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Recolonization and recovery dynamics of the macrozoobenthos after sand extraction in relict sand bottoms of the Northern Adriatic Sea.

R. Simonini ^{a*}, I. Ansaloni ^a, P. Bonini ^a, V. Grandi ^a, F. Graziosi ^a, M. Iotti ^a, G. Massamba-N'Siala ^a, M. Mauri ^a, G. Montanari ^b, M. Preti ^c, N. De Nigris ^c, D. Prevedelli ^a

^a Dipartimento di Biologia Animale, Università degli studi di Modena e Reggio Emilia, via Campi 213/D, 41100 Modena, Italy

^b ARPA Emilia-Romagna, Struttura Oceanografica Daphne, via Vespucci, Cesenatico (FC), Italy

^c ARPA Emilia-Romagna, Ingegneria Ambientale, Vicolo Carega 10, Bologna, Italy

*Corresponding author

Simonini Roberto

Dipartimento di Biologia Animale, Università degli Studi di Modena e Reggio Emilia, Via Campi 213/D, 41100 Modena, Italy

Tel: +3959-2055562; fax: +3959-2055548

E-mail address: simonini.roberto@unimore.it

Abstract

The long-term effects of sand extraction on macrozoobenthic communities were investigated in an offshore area in the Northern Adriatic Sea characterised by relict sands formed during the last Adriatic post-glacial transgression. Surveys were carried out before, during and 1, 6, 12, 18, 24 and 30 months after extraction at three impacted and seven reference stations. The operations did not influence the physical characteristics of the sediment, but they caused almost complete defaunation at dredged sites. Univariate and multivariate analyses highlighted that the macrozoobenthic community responses to the dredging operations were 1) a rapid initial recolonisation phase by the dominant taxa present before dredging, which took place 6-12 months after sand extraction; 2) a slower recovery phase, that ended 30 months after the operations, when the composition and structure of the communities were similar in the dredged and reference areas. This pattern of recolonisation-recovery fits well with the commonly encountered scenario where the substratum merely remains unchanged after marine aggregate extraction.

Keywords: Environmental impact; Dredging; Sand extraction; Benthos; Recolonisation; Recovery; Northern Adriatic Sea; Relict sands

1. Introduction

The impact of marine sand extraction on the characteristics of the seabed and the changes in benthic fauna have been widely investigated (Newell, Seiderer, & Hitchcock, 1998; Sardà, Pinedo, Gremare, & Taboada, 2000; Van Dalfts, Essink, Toxvig Madsen, Birklund, Romero, & Manzanera, 2000). The dredging operations may influence the physical and biological characteristics of the impacted areas both directly, through removal, smothering and damage caused by the dredge head, and indirectly: surface and bottom plumes, changes in plankton bloom seasons, the release of nutrients and chemicals, as well as sound, can affect both the sea bottom and the water column in the immediate area around the dredging site (Boyd, Limpenny, Rees, & Cooper, 2005; Newell et al., 1998; Van Dalfts et al., 2000).

Most studies concerning the impacts of sand dredging on macrozoobenthos were performed in shallow, dynamic sandy bottoms. In these cases, the recovery of the benthic assemblages appeared to be linked to the alterations in the seabed, in terms of grain size, organic content and morphology induced by sand extraction. Major alterations in the sediment and grain size characteristics over a long period favoured the settlement of benthic communities that were different in composition and structure from the pre-operational ones and this difference could be maintained for a long time (Boyd, Limpenny, Rees, Cooper, & Campbell, 2003; Boyd et al., 2005; Van der Veer, Bergman, & Beukema, 1985). In contrast, when there was minimal variation in the seabed characteristics, benthic communities recovered within a few years in terms of biodiversity and biomass (Boyd et al., 2003; Kenny & Rees, 1994,

1996; Kenny, Rees, Greening, & Campbell, 1998; Robinson, Newell, Seiderer, & Simpson, 2005; Van Dalfsen & Essink, 1997).

The largest nourishment operation in the Mediterranean Sea took place in 2002, when approximately 800,000 m³ of sand were dredged from off-shore relict sand deposits in the Northern Adriatic Sea. These were destined for the nourishment of several beaches along the Emilia-Romagna coast (Italy) (Simonini et al., 2005). Besides the geographical location, this operation was unusual because the seabed of the area 1) did not show active sedimentation, 2) was primarily composed of relict sand, 3) was far from the coast (55 km) and 4) at a greater depth (40–42 m) compared with the other previous dredging operations, where the maximum dredging depth was usually around 30 metres (North Sea Foundation, 2005).

