not show a significant  
550 relationship with depth. Only in the Catalan margin a significant decrease of the values with  
551 increasing water depth was observed (see Figure 5C). Biomass did not show any correlation  
552 with the contribution of the mud fraction (Table S3).  
553

554 The individual biomass was determined, highlighting a general decreasing trend moving  
555 eastward. The highest values were observed in the western area (Figure 7A). The average  
556 value for the canyon areas was 1.7± 1.1, 1.4±1.4 and 1.3±1.6 mg ind<sup>-1</sup> for the Catalan,  
557 Ligurian and South Adriatic Margins respectively. A high variability was observed for the  
558 open slope values: 40.4±67.5, 0.3±0.2 and 2.6±2.5 mg ind<sup>-1</sup> for the Catalan, Ligurian and  
559 South Adriatic Margins respectively. The relationship with depth was not significant (R =  
560 0.1735, n = 32, p>0.05; Figure 7B).  
561  
562  
563

#### 564 **4. Discussion**

565 The ANOSIM analysis indicated the presence of significant differences in the structure of the  
566 macrobenthic assemblages among the different sampling regions along the longitudinal  
567 gradient. Such differences were observed also in terms of high turnover diversity among  
568 different investigated regions which resulted in a high  $\delta$ -diversity. Only Oligochaeta and,  
569 within the polychaetes, the family Capitellidae, were reported from all of the investigated  
570 regions. The observed variability at regional scale of the macrofaunal assemblages is  
571  
572  
573  
574  
575  
576  
577  
578  
579  
580  
581  
582



601  
602  
603 apparently linked to the specific environmental features, including differences in primary  
604 productivity and organic carbon fluxes, which decreased moving Eastward, with consequent  
605 decrease of the organic matter available to consumers in deep-sea sediments (Gambi and  
606 Danovaro, 2006; Coll et al., 2010). Also the temporal variability should be considered, the  
607 regions were sampled in different years and different seasons. The importance of food source  
608 in structuring deep-sea assemblages was underlined also by Mamouridis et al., (2011) who  
609 reported seasonal changes in macrofaunal communities of the Besos canyon.  
610  
611

612 The trophic composition of macrofaunal assemblages showed differences between the three  
613 regions, in accordance with the differences reported in terms of food availability (Pusceddu et  
614 al., 2010). In fact, where the quality and quantity of the sedimentary organic matter were  
615 higher, such as in the Catalan margin, the dominant trophic guild were the deposit feeders and  
616 suspension-deposit feeders. Conversely, in the South Adriatic margin where lower food  
617 resources were available the macrofaunal assemblages were dominated by suspension  
618 feeders and predators. The predator contribution to the abundance (Figure 6) increased from  
619 West to East. All meiofaunal variables investigated in the same expeditions decreased moving  
620 Eastward (Bianchelli et al. 2010), we could hypothesize a predatory pressure of the  
621 macrofauna on meiofauna and/or competition between those benthic compartments. On the  
622 other hand, the food resources in the sediment of Mediterranean Sea decrease eastward  
623 reaching a strong oligotrophic condition in the Levantine basin (Danovaro et al. 1999). In  
624 such conditions predators are facilitated in obtaining food due to their higher motility  
625 (Danovaro et al., 2008). The same result was reported by Baldrighi et al. (2014) from the  
626 analysis of the deep macrofaunal assemblages across a longitudinal gradient in Mediterranean  
627 open slopes.  
628  
629  
630  
631  
632  
633  
634  
635  
636  
637  
638  
639  
640  
641  
642  
643  
644  
645  
646  
647  
648  
649  
650  
651  
652  
653  
654  
655  
656  
657  
658  
659  
660

661  
662  
663 The structure of the assemblages within each of the three regions changed also comparing the  
664 two habitats considered (canyon and open slope). Such changes resulted in high values of the  
665  $\beta$ -diversity between the habitats. Duineveld et al. (2001) and Gunton et al. (2015), comparing  
666 the Whittard canyon with the adjacent continental slope, reported significant differences in  
667 macrofaunal assemblages. The macrofaunal dissimilarity between canyons and slopes  
668 reported in the present study was slightly higher than that reported for the rare taxa of the  
669 meiofaunal communities in the Catalan and South Adriatic margins by Bianchelli et al.  
670 (2010), and higher than the dissimilarity for the nematodes assemblages for the same regions  
671 (Danovaro et al., 2009). One possible explanation of the observed patterns is that the fauna  
672 inhabiting each habitat are adapted to the local environmental features such as the dynamic  
673 characteristics of the canyons, where there is a large transport of energy and materials from  
674 the shelf to the deep-sea (Vetter and Dayton, 1998; 1999; Canals et al., 2006; de Stigter et al.,  
675 2007). For instance, we found some individuals of the bivalve *Thyasira flexuosa* inside the  
676 Polcevera canyon and a conspicuous population of the ophiurid *Amphiura filiformis* inside  
677 both canyons, eight-fold higher than in the open slope, in the South Adriatic region. Both  
678 species are considered as indicators of high organic matter loads. In fact, the bivalve belongs  
679 to the first order opportunistic group and the ophiurid to the second order opportunistic group  
680 (Simboura and Zenetos, 2002).  
681  
682  
683  
684  
685  
686  
687  
688  
689  
690  
691  
692  
693  
694  
695  
696  
697  
698  
699  
700

701 The lower number of taxa in canyons than in adjacent slopes observed in the Catalan and  
702 Adriatic margins, are consistent with a number of other studies (Gage et al., 1995; Vetter and  
703 Dayton, 1999; Curdia et al., 2004).  
704  
705  
706  
707

708 Previous studies of macrofauna in canyons reported higher abundance and biomass than in the  
709 adjacent slope (Duineveld et al., 2001). This holds true also for the Ligurian margin, where  
710 the macrofaunal assemblages of the canyon stations showed a higher abundance than those of  
711 the slope, but contrasts with the Catalan and South Adriatic margins where an opposite  
712  
713  
714  
715  
716  
717  
718  
719  
720

721  
722  
723 pattern was reported. As previously mentioned, one of the key factors driving the community  
724 structure of the deep-sea fauna is the food supply (Ruhl and Smith, 2004). Previous studies  
725 conducted in the Catalan and South Adriatic margins (Pusceddu et al. 2010) showed that the  
726 food availability was similar in canyon and open slope habitats. These findings are consistent  
727 with the macrofaunal distribution observed in the present study. In addition, the mud  
728 contribution to the sediment composition was quite similar between the two habitats in all  
729 regions studied. Also water depth played a potentially important role in structuring  
730 macrofaunal assemblages in both habitats at the Ligurian and South Adriatic systems, and this  
731 finding is in accordance with the observations reported in other areas such the Toulon Canyon  
732 (Stora et al., 1999) and the Aviles Canyon (Louzao et al., 2010).

733  
734 The local variability observed also between the topographic habitats in the Catalan and South  
735 Adriatic systems, can contribute to explain the observed differences, as the Catalan canyons  
736 dominated by organisms with mixed trophic strategies (deposit-suspension feeders and  
737 deposit feeders-grazers), pointing to the role of organic materials transported from the shelf to  
738 the deep-sea habitats (such as macrophyte detritus, Rowe et al., 1982; Vetter and Dayton,  
739 1999) during periodic flushing events and sedimentation of the transported material (Canals et  
740 al., 2006). Conversely, the South Adriatic, was dominated by suspension feeders, confirming  
741 the importance of primary production and the export of organic matter to the deep sea  
742 (Pusceddu et al., 2010). In the Ligurian margin the trophic composition of macrofaunal  
743 assemblages did not show differences between canyon stations and slope ones, and this  
744 finding is consistent with the fact that the two habitats showed similar amount of bioavailable  
745 organic matter (Carugati et al., this issue).

746  
747 The trophic characteristics of the sediments (as quality and quantity of sedimentary organic  
748 matter) of the Catalan and South Adriatic margins (Pusceddu et al., 2010) can also contribute

781  
782  
783 to explain the observed patterns in macrofaunal biomass, with highest values in the Catalan  
784 margin and a decreasing pattern Eastward. These findings are consistent with patterns  
785 observed for meiofauna by Gambi and Danovaro (2006) and are reflected also by patterns in  
786 individual size of the organisms, which decreased moving Eastward according to the  
787 depletion of the available food source (Pusceddu et al., 2010). Finally water depth did not  
788 play an important role in driving these faunal variables (see Figure 5C and 7B). We also  
789 observed the lack of any relationships between macrofauna and grain size (Table S3)  
790 suggesting that the textural characteristics did not play any significant key role in controlling  
791 the faunal distribution.  
792  
793  
794  
795  
796  
797  
798  
799  
800  
801  
802  
803

804 In conclusion, the analyses conducted on macrofaunal assemblages along longitudinal  
805 gradients at large spatial-scales (from the Gulf of Lyon to the Ligurian Sea and Southern  
806 Adriatic) in the Mediterranean Sea suggest that canyons by increasing  $\beta$ -diversity (i.e., the  
807 species turnover between canyons and open slopes) due to the spatial and trophic variability  
808 observed between canyons and slopes and among the different biogeographic regions.  
809  
810  
811  
812  
813  
814  
815  
816