The analysis of the short-term impacts of sand extraction on sediment and macrozoobenthos during the first 12 months after dredging, highlighted that the activities did not significantly influence the granulometry and the total organic carbon content (%TOC) of the substratum, but caused almost complete defaunation at the dredging stations. Yet, at the end of this survey period, the recolonisation of the communities at the impacted stations was at an advanced stage (Simonini et al., 2005).

In the present paper, we analysed the impact of relict sand extraction on benthic communities and the degree of recolonisation of macrobenthos at the end of the monitoring program, up to 30 months after the completion of the dredging. The aim was to establish the recolonization-recovery state of the macrozoobenthos and obtain a reference model for the effects of sand extraction on relict sand bottoms.

2. Materials and methods

2.1. Study area and sampling

The dredged area is located approximately 55 Km off the coast of Ravenna, in an area characterized by the occurrence of relict offshore sandy deposits (Fig. 1; Preti, 2000). These deposits, mainly composed of sand and coarse detritus, are the remains of coastal structures that were formed during the last Adriatic marine transgression. This area is outside the main current circuit of the Northern Adriatic, and as a result is characterized by a limited amount of sedimentation of fine material during the summer period that become re-suspended and dispersed during the winter (Preti, 2000, 2002; Simonini et al., 2005). The tidal excursion is negligible and the currents are weak and with variable direction. In fact, the main factors that affect the sediment transport and re-suspension in the area are the storms driven by the NE and SE winds (Bora and Scirocco) that frequently occur in the Northern Adriatic Sea from December to late March and probably lead to significant sediment re-suspension (Matteucci and Frascari, 1997; Wang, 2002). Water temperature in the bottom showed marked seasonal variation ranging from 8°C in February to 26°C in August (Montanari & Pinardi, 2006).

The dredging operations took place in April-May 2002. The monitoring activities started in March 2001 and ended in December 2004. Eight sampling surveys were carried out: March 2001 (before extraction [B.-Ex.]), April 2002 (during extraction [Ex.]), June 2002 (1 month after extraction [A.-Ex. 1]), December 2002 (6 months after extraction [A.-Ex. 6]), June 2003 (12 months after extraction [A.-Ex. 12]),

December 2003 (18 months after extraction [A.-Ex. 18]), June 2004 (24 months after extraction [A.-Ex. 24]) and December 2004 (30 months after extraction [A.-Ex. 30]).

Because of the lack of information on the effects of sand extraction in the Northern Adriatic Sea, most of the efforts were devoted to the after-dredging surveys (7 survey), in order to analyze the major recolonization-recovery patterns of the macrozoobenthos, as opposed to the characterization on the temporal variability of community before dredging (1 survey).

Surveys and sampling activities were performed with the oceanographic M/S “Daphne II” of ARPA (Regional Agency for Prevention and the Environment of Emilia-Romagna). The sampling design provided for samples at three impacted stations within the extraction area and at seven reference stations outside it. The stations were positioned along two transects perpendicular to and intersecting each other in the centre of the dredging zone (Fig. 1). The reference sites were chosen at large enough distances so as not to be affected by the extraction activity and deposition of suspended sediment. The direction and extension of the plume during the extraction were monitored by ARPA Daphne. During the extraction the currents were weak and variable and the plume never reach the inner reference stations. Multibeam profile, Side Scan Sonar and video camera images were used to measure the depth of the excavation pits and holes.

The sampling for macrozoobenthos was carried out using a van Veen grab with a surface area of 1250 cm² and sampling volume of 24 dm³. Three replicates were collected at each station. For each station, analyses for sediment granulometry and %TOC were also performed (ICRAM, 2001).

The samples for macrozoobenthos analysis were sieved (mesh size 0.5 mm) and the retained material was preserved in 4% CaCO₃ buffered formalin in seawater.

2.2. Data analysis

The samples and the data were treated and analysed using the same techniques described in Simonini et al. (2005). The residual material obtained from the sample sieving was sorted in the laboratory and the macrofauna was preserved in 70% ethylic alcohol. Most of the organisms were identified to species level. Data from each sampling station were obtained from the sum of their respective replicates.

The data collected during the different surveys were integrated by obtaining a total matrix of the abundances and the main ecological indices (number of species/taxa, abundance and Shannon–Wiener diversity) were calculated.

Multivariate analyses were performed with original (untransformed) and 4th root transformed abundance data in order to analyze the patterns of variation of benthic assemblages in terms of numerically dominant species and species composition. Outputs from non-metric Multidimensional Scaling (nMDS) ordination models of the Bray-Curtis similarity matrix were obtained for all the stations/surveys. Differences among the extraction and reference stations during different surveys were tested using a two-way crossed ANOSIM (ANalysis Of SIMilarity) randomisation–permutation test. One-way ANOSIM was performed to compare *a posteriori* the extraction area stations with the reference stations, and the reference stations of the various surveys. The species making the greatest contribution to dissimilarity among the impacted and reference stations for the different surveys were investigated using the SIMilarity

PERcentage breakdown procedure (SIMPER). The analyses were carried out using PRIMER V5 software (Clarke & Warwick, 2001).

3. Results

3.1. Sediment characteristics

Both the reference and impacted sites did not differ for the sediment characteristics during the whole examined period. In all the samples analysed the percentage of sand varied from 85% to 98%, with the prevalence of fine sands (250-125 μm), and the %TOC was low (0.3–0.8%). The recorded sailing tracks of the dredger and the bathymetric changes in the sea bed after sand extraction revealed that the operations were carried out mainly in the southern and central parts of the dredged area, corresponding to sampling stations 4 and 3 (Fig. 1, 2). During the extraction period the currents were weak and directed toward S9, but the dredged-sediment plumes were confined to within 1 km of the sand extraction area. The reconstruction of the bottom profile based on multi-beam data collected immediately after the extraction showed that pits with a depth of 1.5 m had formed on the seabed of the impacted area, due to the extraction activities (Fig. 3). The surface area and the depth of these “holes” tended to decrease over time: their mean depth was about 0.30 m in the the A.-Ex. 24 months surveys (Fig. 3).

3.2. Taxonomic analysis

The taxonomic analysis of the samples collected during the eight surveys led to the identification of about 42000 individuals, belonging to 183 taxa and distributed

among 12 phyla (in order of abundance: annelids, arthropods, molluscs, nemertines, echinoderms, sipunculids, phoronids, poripherans, cnidarians, echiurians, chordates and hemichordates). The polychaetous annelids were the most abundant group (43% of the taxa, 59% of the individuals), followed by the molluscs (mainly bivalves and gastropods, 24% of the taxa and 4% of the individuals), the arthropods (mostly crustaceans, 18% of the taxa and 33% of the individuals) and the echinoderms (echinoids, stellaroids and ophiuroids, 9% of the taxa and 2% of the individuals). All the other phyla together represented 7% of the taxa, but only 1% of the individuals. The species were characteristic of offshore sandy–muddy bottoms. No mass settlement of opportunistic species or species typical of disturbed bottoms was observed in the impacted areas. The list of the taxa found during the whole monitoring program was reported in Appendix 1.

3.3. *Abundance data analyses*

The multivariate analyses performed on original and 4th root transformed data highlighted that the sand extraction affected the macrozoobenthic communities of the impacted stations during and after sand extraction, both in terms of species composition and of numerically dominant species (ANOSIM, Table 1).

The MDS ordination plots of the untransformed and 4th transformed data relating to each station and sampling date (Fig. 4a, b) showed the stations 3 and 4 of the Ex. and A.-Ex. 1 surveys to be segregated towards one side, while the reference and the dredging station of the last three A.-Ex. Surveys (A.-Ex. 18, A.-Ex. 24 and A.-Ex. 30 surveys) were grouped on the other side, very close to each other. Between these two

extremes, there were the points indicative of stations 2–4 of the A.-Ex. 6 and A.-Ex. 12 months surveys) (Fig. 4a, b). The MDS sub-plot of the impacted station from the different surveys (Fig. 4c, d) showed the same patterns as the whole plot.

In the Ex and A.-Ex. 1 survey, the average dissimilarities of impacted station with respect to the reference station and B-Ex Imp. stations were very high and significant (80-90% with untransformed data, 70-75% with 4th root transformed data) (Table 2-3). The stations 3 and 4 of the Ex. and A.-Ex. 1 surveys underwent almost complete defaunation after the sand extraction, evidenced by the drastic reduction in the number of taxa, abundances and Shannon-Wiener diversity with respect to reference stations (Table 2; Fig. 5). Station 2, located in the northern part of the extraction area, was subject to a lower impact and showed a smaller reduction in the number of taxa and abundances than station 3 and 4 (Table 4; Fig. 5).