### 817 **Acknowledgements**

818  
819 This study has been conducted in the framework of the Projects: HERMES (Hotspot  
820 Ecosystem Research on the Margin of European Seas), MERCES (Marine Ecosystem  
821 Restoration in Changing European Seas; Grant Agreement No. 689518), DG-ENV project  
822 IDEM: Implementation of the MSFD to the Deep Mediterranean Sea (contract EU No  
823 11.0661 /2017/750680/SUB/EN V.C2), and RITMARE (Ricerca Italiana per il Mare) funded  
824 by the Ministero dell'Istruzione, dell'Università e della Ricerca. The authors are indebted to  
825 S. Bianchelli, L. Bongiorno, M. Molari, G. Quero and D. Zeppilli as well as the crews of the  
826  
827  
828  
829  
830  
831  
832  
833  
834  
835  
836  
837  
838  
839  
840

841  
842  
843 ships Universitatis, Urania and Minerva II (Italy) and Tethys II (France) for their valuable  
844  
845 help in the sampling procedures.  
846  
847  
848

## 849 **References**

850  
851  
852 Baldrighi E., Lavaleye M., Aliani S., Conversi A., Manini E. 2014. Large Spatial Scale  
853  
854 Variability in Bathyal Macrobenthos Abundance, Biomass,  $\alpha$ - and  $\beta$ -Diversity along the  
855  
856 Mediterranean Continental Margin. PLoS ONE 9(9): e107261. doi:  
857  
858 10.1371/journal.pone.0107261.  
859

860 Bas C. 2009. The Mediterranean: a synoptic overview. Contributions to Science 5, 25–39.

861  
862 Bellan-Santini D., Karaman G., Krapp-Schieckel G., Ledoyer M., Myers A.A., Ruffo S.,  
863  
864 Schiecke S. 1982. The Amphipoda of the Mediterranean. Part 1 Gammaridea  
865  
866 (Acanthonotozomatidae to Gammaridae). Memoires de l'Institut Océanographique de  
867  
868 Monaco (Ed. S. Ruffo), 13: 1-364.  
869

870  
871 Bellan-Santini D., Diviacco G., Krapp-Schickel G., Myers A.A., Ruffo S. 1989. The  
872  
873 Amphipoda of the Mediterranean. Part 2 Gammaridea (Haustoriidae to Lysianassidae).  
874  
875 Memoires de l'Institut Océanographique de Monaco (Ed. S. Ruffo), 13: 365-576.  
876

877  
878 Bellan-Santini D., Karaman G., Krapp-Schieckel G., Ledoyer M., Ruffo S. 1993. The  
879  
880 Amphipoda of the Mediterranean. Part 3 Gammaridea (Melphidipidae to Talitridae).  
881  
882 Memoires de l'Institut Océanographique de Monaco (Ed. S. Ruffo), 13: 577-809.  
883

884  
885 Bianchelli S., Gambi C., Zeppilli D., Danovaro R. 2010. Metazoan meiofauna in deep-sea  
886  
887 canyons and adjacent open slopes: A large-scale comparison with focus on the rare taxa.  
888  
889 Deep-Sea Research I 57, 420-433.

890  
891 Canals M., Puig P., deMadron X.D., Heussner S., Palanques A., Fabres J. 2006. Flushing  
892  
893 submarine canyons. Nature 444, 354–357.  
894  
895  
896  
897  
898  
899  
900

- 901  
902  
903 Carugati L., Lo Martire M., Danovaro R. 2018. Patterns and drivers of meiofaunal  
904 assemblages in the canyons Polcevera and Bisagno of the Ligurian Sea (NW Mediterranean  
905 Sea) (this issue).  
906  
907  
908  
909  
910 Clarke K.R., Warwick R.M. 2001. Changes in marine communities: an approach to statistical  
911 analysis and interpretation, 2nd edition. PRIMER-E, Plymouth.  
912  
913  
914 Coll M, Piroddi C, Steenbeek J, Kaschner K, Ben Rais Lasram F, et al. 2010. The biodiversity  
915 of the Mediterranean Sea: estimates, patterns, and threats. PLoS ONE 5(8), e11842.  
916 doi:10.1371/journal.pone.0011842.  
917  
918  
919  
920 Curdia J., Carvalho S., Ravara A., Gage J.D., Rodrigues A.M., Quintino V. 2004. Deep  
921 macrobenthic communities from Nazaré submarine canyon (NW Portugal). Scientia Marina  
922 68, 171-180.  
923  
924  
925  
926  
927 Danovaro R., Dinet A., Duineveld G., Tselepides A. 1999. Benthic response to particulate  
928 fluxes in different trophic environments: a comparison between the Gulf of Lions–Catalan  
929 Sea (Western-Mediterranean) and the Cretan Sea (Eastern-Mediterranean). Progress in  
930 Oceanography 44, 287–312.  
931  
932  
933  
934  
935 Danovaro R., Gambi C., Lampadariou N., Tselepides A. 2008. Deep-sea nematode  
936 biodiversity in the Mediterranean basin: testing for longitudinal, bathymetric and energetic  
937 gradients. Ecography 31, 231–244.  
938  
939  
940  
941 Danovaro R., Bianchelli S., Gambi C., Mea M., Zeppilli D. 2009.  $\alpha$ -,  $\beta$ -,  $\gamma$ -,  $\delta$ - and  $\epsilon$ -diversity  
942 of deep-sea nematodes in canyons and open slopes of northeast Atlantic and Mediterranean  
943 margins. Marine Ecology Progress Series 396, 197-209.  
944  
945  
946  
947  
948 deStigter H.C., Boer W., de Jesus Mendes P.A., Jesus C.C., Thomsen L., van den Bergh  
949 G.D., van Weering T.C.E. 2007. Recent sediment transport and deposition in the Nazaré  
950 Canyon, Portuguese continental margin. Marine Geology 246, 144–164.  
951  
952  
953  
954  
955  
956  
957  
958  
959  
960

- 961  
962  
963 Duineveld G., Lavaleye M., Berghuis E., de Wilde P. 2001. Activity and composition of the  
964 benthic fauna in the Whittard Canyon and the adjacent continental slope (NE Atlantic).  
965  
966  
967 Oceanologica Acta 24, 69–83.  
968  
969  
970 Epping E., van der Zee C., Soetaert K., Helder W., 2002. On the oxidation and burial of  
971 organic carbon in sediments of the Iberian margin and Nazaré Canyon (NE Atlantic). Progress  
972 in Oceanography 52, 399-431.  
973  
974  
975  
976 Fauchald K., Jumars P.A. 1979. The diet of worms: a study of polychaete feeding guilds.  
977  
978 Oceanography and Marine Biology Annual Review 17, 193–284.  
979  
980  
981 Fauvel P. 1923. Faune de France, N° 5-Polychètes errantes. Federation française des Sociétés  
982 de Sciences Naturelles, France.  
983  
984  
985 Fauvel P. 1927. Faune de France, N° 16-Polychètes sédentaires. Federation française des  
986 Sociétés de Sciences Naturelles, France.  
987  
988  
989 Fernández-Arcaya U., Ramírez-Llodra E., Aguzzi J., Allcock A.L., Davies J.S., Dissanayake  
990 A., Harris P., Howell K., Huvenne V.A.I., Macmillan-Lawler M., Martín J., Menot L.,  
991  
992 Nizinski M., Puig P., Rowden A.A., Sánchez F., Van den Beld I.M.J. 2017. Ecological role of  
993 submarine canyons and need for canyon conservation: A review. *Frontiers in Marine Science* |  
994  
995  
996  
997 doi: 10.3389/fmars.2017.00005.  
998  
999  
1000 Gage J.D., Lamont P.A., Tyler P.A. 1995. Deep-sea microbenthic communities at contrasting  
1001 sites off Portugal, preliminary results. 1. Introduction and diversity comparisons.  
1002  
1003 *Internationale Revue der Gesamten Hydrobiologie* 80, 235-250.  
1004  
1005  
1006 Gage J.D. 2003. Food inputs, utilisation, carbon flow and energetics. In: Tyler P.A. (Ed.),  
1007  
1008 *Ecosystems of the Deep Oceans. Ecosystems of the World*. Elsevier, Amsterdam, 313–426.  
1009  
1010  
1011 Gambi C., Danovaro R. 2006. A multiple-scale analysis of metazoan meiofaunal distribution  
1012 in the deep Mediterranean Sea. *Deep-Sea Research I* 53, 1117-1134.  
1013  
1014  
1015  
1016  
1017  
1018  
1019  
1020