The intermediate position of the stations 2–4 of the A.-Ex. 6 and A.-Ex. 12 surveys in the MDS plots of Fig. 3a-d suggested that recolonisation was taking place at these stations. In fact the average dissimilarity of impacted station in the Ex and A.-Ex. 1 survey, with respect to B-Ex Imp. stations was 63-83% (untransformed data) and 45-66% (4th root transformed data) (Table 2-3). Moreover, the dredged stations showed abundances that were about 40-50% lower than the reference stations, whereas during the A.-Ex. 1 survey the mean difference was about 90%. In addition, the number of taxa discovered at the extraction area stations was 60–70% of that observed at the reference stations, whereas during the surveys carried out during and 1 month after extraction this percentage was only 26% (Table 4; Fig. 5).

Although the reference sites of each survey are tightly grouped on one side of the global MDS plots (Fig. 4a, b), the MDS subplots of the reference station from the different surveys highlighted the great temporal variability of the community, throughout the eight surveys both in term of species composition and, overall, in term of numerically dominant species (Fig. 4 e-f). In particular, the MDS plots showed the reference station of the A-Ex 12 survey segregated separately and more dispersed than the others, due to the lower abundances and number of taxa found (Table 4).

With the exception of the A-Ex 12 survey, the dissimilarities among surveys were lower than could be ascribed to the impact of the extraction activity, ranging from 25 to 42% (4th root transformed data) and from 33 to 62% (untransformed data). Yet the results of the one-way ANOSIM tests between the references station of the eight surveys showed that the communities varied from one survey to another (Table 2, 3), probably because the high within-groups similarities (60-70% with untransformed data, 70-80% with 4th root transformed data). Most of the dissimilarities among the reference stations of different surveys were due to variations in the abundances of some of the more highly represented and/or characteristic species, and to the occurrence of low-density, rare species that characterize one or more surveys (SIMPER, Simonini et al, 2005). In fact only about the 50%-60% of the species found throughout the whole investigation were common to the reference station of all the surveys but these species represented 90-95% of the total abundance.

The high R-values suggested that the community of the B-Ex. Imp. station and the impacted station from the other survey are different from one survey to another, despite their dissimilarity decrease over time after the dredging (Table 2, 3). The

SIMPER analysis performed on the 4th root-transformed data from the B-Ex or the A-Ex 30 impacted stations has emphasized the role of some rare species with very low density (Appendix 2). The same analysis performed without data transformation highlighted that the differences in the communities of the impacted station before and 30 month after dredging were mostly due to variations in the abundances of the more highly represented and characteristic species of the relict sands, such as the polychaetes *Aricidea claudiae*, *Spiophanes kroyeri kroyeri*, *Monticellina dorsobranchialis* and *Jasmineira elegans*, and the crustaceans *Ampelisca diadema* and *Apseudes latreilli* (Appendix 2). As observed for the reference station of the different survey, the communities of the impacted station before and 30 month after sand extraction have a number (52%) of species in common. These species contributed to the 89% and the 91% of the total abundances of the B-Ex impacted station and the A-Ex. 30 impacted station.

4. Discussion

Despite the extraction depth and the considerable volume of sand removed, negligible change in sediment structure were observed at the relict sand extraction sites in the Northern Adriatic Sea. The deep depressions created by the extraction have gradually been levelled out by the winter storm activity and by the sedimentary regime (predominantly erosive) of the area that did not allow the accumulation of appreciable quantities of fine material in the depressions.

The variability in the composition and structure of the macrozoobenthic communities at the reference station, throughout the different surveys makes it difficult to analyze the recovery of the benthic community. A number of studies have highlighted the considerable intra- and inter-annual variability in composition and density of the community of sandy and muddy bottoms of the Northern Adriatic (Ambroggi et al., 1990; Bonvicini Pagliai et al., 1996; Bonvicini Pagliai et al., 1999; Prevedelli et al., 2001; Occhipinti Ambroggi et al., 2002; Simonini et al. 2004). It is likely that the fluctuations in density observed during the different surveys are linked to the biological cycles of different species and to the high seasonal variation in physical conditions in the Northern Adriatic Sea. For example, the reduction in terms of abundances and number of taxa that was observed during the A-Ex 12 survey can be related to the anomalous low temperature of the water (4°C in the bottom) that occurred in the northern Adriatic sea during the winter 2002 and caused the mortality of several benthic species (Gismondi e Rinaldi, 2002).

The temporal variability among the surveys was low and could not be attributed to the impact of the extraction activity. The effects of sand extraction on the macrozoobenthic communities were particularly intense, especially during and immediately after the extraction at station 3 and 4, and were mainly due to the removal of fauna from the seabed. The recolonisation at the impacted station began early by larval settlement (most polychaetes, molluscs, echinoderms and other sedentary organisms) and adult migration (vagile taxa, such as the dominant crustaceans *A. latreilli* and *A. diadema*). In other cases larval settlements were noted

to have been the most important source of new recruits, rather than the migration of adults from nearby (Boyd & Rees, 2003).

In contrast to the MDS and dissimilarity analyses, the high R value derived from the pair-wise ANOSIM test suggested that the community at the A-Ex impacted station were different from those found prior to the dredging operation, even 30 months after sand extraction. We suggest that the failure of the R statistic in detecting the recovery of the community could be the result of the low (spatial) variability among stations for each sampling survey relative to the natural seasonal and inter-annual variability of the benthic communities.

The % dissimilarity between Before-Extraction and After-Extraction impacted station decreased over time and, starting from 18 months after dredging, the community of impacted station grouped together with the reference station and B-Ex station on one side of the MDS plots. These findings suggested that starting from 24 months after dredging the composition and structure of the macrozoobenthic assemblages becomes similar to that observed prior to sand extraction. In particular, the difference in the community of the B-Ex and the A-Ex 30 impacted stations are due to 1) some rare species exclusive to the first or the last survey; and 2) differences in the abundances of common species, characteristic of the relict sands biocoenosis, that can be related to the natural temporal variability of the communities. The species in common between the two surveys constituted about 90% of the whole abundances at both B-Ex and the A-Ex 30 impacted stations. Therefore it could be assumed that the communities of the dredged area returned to the original condition 30 months after sand extraction.

This pattern of recolonisation-recovery fits well with the commonly encountered scenario where the substratum merely remains unchanged after marine aggregate excavation (Boyd et al., 2003, 2005; ICES, 2000). In most of these cases, the extraction area exhibited a strong hydrodynamic activity, which could obstruct the deposition of fine sediment, or the depth of the extraction was low, so the change in sediment structure of the dredged areas was limited over time and the complete recovery of the communities required a period of between 2 and 4 years.

In order to contrast coastal erosion along the Mediterranean coasts, the local governments and the environmental protection agencies of several regions and countries are planning new nourishment operations exploiting the relict sands (see INTEREG IIIC-BEACHMED project at <http://www.beachmed.it>). This monitoring programme has provided the opportunity to derive useful information for the evaluation of the consequences of relict sand extraction over sufficiently long time scales, from impact to recovery. Firstly, in order to separate the effects of sand extraction from the effects of the high inter- and intra-annual variability, the monitoring survey must be repeated a number of times both before and after dredging (Underwood, 1994). And secondly, to evaluate the outcome of the recolonisation-recovery processes the post-dredging survey might be performed 2-3 years after sand extraction

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Table 1. Results of two-way ANOSIM for the effects of sand extraction at different surveys, performed on untransformed and 4th root transformed macrofaunal abundance data.

Transformation Statistics Factors	none		4 th	
	R	P	R	P
Extraction	0.532	<0.001	0.635	<0.001
Survey	0.791	<0.001	0.732	<0.001

Table 2. Pair-wise test (One-way ANOSIM, untransformed data) and % average dissimilarity between the impacted stations vs reference stations for each survey, the impacted station from the different surveys vs the pre-dredging station (B-Ex Imp) and the reference stations of the different surveys. Significant *R* values ($P < 0.05$) are reported in bold.