- 1021  
1022  
1023 Gambi, M.C., Giangrande, A. 1985. Caratterizzazione e distribuzione delle categorie trofiche  
1024 dei policheti nei fondi mobili del Golfo di Salerno. *Oebalia* 11, 223-240.  
1025  
1026  
1027 Grahame, J. 1983. Adaptive aspects of feeding mechanisms. *The Biology of Crustacea* 8,  
1028 65-107.  
1029  
1030  
1031 Gray J.S. 2000. The measurement of marine species diversity, with an application to the  
1032 benthic fauna of the Norwegian continental shelf. *Journal of Experimental Marine Biology*  
1033 *and Ecology* 250, 23-49.  
1034  
1035  
1036  
1037 Gunton L.M., Gooday A.J., Glover A.G., Bett B.J. 2015. Macrofaunal abundance and  
1038 community composition at lower bathyal depths in different branches of the Whittard Canyon  
1039 and on the adjacent slope (3500 m; NE Atlantic). *Deep-Sea Research* 97, 29-39.  
1040  
1041  
1042  
1043 Jangoux M., Lawrence J.M. (Eds.) 1982. *Echinoderm Nutrition*. A.A. Balkema Publishers,  
1044 Rotterdam, 654 pp.  
1045  
1046  
1047 Louzao M., Anadon N., Arrontes J., Alvarez-Claudio C., Fuente D.M. 2010. Historical  
1048 macrobenthic community assemblages in the Avilés Canyon, N Iberian Shelf: Baseline  
1049 biodiversity information for a marine protected area. *Journal of Marine Systems* 80, 47-56.  
1050  
1051  
1052  
1053 Martín J., Palanques A., Puig. P. 2006. Composition and variability of downward particulate  
1054 matter fluxes in the Palamó's submarine canyon (NW Mediterranean). *Journal of Marine*  
1055 *Systems* 60, 75-97  
1056  
1057  
1058  
1059 Mamouridis V., Cartes J.E., Parra S., Fanelli E., Saiz Salinas J.I. 2011. A temporal analysis  
1060 on the dynamics of deep-sea macrofauna: Influence of environmental variability off Catalonia  
1061 coasts (western Mediterranean). *Deep-Sea Research I* 58, 323–337.  
1062  
1063  
1064  
1065 Pancucci-Papadopoulou M.A., Murina G.V.V., Zenetos A. 1999. The phylum Sipuncula in  
1066 the Mediterranean Sea. Ed. National Centre for Marine Research, Athens.  
1067  
1068  
1069  
1070  
1071 Parnizan P. 1976. *Carta d'identità delle conchiglie del Mediterraneo*. Ed. Bios Taras, Taranto.  
1072  
1073  
1074  
1075  
1076  
1077  
1078  
1079  
1080



1081  
1082  
1083 Pusceddu A., Bianchelli S., Canals M., Sánchez -Vidal A., Durrieu De Madron X., Heussner  
1084 S., Likousis V., de Strigter H., Trincardi F., Danovaro R. 2010. Organic matter in sediments  
1085 of canyons and open slopes of the Portuguese Catalan, South Adriatic and Cretan Sea  
1086 margins. *Deep-Sea Research I* 57, 441-457.  
1087  
1088  
1089  
1090  
1091 Rowe G.T, Polloni P.T., Haedrich R.L. 1982. The deep-sea macrobenthos on the continental  
1092 margin of the northwest Atlantic Ocean. *Deep Sea Research* 29(2A), 257-278.  
1093  
1094  
1095  
1096 Ruhl H.A., Smith K.L. 2004. Shifts in deep-sea community structure linked to climate and  
1097 food supply. *Science* 305, 513–515.  
1098  
1099  
1100 Russell-Hunter, W.D. (Ed.), 1983. *Ecology. The Mollusca*, vol. 6. Academic Press, Orlando,  
1101 695 pp.  
1102  
1103  
1104 Salihoglu I., Saydam C., Bastürk Ö., Yilmaz K., Gö D., Hatipoglu E., Yilmaz A. 1990.  
1105 Transport of nutrients and chlorophyll-a by mesoscale eddies in the northeastern  
1106 Mediterranean. *Marine Chemistry* 29, 375-390.  
1107  
1108  
1109  
1110 Sánchez-Vidal A., Pasqual C., Kerhervé P., Calafat A., Heussner S., Palanques A., Durrieu de  
1111 Madron X., Canals M., Puig P. 2008. Impact of dense water cascading on the transfer of  
1112 organic matter to the deep western Mediterranean basin. *Geophysical Research Letters* 35, 1-  
1113 5.  
1114  
1115  
1116  
1117  
1118  
1119 Simboura N., Zenetos A. 2002. Benthic indicators to use in ecological quality classification of  
1120 Mediterranean soft bottom marine ecosystems, including a new Biotic Index. *Mediterranean*  
1121 *Marine Science* 3/2, 77-111.  
1122  
1123  
1124  
1125 Stora G., Bourcier M., Arnoux A., Gerino M., Le Campion J., Gilbert F., Durbec J.P. 1999.  
1126 The deep-sea macrobenthos on the continental slope of the northwestern Mediterranean Sea: a  
1127 quantitative approach. *Deep-Sea Research I* 46, 1339-1368.  
1128  
1129  
1130  
1131  
1132  
1133  
1134  
1135  
1136  
1137  
1138  
1139  
1140

- 1141  
1142  
1143 Tecchio S., Ramírez-Llodra E., Sardà F., Company J.B., Palomera I., Mecho A., Pedrosa-  
1144 Pamies R., Sanchez-Vidal A. 2011. Drivers of deep Mediterranean megabenthos communities  
1145 along longitudinal and bathymetric gradients. *Marine Ecology Progress Series* 439, 181–192.  
1146  
1147  
1148  
1149 Tecchio S., Ramírez-Llodra E., Aguzzi J., Flexas M.M., Company J.B., Sardà F. 2013.  
1150 Seasonal fluctuations of deep megabenthos: Finding evidence of standing stock accumulation  
1151 in a flux-rich continental slope. *Progress in Oceanography* 118, 188-198.  
1152  
1153  
1154  
1155  
1156 Thomsen L., Aguzzi J., Costa C., De Leo F., Ogston A., Purser A. 2017. The oceanic  
1157 biological pump: rapid carbon transfer to depth at continental margins during Winter.  
1158 *Scientific Reports* | 7: 10763 | DOI:10.1038/s41598-017-11075-6.  
1159  
1160  
1161  
1162 Thistle D., Levin, L.A. 1998. The effect of the experimentally increased near-bottom flow on  
1163 metazoan meiofauna at the deep-sea site, with comparison data on macrofauna. *Deep Sea*  
1164 *Research I* 45, 625-685.  
1165  
1166  
1167  
1168 Tselepides A., Papadopoulou K.N., Podaras D., Plaiti W., Koutsoubas D. 2000. Macrobenthic  
1169 community structure over the continental margin of Crete (South Aegean Sea, NE  
1170 Mediterranean). *Prog Oceanogr* 46: 401–428.  
1171  
1172  
1173  
1174  
1175 Vetter E.W., Dayton P.K. 1998. Macrofaunal communities within and adjacent to a detritus-  
1176 rich submarine canyon system. *Deep Sea Research II* 45, 25–54.  
1177  
1178  
1179  
1180 Vetter E.W., Dayton P.K. 1999. Organic enrichment by macrophyte detritus, and abundance  
1181 patterns of megafaunal populations in submarine canyons. *Marine Ecology Progress Series*  
1182 186, 137–148.  
1183  
1184  
1185  
1186  
1187  
1188  
1189  
1190  
1191  
1192  
1193  
1194  
1195  
1196  
1197  
1198  
1199  
1200

1201  
1202  
1203 **Figure captions**  
1204

1205 Figure 1. Location of the sampling stations in the three sites of Western and Central  
1206 Mediterranean Sea. Black circles: open slope stations, grey circles: canyon stations.  
1207

1208  
1209 Figure 2. The multidimensional scaling (MDS) applied on presence/absence data of the three  
1210 regions considered. The upper panel represents the analysis of all the data, in the lower panel  
1211 are presented the results of analysis excluding the three outliers.  
1212  
1213

1214  
1215 Figure 3. Results of the SIMPER analysis between the three regions and characteristic taxa .  
1216  
1217

1218 Figure 4. Average values  $\pm$  standard error of A: abundance, B: number of taxa, C: biomass.  
1219

1220 Fig. 5. Linear regression between depth and macrofaunal parameters in the three sites. A:  
1221 abundance, B: number of taxa and C: biomass.  
1222  
1223

1224 Fig. 6. Relative abundance of the trophic groups in the assemblages of the two topographic  
1225 habitats (canyon and open slope) in the three sites studied. Susp: suspension feeders, dep-  
1226 susp: deposit-suspension feeders, dep: deposit feeders, dep-graz: deposit feeders-grazers, dep-  
1227 pred: deposit feeders-predators, pred: predators.  
1228  
1229  
1230  
1231

1232 Fig. 7. Relationship between organism individual size vs longitude (A) and vs depth (B).  
1233 Black triangles represent the average value; standard error is indicated, when not visible it is  
1234 included in the marker size.  
1235  
1236  
1237  
1238  
1239  
1240  
1241  
1242  
1243  
1244  
1245  
1246  
1247  
1248  
1249  
1250  
1251  
1252  
1253  
1254  
1255  
1256  
1257  
1258  
1259  
1260