Groups	<i>R</i>	% dissimilarity	Groups	<i>R</i>	% dissimilarity
A-Ex 30: Imp. vs Ref.	0.075	33.5	A-Ex 30 vs A-Ex 24	0.841	41.3
A-Ex 24: Imp. vs Ref.	0.460	40.6	A-Ex 30 vs A-Ex 18	0.21	33.0
A-Ex 18: Imp. vs Ref.	0.508	35.8	A-Ex 30 vs A-Ex 12	1.000	78.1
A-Ex 12: Imp. vs Ref.	0.611	57.2	A-Ex 30 vs A-Ex 6	0.760	51.0
A-Ex 6: Imp. vs Ref.	0.742	57.7	A-Ex 30 vs A-Ex 1	1.000	61.0
A-Ex 1: Imp. vs Ref.	0.948	83.0	A-Ex 30 vs Ex	1.000	59.1
Ex: Imp. vs Ref.	0.861	84.5	A-Ex 30 vs B-Ex	0.995	56.3
B-Ex: Imp. vs Ref.	-0.136	40.0	A-Ex 24 vs A-Ex 18	0.653	38.5
			A-Ex 24 vs A-Ex 12	1.000	79.4
			A-Ex 24 vs A-Ex 6	0.832	55.5
B-Ex Imp. vs Ex Imp.	0.444	89.2	A-Ex 24 vs A-Ex 1	1.000	61.7
B-Ex Imp. vs Ex 1 Imp.	1.000	91.1	A-Ex 24 vs Ex	1.000	60.3
B-Ex Imp. vs Ex 6 Imp.	0.850	65.8	A-Ex 24 vs B-Ex	1.000	57.8
B-Ex Imp. vs Ex 12 Imp.	1.000	83.2	A-Ex 18 vs A-Ex 12	1.000	78.0
B-Ex Imp. vs Ex 18 Imp.	1.000	51.9	A-Ex 18 vs A-Ex 6	0.715	46.2
B-Ex Imp. vs Ex 24 Imp.	1.000	58.9	A-Ex 18 vs A-Ex 1	1.000	59.0
B-Ex Imp. vs Ex 30 Imp.	1.000	54.3	A-Ex 18 vs Ex	1.000	56.7
			A-Ex 18 vs B-Ex	0.998	53.0
			A-Ex 12 vs A-Ex 6	0.949	72.1
			A-Ex 12 vs A-Ex 1	0.922	64.3
			A-Ex 12 vs Ex	0.962	67.2
			A-Ex 12 vs B-Ex	0.988	75.8
			A-Ex 6 vs A-Ex 1	0.741	50.7
			A-Ex 6 vs Ex	0.771	53.1
			A-Ex 6 vs B-Ex	0.631	48.6
			A-Ex 1 vs Ex	0.896	49.3
			A-Ex 1 vs B-Ex	0.963	53.8
			Ex vs B-Ex	0.954	53.1

Table 3. Pair-wise test (One-way ANOSIM, 4th root transformed data) and % average dissimilarity between the impacted stations vs reference ones of each survey, the impacted station from the different surveys vs the pre-dredging station (B-Ex Imp) and the reference stations of the different surveys. Significant *R* values ($P < 0.05$) are reported in bold.

Groups	<i>R</i>	% dissimilarity	Groups	<i>R</i>	% dissimilarity
A-Ex 30: Imp, vs Ref,	0,046	22.7	A-Ex 30 vs A-Ex 24	1.000	29.8
A-Ex 24: Imp, vs Ref,	0,698	27.7	A-Ex 30 vs A-Ex 18	0.697	25.0
A-Ex 18: Imp, vs Ref,	0,690	27.8	A-Ex 30 vs A-Ex 12	0.857	57.1
A-Ex 12: Imp, vs Ref,	0,683	56.1	A-Ex 30 vs A-Ex 6	0.890	37.6
A-Ex 6: Imp, vs Ref,	0,651	37.9	A-Ex 30 vs A-Ex 1	1.000	41.7
A-Ex 1: Imp, vs Ref,	0,917	68.2	A-Ex 30 vs Ex	1.000	39.2
Ex: Imp, vs Ref,	0,849	72.8	A-Ex 30 vs B-Ex	0.996	39.0
B-Ex: Imp, vs Ref,	-0,062	32.2	A-Ex 24 vs A-Ex 18	0.974	29.7
			A-Ex 24 vs A-Ex 12	0.992	59.0
			A-Ex 24 vs A-Ex 6	0.983	41.5
B-Ex Imp. vs Ex Imp.	0.444	74.1	A-Ex 24 vs A-Ex 1	1.000	42.7
B-Ex Imp. vs Ex 1 Imp.	0.815	74.3	A-Ex 24 vs Ex	1.000	41.0
B-Ex Imp. vs Ex 6 Imp.	0.815	45.5	A-Ex 24 vs B-Ex	1.000	41.6
B-Ex Imp. vs Ex 12 Imp.	0.963	65.6	A-Ex 18 vs A-Ex 12	0.863	57.0
B-Ex Imp. vs Ex 18 Imp.	1.000	39.1	A-Ex 18 vs A-Ex 6	0.688	34.7
B-Ex Imp. vs Ex 24 Imp.	1.000	40.5	A-Ex 18 vs A-Ex 1	1.000	41.4
B-Ex Imp. vs Ex 30 Imp.	1.000	40.3	A-Ex 18 vs Ex	1.000	38.2
			A-Ex 18 vs B-Ex	0.992	37.8
			A-Ex 12 vs A-Ex 6	0.734	52.6
			A-Ex 12 vs A-Ex 1	0.688	50.8
			A-Ex 12 vs Ex	0.715	51.5
			A-Ex 12 vs B-Ex	0.729	53.2
			A-Ex 6 vs A-Ex 1	0.793	35.1
			A-Ex 6 vs Ex	0.741	34.4
			A-Ex 6 vs B-Ex	0.642	33.8
			A-Ex 1 vs Ex	0.773	32.9
			A-Ex 1 vs B-Ex	0.821	34.3
			Ex vs B-Ex	0.704	32.2

Table 4. Number of taxa, density and Shannon diversity index calculated at each station of the different survey.

		Number of taxa							
Survey		B-Ex	Ex	A-Ex 1	A-Ex 6	A-Ex 12	A-Ex 18	A-Ex 24	A-Ex 30
Extraction	Station								
impact	2	62	42	28	60	31	72	91	73
	3	65	5	10	42	22	75	80	60
	4	71	6	14	31	17	73	100	78
reference	1	68	81	60	70	54	77	90	77
	5	69	54	68	59	45	83	95	75
	6	72	64	73	60	31	82	103	72
	7	54	69	55	52	32	76	97	78
	8	58	72	56	60	48	68	91	79
	9	72	64	62	63	22	79	92	75
	10	66	67	65	70	35	80	90	75
		Density [ind m ⁻²]							
Survey		B-Ex	Ex	A-Ex 1	A-Ex 6	A-Ex 12	A-Ex 18	A-Ex 24	A-Ex 30
Extraction	Station								
impact	2	1506	358	189	858	233	1878	2450	2481
	3	1561	14	42	483	219	1856	2911	2492
	4	2406	19	67	408	175	2236	2897	2308
reference	1	1492	1389	881	1864	533	1944	3508	1903
	5	1442	767	942	1369	550	2533	2858	2919
	6	1811	861	939	1492	214	2083	2800	1906
	7	2058	1156	661	550	286	2053	2392	1797
	8	1556	1161	703	1497	403	2333	2408	2678
	9	1817	1250	986	1600	258	2617	2581	2036
	10	1768	869	1092	1786	342	2106	2664	1856
		Shannon diversity H'							
Survey		B-Ex	Ex	A-Ex 1	A-Ex 6	A-Ex 12	A-Ex 18	A-Ex 24	A-Ex 30
Extraction	Station								
impact	2	3.35	3.31	2.95	3.45	2.76	2.94	3.64	3.36
	3	3.25	1.61	2.15	3.19	2.35	3.32	2.48	3.15
	4	3.19	1.75	2.40	3.11	2.16	2.93	2.87	3.25
reference	1	3.32	3.75	3.42	3.44	3.28	3.52	2.99	3.61
	5	3.41	3.56	3.69	3.44	2.81	3.28	3.17	3.35
	6	3.22	3.71	3.81	3.30	2.99	3.46	3.36	3.41
	7	3.11	3.78	3.53	3.46	3.00	3.62	3.69	3.65
	8	3.35	3.66	3.57	3.28	3.43	3.39	3.63	3.47
	9	3.38	3.54	3.65	3.42	2.54	3.46	3.74	3.57
	10	3.28	3.65	3.55	3.62	3.04	3.55	3.60	3.56

Figure legend

Fig. 1. Location of the sand extraction area with a map of the sampling stations and the sailing tracks of the dredger. Grey area = sand extraction area; grey points = stations in extraction area; white points = reference stations. The effects on the bottom profile was examined by side-scan sonar along the A-B axis.

Fig. 2. Grain size, and % total organic carbon (TOC; mean $\pm$ 2 S.D.) at impacted and reference stations at each survey.

Fig. 3. Reconstruction of the bottom profile immediately after (A Ex 1 survey) and 24 months (A Ex 24 survey) after sand extraction (based on multi-beam data).

Fig. 4. 2D-nMDS ordination of: a) impacted (Imp) and reference (Ref) station at all surveys, untransformed abundance data; b) impacted and reference station at all surveys, 4th root transformed abundance data; c) only impacted station at all surveys, untransformed abundance data; d) only impacted station at all surveys, 4th root transformed abundance data; e) only reference station at all surveys, untransformed abundance data; f) only reference station at all surveys, 4th root transformed abundance data. Abbreviations: B Ex – Before extraction; A Ex – After extraction followed by the time in months (1, 6, 12, 18, 24 and 30 months).