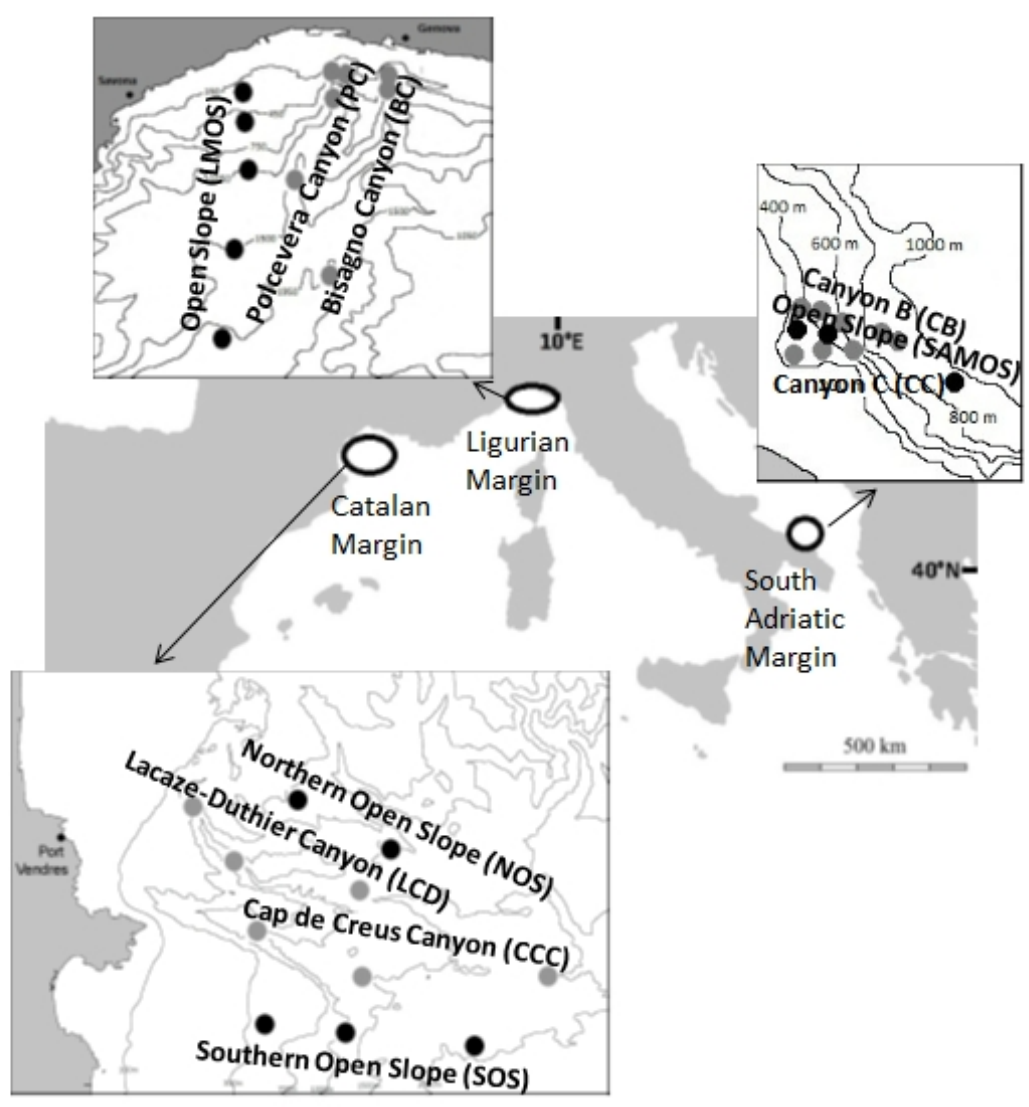
1261  
1262  
1263 **Supporting Material**  
1264

1265 Table S1. Complete list of taxa found in the three studied areas.  
1266

1267 Table S2. Species contributions, in percent, to the dissimilarity ( $\beta$ -diversity), between canyon  
1268 and open slope within each site.  
1269

1270 Table S3. Result of the regression analysis between the contribute of the mud fraction to the  
1271 sediment composition and the principal macrofaunal parameters.  
1272  
1273  
1274  
1275  
1276  
1277  
1278  
1279  
1280  
1281  
1282  
1283  
1284  
1285  
1286  
1287  
1288  
1289  
1290  
1291  
1292  
1293  
1294  
1295  
1296  
1297  
1298  
1299  
1300  
1301  
1302  
1303  
1304  
1305  
1306  
1307  
1308  
1309  
1310  
1311  
1312  
1313  
1314  
1315  
1316  
1317  
1318  
1319  
1320

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55

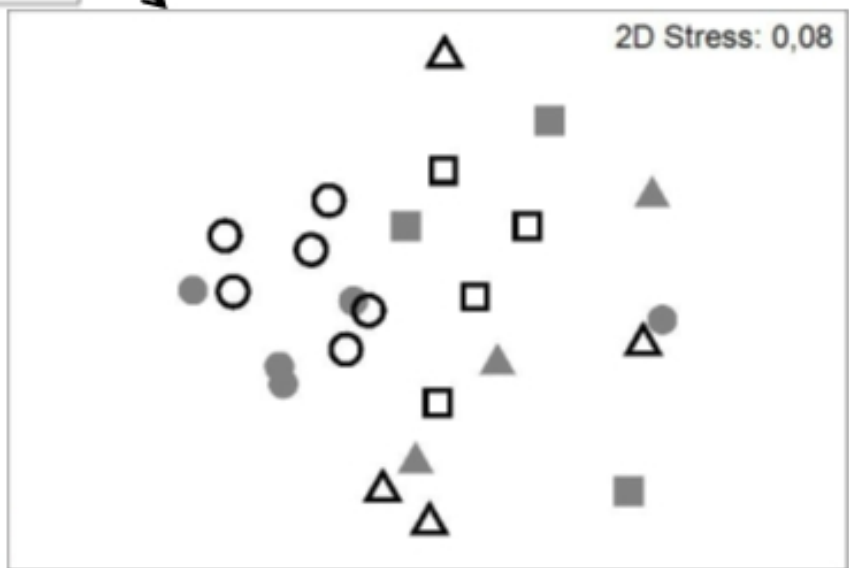


1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36

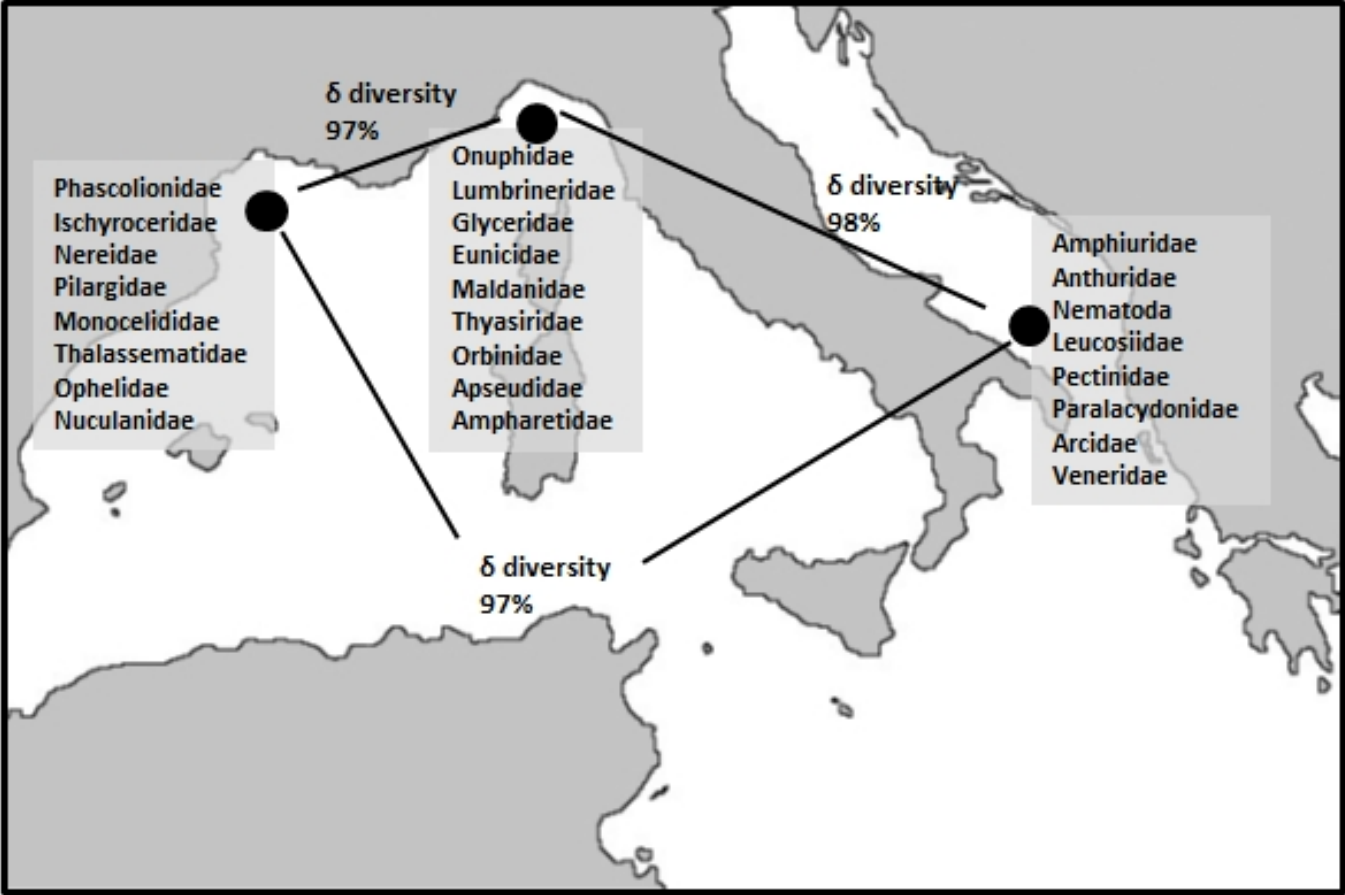


Transform: Presence/absence  
Resemblance: S17 Bray Curtis similarity

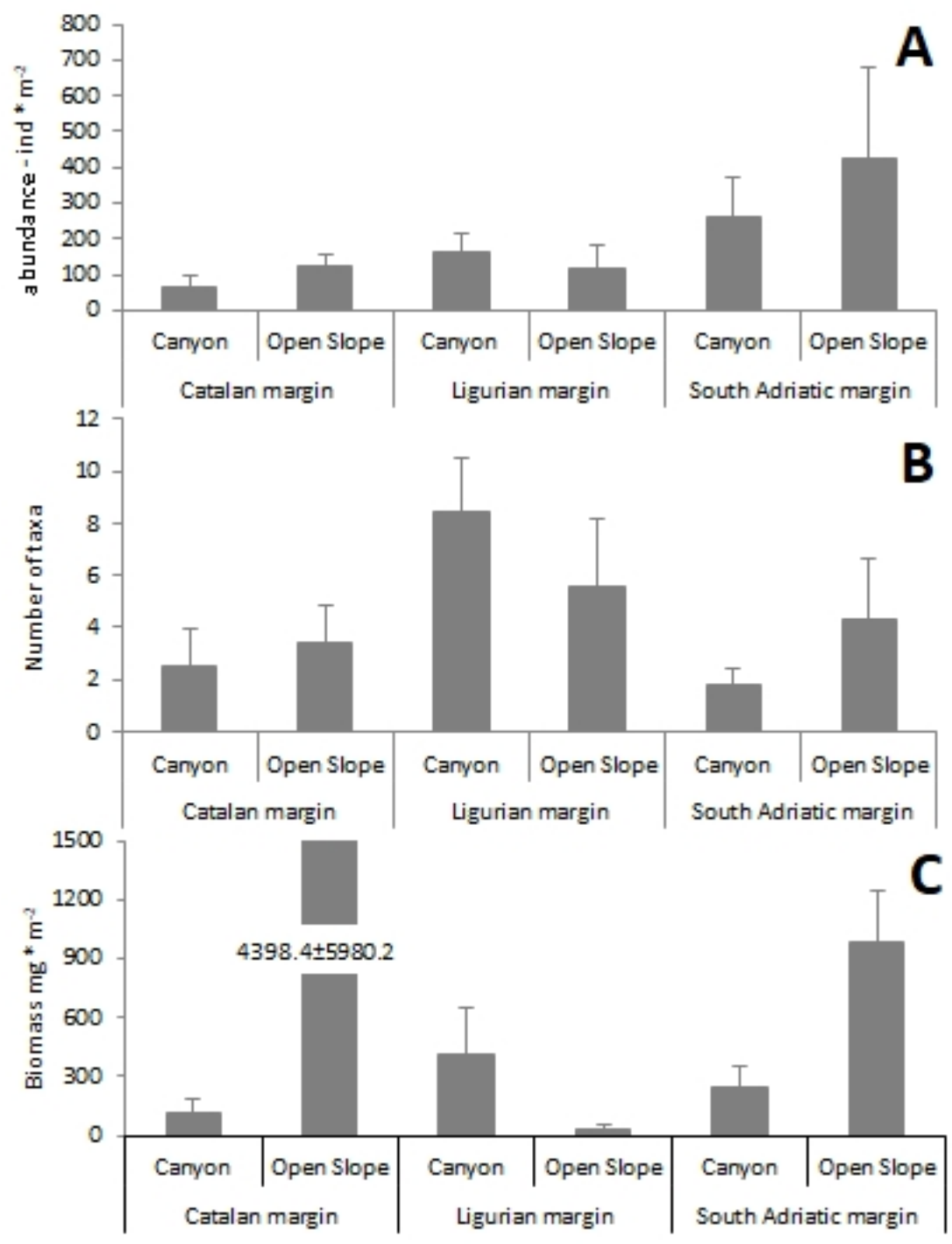
- Ligurian Margin Canyon
- Ligurian Margin Open Slope
- Catalan Margin Canyon
- Catalan Margin Open Slope
- △ South Adriatic Margin Canyon
- ▲ South Adriatic Margin Open Slope



1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36

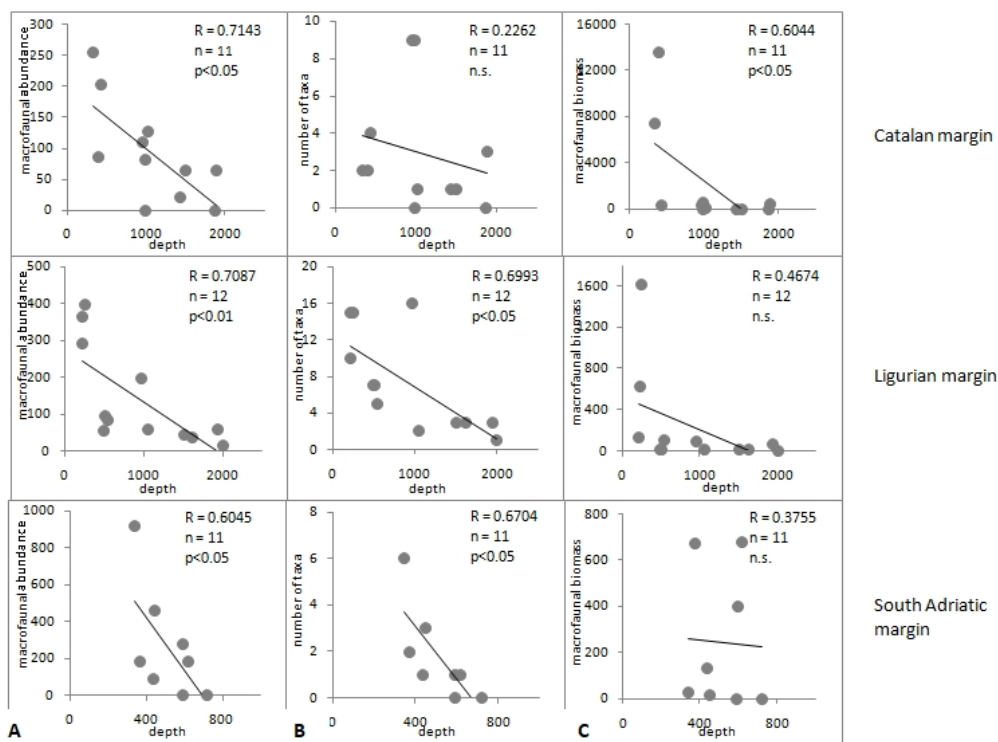


1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55

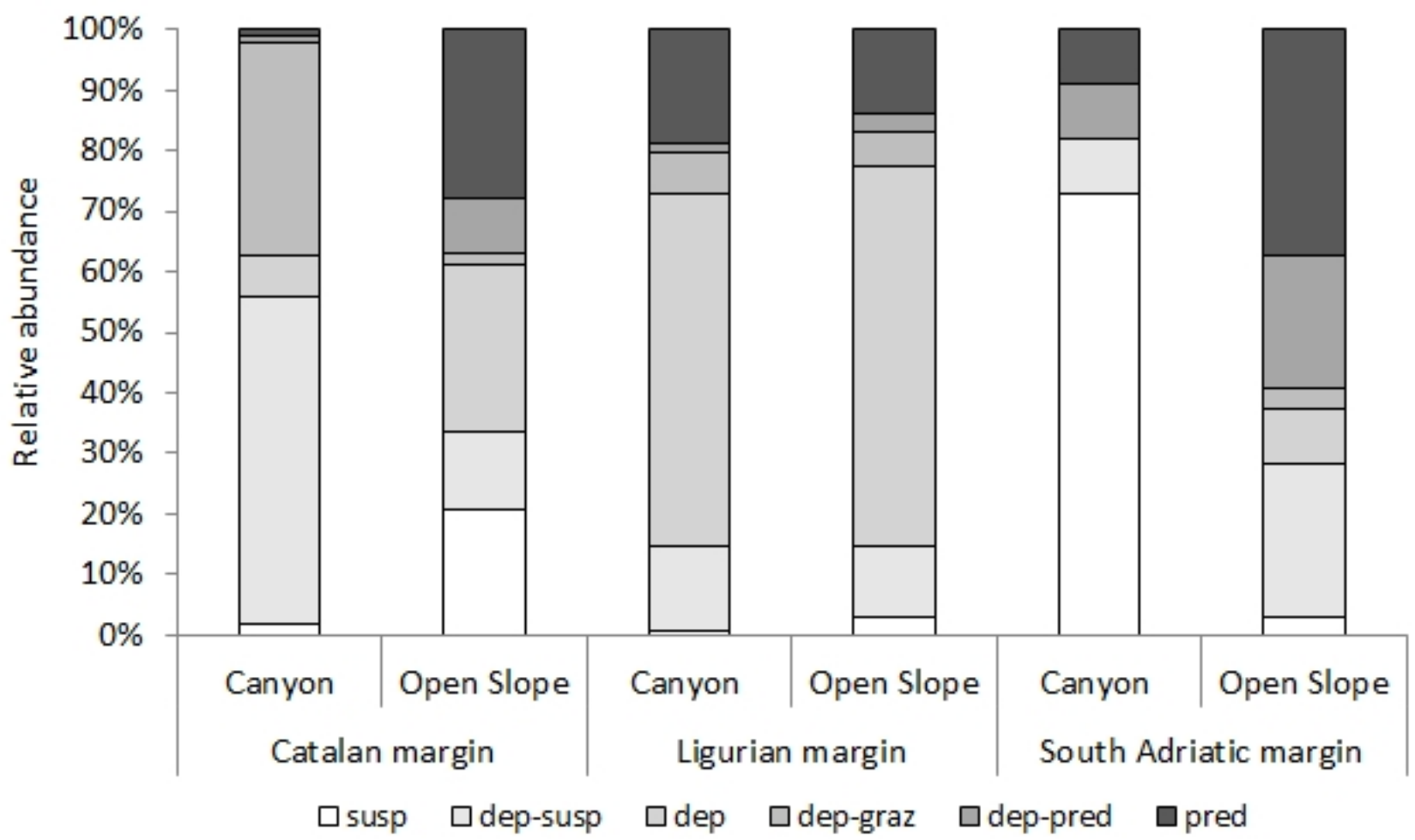




1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55



1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36



1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55

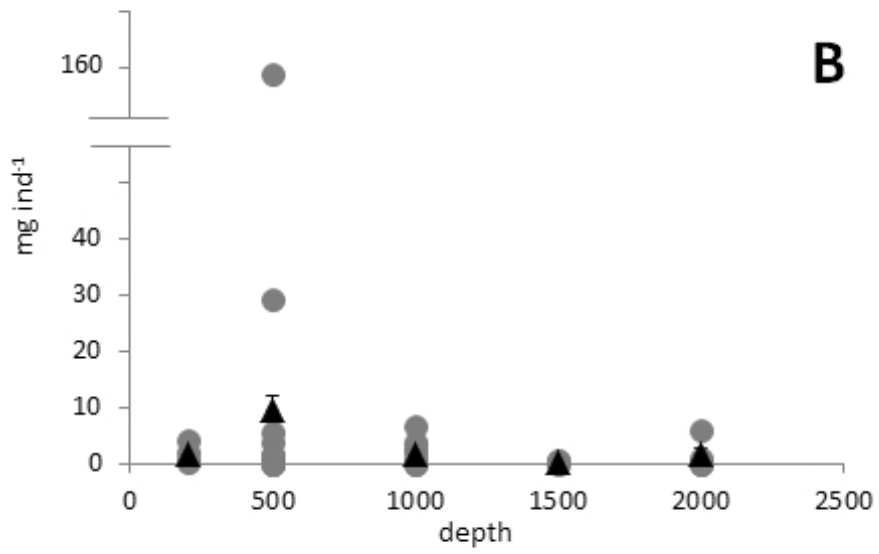
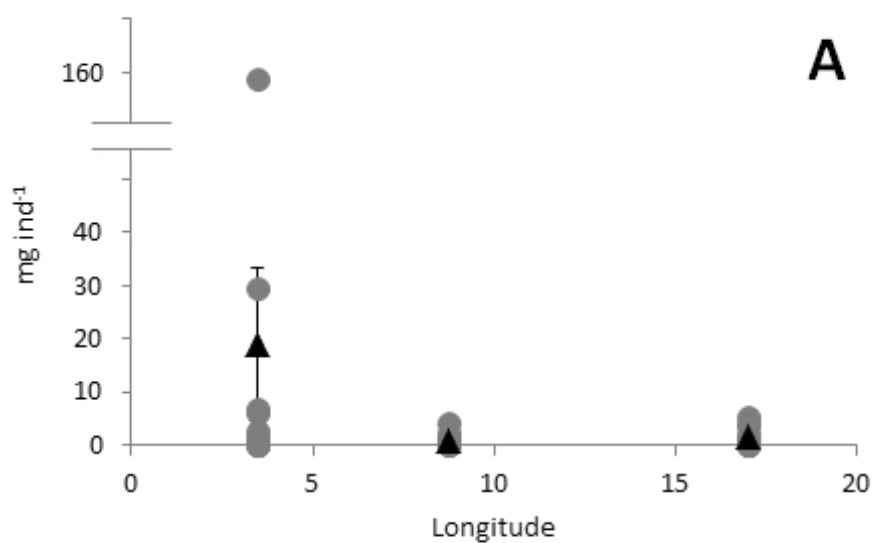


Table 1. Station depth, geographic coordinates, sampling device and percentage of mud in the sediment texture. N.d.: not determined.

		Depth (m)	Latitude N	Longitude E	Sampling device	Mud %
Catalan margin	Southern Open Slope (SOS)	398	42° 08.85'	03° 35.06'	multi-corer	69
		985	42° 07.72'	03° 46.63'		87
		1887	42° 07.05'	04° 02.74'		77
R/V Universitatis October 2005	Cap de Creus Canyon (CCC)	960	42° 18.47'	03° 36.60'		80
		1434	42° 12.64'	03° 49.22'		86
R/V Thetys II August 2006	Lacaze-Duthiers Canyon (LDC)	1870	42° 12.88'	04° 15.43'		68
		434	42° 34.44'	03° 24.04'		76
		990	42° 26.56'	03° 31.83'		80
		1497	42° 21.96'	03° 49.41'		56
	Northern Open Slope (NOS)	334	42° 34.13'	03° 39.19'		71
		1022	42° 26.49'	03° 51.32'		77
Ligurian margin	Open Slope (LMOS)	222	44° 18.79'	08° 40.22'	box-corer	98
		507	44° 16.22'	08° 40.30'		99
		1054	44° 11.37'	08° 40.52'		99
R/V Minerva Uno May 2013		1516	44° 03.40'	08° 39.16'		99
		2005	43° 55.02'	08° 38.17'		96
	Polcevera Canyon (PC)	252	44° 21.87'	08° 49.33'		99
		540	44° 21.23'	08° 50.19'		n.d.
		963	44° 18.72'	08° 49.88'		94
		1623	44° 10.58'	08° 45.71'		99
	Bisagno Canyon (BC)	225	44° 21.13'	08° 54.74'		96
		496	44° 20.12'	08° 54.64'		92
		1946	44° 00.71'	08° 49.10'		98
South Adriatic margin	Canyon B (CB)	370	41° 22.10'	17° 06.70'	multi-corer	39
		446	41° 21.71'	17° 07.75'		65
		590	41° 20.63'	17° 11.02'		19
N/O Urania May 2006	Open Slope (SAMOS)	196	41° 21.30'	17° 05.96'		11
		406	41° 20.08'	17° 10.32'		44
	Canyon C (CC)	908	41° 13.70'	17° 35.15'		68
		341	41° 19.07'	17° 05.15'		1
		435	41° 19.47'	17° 09.75'		63
		593	41° 18.18'	17° 12.51'		57
		618	41° 18.84'	17° 14.66'		48
		721	41° 18.43'	17° 15.61'		58

Table 2. Results of Analysis of similarities (ANOSIM), applied on the presence/absence data, between sites (Catalan margin, Ligurian margin and South Adriatic margin) and between Canyon and Open Slope inside each site. A): abundance of taxa matrix, B): abundance of trophic groups matrix.

<b>A)</b>		R	p
Catalan margin vs Ligurian margin		0.409	0.001
Catalan margin vs South Adriatic margin		0.150	0.018
Ligurian margin vs South Adriatic margin		0.505	0.001
<hr/>			
Catalan margin			
	Canyon vs Open Slope	-0.066	0.667
Ligurian margin			
	Canyon vs Open Slope	0.142	0.138
South Adriatic margin			
	Canyon vs Open Slope	-0.142	0.786
<hr/>			
<b>B)</b>		R	p
Catalan margin vs Ligurian margin		0.222	0.005
Catalan margin vs South Adriatic margin		0.072	0.186
Ligurian margin vs South Adriatic margin		0.392	0.002
<hr/>			
Catalan margin			
	Canyon vs Open Slope	0.263	0.079
Ligurian margin			
	Canyon vs Open Slope	0.058	0.290
South Adriatic margin			
	Canyon vs Open Slope	0.194	0.190

119  
120  
121  
122  
123  
124  
125  
126  
127  
128  
129  
130  
131  
132  
133  
134  
135  
136  
137  
138  
139  
140  
141  
142  
143  
144  
145  
146  
147  
148  
149  
150  
151  
152  
153  
154  
155  
156  
157  
158  
159  
160  
161  
162  
163  
164  
165  
166  
167  
168  
169  
170  
171  
172  
173  
174  
175  
176  
177

Table 3. Station depth, total abundance (mean±standard error), number of taxa and percentage of the contribution of the main taxa to the total abundance. An: Annelida, Cr: Crustacea, Mo: Mollusca, Si: Sipuncula, Ec: Echinodermata, Ot: others.

		Depth	Total abundance ind * m <sup>-2</sup>	N. taxa	An %	Cr %	Mo %	Si %	Ec %	Ot %
Catalan margin	Southern Open Slope (SOS)	398	84.9±11.5	2	50	50	0	0	0	0
		985	81.2±44.4	9	48	4	9	26	0	13
		1887	63.7±21.2	3	33.34	0	0	33.32	0	33.34
	Cap de Creus Canyon (CCC)	960	109.4±17.9	9	56	15	21	8	0	0
		1434	21.2±11.5	1	100	0	0	0	0	0
		1870	0	0	0	0	0	0	0	0
	Lacaze-Duthiers Canyon (LDC)	434	203.7±28.9	4	62	0	13	25	0	0
		990	0	0	0	0	0	0	0	0
		1497	63.7±17.3	1	100	0	0	0	0	0
	Northern Open Slope (NOS)	334	254.6±28.9	2	50	0	0	0	50	0
1022		127.3±23.1	1	0	100	0	0	0	0	
Ligurian margin	Open Slope (LMOS)	222	364.6±57.2	15	85	11	2	0	2	0
		507	95.1±41.2	7	83	17	0	0	0	0
		1054	59.5±11.7	2	80	0	0	20	0	0
		1516	44.6±14.9	3	67	33	0	0	0	0
		2005	15.0±15.0	1	100	0	0	0	0	0
	Polcevera Canyon (PC)	252	396.4±97.4	15	76	10	8	2	4	0
		540	83.2±54.8	5	57	29	14	0	0	0
		963	198.2±28.6	16	56	24	0	20	0	0
	Bisagno Canyon (BC)	1623	38.6±8.9	3	100	0	0	0	0	0
		225	291.3±48.7	10	85	7	8	0	0	0
South Adriatic margin	Canyon B (CB)	496	55.5±28.6	7	72	14	14	0	0	0
		1946	57.7±57.7	3	33	67	0	0	0	0
	Open Slope (SAMOS)	370	183.0±83.3	2	50	0	50	0	0	0
		446	457.9±183.3	3	20	0	0	20	60	0
		590	0	0	0	0	0	0	0	0
		196	2747.3±1066.7	9	23	47	0	7	10	13
		406	183.0±57.7	2	50	0	0	0	50	0
		908	274.7±126.7	2	67	33	0	0	0	0
Canyon C (CC)	341	915.8±366.7	6	10	0	70	0	0	20	
	435	91.5±46.2	1	0	100	0	0	0	0	
	593	274.5±144.3	1	0	0	0	0	100	0	
	618	183.0±86.6	1	0	0	0	0	100	0	
		721	0	0	0	0	0	0	0	

178  
 179  
 180  
 181  
 182  
 183  
 184  
 185  
 186  
 187  
 188  
 189  
 190  
 191  
 192  
 193  
 194  
 195  
 196  
 197  
 198  
 199  
 200  
 201  
 202  
 203  
 204  
 205  
 206  
 207  
 208  
 209  
 210  
 211  
 212  
 213  
 214  
 215  
 216  
 217  
 218  
 219  
 220  
 221  
 222  
 223  
 224  
 225  
 226  
 227  
 228  
 229  
 230  
 231  
 232  
 233  
 234  
 235  
 236

Table 4. Station depth, total biomass (mean±standard error) and percentage of the contribution of the main taxa to the total biomass. An: Annelida, Cr: Crustacea, Mo: Mollusca, Si: Sipuncula, Ec: Echinodermata, Ot: others.

		Depth	Total biomass mg *m <sup>-2</sup>	An %	Cr %	Mo %	Si %	Ec %	Ot %
Catalan margin	Southern Open Slope (SOS)	398	13546.8±7505.6	1	99	0	0	0	0
		985	545.2±397.4	15	1	0	2	0	82
		1887	385.6±172.5	8	0	0	28	0	64
	Cap de Creus Canyon (CCC)	960	314.9±187.9	51	16	1	32	0	0
		1434	12.4±6.4	100	0	0	0	0	0
		1870	0	0	0	0	0	0	0
	Lacaze-Duthiers Canyon (LDC)	434	355.2±173.2	85	0	9	6	0	0
		990	0	0	0	0	0	0	0
		1497	1.9±0.6	100	0	0	0	0	0
	Northern Open Slope (NOS)	334	7471.4±692.8	0.03	0	0	0	99.97	0
1022		43.3±23.1	0	100	0	0	0	0	
Ligurian margin	Open Slope (LMOS)	222	132.6±67.3	94	2	4	0	0	0
		507	11.6±7.8	62	38	0	0	0	0
		1054	21.2±12.7	81	0	0	19	0	0
		1516	23.2±19.6	84	16	0	0	0	0
		2005	0.1±0.1	100	0	0	0	0	0
	Polcevera Canyon (PC)	252	1620.4±885.7	45	53	0	0	2	0
		540	108.9±82.6	12	86	2	0	0	0
		963	93.8±31.1	85	2	0	13	0	0
	Bisagno Canyon (BC)	1623	17.5±14.9	100	0	0	0	0	0
		225	628.1±79.3	99	1	0	0	0	0
496		13.5±9.8	97	1	2	0	0	0	
1946	65.6±65.6	37	63	0	0	0	0		
South Adriatic margin	Canyon B (CB)	370	675.4±115.5	76	0	24	0	0	0
		446	17.4±15.6	6	0	0	21	73	0
		590	0	0	0	0	0	0	0
	Open Slope (SAMOS)	196	1435.8±1132.1	4	95	0	0.4	0.5	0.1
		406	999.3±346.4	57	0	0	0	43	0
		908	527.4±317.5	7	93	0	0	0	0
	Canyon C (CC)	341	27.7±20.2	3	0	95	0	0	2
		435	131.8±49.1	0	100	0	0	0	0
		593	400.8±109.7	0	0	0	0	100	0
		618	680.9±173.2	0	0	0	0	100	0
721	0	0	0	0	0	0	0		

## Supporting Material

Table S1. Complete list of taxa found in the three studied areas. The taxonomical level used for statistical analysis is reported in bold. Und.: undetermined or damaged specimens that cannot be determined for their conditions; juv.: juvenile specimens; “+ and -”: are for present and absent.

	Catalan margin		Ligurian margin		South Adriatic margin	
	Canyon	Open Slope	Canyon	Open Slope	Canyon	Open Slope
Phylum Arthropoda						
Subphylum Crustacea						
Class Malacostraca						
Order Decapoda						
Infraorder Caridea						
<b>Caridea und.</b>	-	-	+	-	-	-
<b>Family Acanthephyridae</b>						
<i>Acanthephyra eximia</i>	-	-	-	-	-	+
Infraorder Brachyura						
<b>Family Leucosiidae</b>						
<i>Ebalia</i> sp.	-	-	-	-	-	+
<b>Family Xanthidae</b>						
<i>Xantho pilipes</i>	-	+	-	-	-	+
Infraorder Gebiidea						
<b>Family Thalassinidae</b>						
Order Amphipoda						
Amphipoda und.	-	-	+	-	-	-
Infraorder Lysianassida						
<b>Family Ampeliscidae</b>						
<i>Ampelisca</i> sp.	-	-	-	+	-	-
<b>Family Phoxocephalidae</b>						
<i>Harpinia</i> cf. <i>dellavallei</i>	-	-	+	-	-	-
<i>Harpinia truncata</i>	-	-	-	+	-	-
<i>Paraphoxus oculatus</i>	-	-	-	+	-	-
<b>Family Urothoidae</b>						
<i>Urothoe</i> cf. <i>elegans</i>	-	-	+	+	-	-
Infraorder Corophiida						
<b>Family Ischyroceridae</b>						
<i>Jassa marmorata</i>	-	+	-	-	-	-
<b>Family Corophiidae</b>						
<i>Leptocheirus</i> cf. <i>mariae</i>	-	-	+	-	-	-
Infraorder Hadziida						
<b>Family Maeridae</b>						
<i>Othomaera schmidtii</i>	-	-	-	+	-	-
Infraorder Amphilochida						
<b>Family Oedicerotidae</b>						
<i>Westwoodilla caecula</i>	-	-	+	-	-	-



60  
61  
62  
63  
64  
65  
66  
67  
68  
69  
70  
71  
72  
73  
74  
75  
76  
77  
78  
79  
80  
81  
82  
83  
84  
85  
86  
87  
88  
89  
90  
91  
92  
93  
94  
95  
96  
97  
98  
99  
100  
101  
102  
103  
104  
105  
106  
107  
108  
109  
110  
111  
112  
113  
114  
115  
116  
117  
118

Order Isopoda						
Superfamily Anthuroidea						
<b>Family Antheluridae</b>						
<i>Pilosanthura cf. fresii</i>	-	-	+	-	-	-
<b>Family Anthuridae</b>	-	-	-	-	+	-
Superfamily Cymothooidea						
<b>Family Cirolanidae</b>						
<i>Natatolana borealis</i>	+	-	+	-	-	-
<b>Infraorder Epicaridea</b>	+	-	-	-	-	-
Order Cumacea						
Cumacea und.	+	-	-	-	-	-
<b>Family Lampropidae</b>						
<i>Platysympus typicus</i>	-	-	+	-	-	-
<b>Family Leuconidae</b>						
<i>Leucon (Epileucon) longirostris</i>	-	-	-	+	-	-
Order Tanaidacea						
Superfamily Apseudoidea						
<b>Family Apseudidae</b>						
<i>Fageapseudes retusifrons</i>	-	-	-	+	-	-
Class Hexanauplia						
Subclass Copepoda	-	-	+	+	-	+
Class Ostracoda						
Order Podocopida						
<b>Family Cyprididae</b>	-	-	+	-	-	-
Phylum Annelida						
Class Polychaeta						
Polychaeta und.	+	-	-	+	+	+
Subclass Sedentaria						
Infraclass Canalipalpata						
<b>Canalipalpata und.</b>	+	+	-	-	+	-
<b>Family Spionidae</b>	-	-	+	+	-	+
<b>Family Ampharetidae</b>	-	-	+	+	-	-
<b>Family Terebellidae</b>	+	-	-	-	-	-
<i>Polycirrus</i> sp.	-	-	-	-	-	+
Infraclass Scolecida						
<b>Family Capitellidae</b>	-	+	+	+	-	+
<i>Notomastus</i> sp.	+	-	-	-	-	-
<b>Family Maldanidae</b>	-	-	+	+	-	-
<b>Family Ophelidae</b>	+	-	-	-	-	-
<b>Family Orbinidae</b>	-	-	+	-	-	-
<b>Family Paraonidae</b>	-	+	+	+	-	-
<b>Family Chaetopteridae</b>	-	+	-	-	-	-
Subclass Echiura						
Order Echiuroidea						
<b>Family Thalassematidae</b>						

119  
120  
121  
122  
123  
124  
125  
126  
127  
128  
129  
130  
131  
132  
133  
134  
135  
136  
137  
138  
139  
140  
141  
142  
143  
144  
145  
146  
147  
148  
149  
150  
151  
152  
153  
154  
155  
156  
157  
158  
159  
160  
161  
162  
163  
164  
165  
166  
167  
168  
169  
170  
171  
172  
173  
174  
175  
176  
177

<i>Maxmuelleria gigas</i>	-	+	-	-	-	-
Subclass Errantia						
Order Eunicida						
<b>Family Eunicidae</b>	-	-	+	+	-	-
<b>Family Lumbrineridae</b>	-	-	+	+	-	-
<b>Family Onuphidae</b>	-	-	+	+	-	-
Order Phyllodocida						
<b>Family Acoetidae</b>	-	-	+	-	-	-
<b>Family Sigalionidae</b>	-	-	+	-	-	-
<b>Family Glyceridae</b>	-	-	+	+	-	-
<b>Family Paralacydonidae</b>	-	-	-	-	+	-
<b>Family Pilargidae</b>						
<i>Ancistrosyllis cf. groenlandica</i>	-	+	-	-	-	-
<b>Family Nereididae</b>	-	+	-	-	-	-
<b>Family Syllidae</b>	-	-	-	+	-	-
<b>Family Phyllodocidae</b>	-	-	+	-	-	-
Class Clitellata						
Subclass Oligochaeta	+	-	+	+	-	+
Phylum Mollusca						
Class Bivalvia						
Order Pectinida						
<b>Family Pectinidae</b>						
<i>Aequipecten opercularis</i>	-	-	-	-	+	-
Order Arcida						
<b>Family Arcidae</b>						
<i>Bathyarca philippiana</i>	-	-	-	-	+	-
Order Venerida						
<b>Family Veneridae</b>						
<i>Clausinella fasciata</i>	-	-	-	-	+	-
<i>Pitar rude</i>	-	-	-	-	+	-
<i>Lajonkairia lajonkairii</i>	-	-	-	-	+	-
Superfamily Galionmatoidea						
<b>Family Lasaidae</b>						
<i>Kellia suborbicularis</i>	+	-	-	-	-	-
Order Nuculida						
<b>Family Nuculidae</b>						
<i>Ennucula tenuis</i>	-	-	+	+	-	-
<i>Nucula sulcata</i>	+	-	+	-	-	-
Order Nuculanida						
<b>Family Nuculanidae</b>						
<i>Nuculana fragilis</i>	-	+	-	-	-	-
<i>Nuculana</i> sp. juv.	+	-	-	-	-	-
Order Lucinida						
<b>Family Thyasiridae</b>						
<i>Thyasira flexuosa</i>	-	-	+	-	-	-

178  
179  
180  
181  
182  
183  
184  
185  
186  
187  
188  
189  
190  
191  
192  
193  
194  
195  
196  
197  
198  
199  
200  
201  
202  
203  
204  
205  
206  
207  
208  
209  
210  
211  
212  
213  
214  
215  
216  
217  
218  
219  
220  
221  
222  
223  
224  
225  
226  
227  
228  
229  
230  
231  
232  
233  
234  
235  
236

Phylum Echinodermata						
Class Ophiuroidea						
Ophiuridea juv.	-	-	-	+	-	-
Order Amphilepidida						
<b>Family Amphiuridae</b>						
<i>Amphiura chiajei</i>	-	-	-	-	-	+
<i>Amphiura filiformis</i>	-	-	-	-	+	+
Order Ophiurida						
<b>Family Ophiuridae</b>						
<i>Ophiura lacertosa</i>	-	-	+	-	-	-
Class Echinoidea						
Subclass Euechinoidea	-	+	-	-	-	-
Phylum Sipuncula					-	
Sipuncula und.	-	-	-	-	-	+
Class Sipunculidea						
Order Golfingiida						
<b>Family Golfingiidae</b>						
<i>Golfingia (Golfingia) elongata</i>	-	-	+	-	-	-
<i>Golfingia</i> sp.	-	+	-	-	-	-
<i>Nephasoma (Nephasoma) constrictum</i>	-	-	+	-	-	-
<b>Family Phascolionidae</b>						
<i>Onchnesoma steenstrupii steenstrupii</i>	+	+	-	-	-	-
<i>Onchnesoma</i> sp.	+	-	-	-	-	-
Class Phascolosomatidea						
Order Phascolosomatida						
<b>Family Phascolosomatidae</b>						
<i>Phascolosoma (Fisherana) capitatum</i>	-	+	-	-	-	-
<i>Phascolosoma (Phascolosoma) agassizii</i>	-	-	+	-	+	-
<i>Phascolosoma (Phascolosoma) granulatum</i>	-	-	+	+	-	-
Phylum Platyhelminthes						
Class Rhabditophora						
Order Prosejata						
<b>Family Monocelididae</b>						
<i>Boreocelis</i> sp.	-	+	-	-	-	-
Phylum Nematoda	-	-	-	-	+	+
Phylum Nemertina	-	+	-	-	-	-

237  
238  
239  
240  
241  
242  
243  
244  
245  
246  
247  
248  
249  
250  
251  
252  
253  
254  
255  
256  
257  
258  
259  
260  
261  
262  
263  
264  
265  
266  
267  
268  
269  
270  
271  
272  
273  
274  
275  
276  
277  
278  
279  
280  
281  
282  
283  
284  
285  
286  
287  
288  
289  
290  
291  
292  
293  
294  
295

Table S2. Percentage contribution of the various taxa to the dissimilarity ( $\beta$ -diversity) between canyon and open slope within each sampling site. Rel. abund.: relative abundance; Cum. cont.: cumulative contribution; 0: absence; “+”: higher abundance; “-“: lower abundance.

Catalan margin			
$\beta$ -diversity = 95%			
	Open Slope	Canyon	
	Rel. abund.	Rel. abund.	Cum. cont.%
Oligochaeta	0	+	11.70
Ischyroceridae	+	0	19.96
Phascolionidae	-	+	27.98
Capitellidae	-	+	35.40
Euechinoidea	+	0	41.05
Pilargidae	+	0	46.70
Nereidae	+	0	51.74
Xanthidae	+	0	56.79
Terebellidae	0	+	61.70
Monocelididae	+	0	65.40
Thalassematidae	+	0	69.11
Nuculidae	0	+	72.58
Ophelidae	0	+	75.57
Canalipalpata	-	+	78.41
Nuculanidae	0	+	80.68
Ligurian margin			
$\beta$ -diversity = 82%			
	Open Slope	Canyon	
	Rel. abund.	Rel. abund.	Cum. cont.%
Onuphidae	+	-	8.41
Paraonidae	-	+	15.72
Lumbrineridae	-	+	21.17
Maldanidae	+	-	26.32
Eunicidae	+	-	30.67
Nuculidae	+	-	34.64
Glyceridae	-	+	38.44
Phascolosomatidae	+	-	42.04
Thyasiridae	+	0	45.64
Capitellidae	+	-	48.61
Syllidae	0	+	51.34
Caridea	+	0	54.01
Apseudidae	0	+	56.65
Orbinidae	+	0	59.27
Ampharetidae	+	-	61.88
Cirolanidae	+	0	64.48
Oligochaeta	+	-	67.01
Sigalionidae	+	0	69.26

296  
297  
298  
299  
300  
301  
302  
303  
304  
305  
306  
307  
308  
309  
310  
311  
312  
313  
314  
315  
316  
317  
318  
319  
320  
321  
322  
323  
324  
325  
326  
327  
328  
329  
330  
331  
332  
333  
334  
335  
336  
337  
338  
339  
340  
341  
342  
343  
344  
345  
346  
347  
348  
349  
350  
351  
352  
353  
354

Copepoda	+	-	71.39
Spionidae	-	+	73.25
Cypridinidae	+	0	75.09
Phoxocephalidae	-	+	76.92
Urothoidae	-	+	78.63
Ophiuridae	-	+	80.32
South Adriatic margin			
$\beta$ -diversity = 88%			
	Open Slope	Canyon	
	Rel. abund.	Rel. abund.	Cum. cont. %
Amphiuridae	+	-	16.08
Terebellidae	+	0	27.80
Oligochaeta	+	0	36.99
Leucosiidae	+	0	45.28
Acanthephyridae	+	0	52.39
Nematoda	+	-	58.48
Veneridae	0	+	63.07
Anthuridae	0	+	67.44
Spionidae	+	0	71.73
Canalipalpata	0	+	75.12
Pectinidae	0	+	78.51
Capitellidae	+	0	81.54

Table S3. Result of the regression analysis between the mud fraction to the sediment composition and the principal macrofaunal parameters. ns: not significant.

	R	n	p
<b>Catalan margin</b>			
Density vs Mud fraction	0.2588	9	ns
Number of taxa vs Mud fraction	0.4569	9	ns
Biomass vs Mud fraction	0.2717	9	ns
<b>Ligurian margin</b>			
Density vs Mud fraction	0.0265	10	ns
Number of taxa vs Mud fraction	0.2280	10	ns
Biomass vs Mud fraction	0.1517	10	ns
<b>South Adriatic margin</b>			
Density vs Mud fraction	0.5643	9	ns
Number of taxa vs Mud fraction	0.6192	9	< 0.05
Biomass vs Mud fraction	0.2149	9	ns