Fig. 5. Number of taxa, total abundance and Shannon-Wiener diversity index (mean $\pm$ 2 S.D.) at impacted and reference stations of each survey.

FIGURE 1 Simonini et al.

Fig. 2 Simonini et al.

Fig. 3 Simonini et al.

Fig. 4 Simonini et al.

FIGURE 5 Simonini et al.

Appendix 1. List of the taxa found during the eight surveys. Class (Cl.): A= amphipods; An= anthozoans; Ap= aplacophorans; As=ascidians; B= bivalves; C= cumaceans; D= decapods; De= demosponges; E= Echinoids; Ec= Echiurids; En= enteropneust; G= gastropods; H= hydrozoans; Ho= holothurians; I= isopods; N=nemerteans; O=ostracods; Op=ophiurans; P= Polychaetes; Ph= phoronids; Po=polyplacophorans; S=stomatopods; Si=sipunculids; ST=sea stars; Sy=sylicosponges; T=tanaidaceans.

Taxon	Cl.	Taxon	Cl.	Taxon	Cl.
<i>Amage adspersa</i>	P	<i>Prionospio cirrifera</i>	P	<i>Holothuria forskali</i>	Ho
<i>Ampharete acutifrons</i>	P	<i>Prionospio malmgreni</i>	P	<i>Labidoplax digitata</i>	Ho
<i>Aonides oxycephala</i>	P	<i>Prionospio steenstrupi</i>	P	<i>Leptosynapta inhaerens</i>	Ho
<i>Aphelochaeta marioni</i>	P	<i>Protodervillea atlantica</i>	P	<i>Ophiorthrix fragilis</i>	Op
<i>Aphelochaeta multibranchiis</i>	P	<i>Sabella pavonina</i>	P	<i>Ophiura albida</i>	Op
<i>Aphrodita aculeata</i>	P	<i>Sabellaria spinulosa</i>	P	<i>Phyllophorus urna</i>	Ho
<i>Aponuphis brementi</i>	P	<i>Sabellides octocirrata</i>	P	<i>Schizaster canaliferus</i>	E
<i>Aricidea claudiae</i>	P	<i>Scalibregma inflatum</i>	P	<i>Trachythone elongata</i>	Ho
<i>Armandia cirrhosa</i>	P	<i>Scolecopsis cantabra</i>	P	<i>Maxmuelleria gigas</i>	Ec
<i>Axionice maculata</i>	P	<i>Sigambra tentaculata</i>	P	<i>Glossobalanus minutus</i>	En
<i>Branchioma lucullanum</i>	P	<i>Sphaerosyllis bulbosa</i>	P	<i>Abra prismatica</i>	B
<i>Dasybranchus caducus</i>	P	<i>Spio filicornis</i>	P	<i>Achantocardia tuberculata</i>	B
<i>Ditrupa arietina</i>	P	<i>Spio multioculata</i>	P	<i>Anomia ephippium</i>	B
<i>Eteone flava</i>	P	<i>Spiophanes kroyeri kroyeri</i>	P	<i>Aporrhais pespelecani</i>	G
<i>Euchone</i> sp.	P	<i>Sthenolepis yhleni</i>	P	<i>Atrina pectinata</i>	B
<i>Euclymene lumbricoides</i>	P	<i>Syllis amica</i>	P	<i>Azorus chamasolen</i>	B
<i>Euclymene oerstedii</i>	P	<i>Syllis cornuta</i>	P	<i>Calyptaea chinensis</i>	G
<i>Eunice pennata</i>	P	<i>Tachytrypae jeffreysii</i>	P	<i>Chlamys varia</i>	B
<i>Eunice vittata</i>	P	<i>Terebellides stroemi</i>	P	<i>Clausinella brongniartii</i>	B
<i>Filogranula annulata</i>	P	<i>Thalenessa dendrolepis</i>	P	<i>Corbula gibba</i>	B
<i>Glycera alba</i>	P	<i>Acheus gracilis</i>	D	<i>Cuspidaria rostrata</i>	B
<i>Glycera capitata</i>	P	<i>Ampelisca diadema</i>	A	<i>Cylichna cylindracea</i>	G
<i>Glycera tridactyla</i>	P	<i>Apseudes latreillii</i>	T	<i>Dosinia lupinus</i>	B
<i>Goniada maculata</i>	P	<i>Atelecycclus rotundatus</i>	D	<i>Ensis ensis</i>	B
<i>Harmothoe imbricata</i>	P	<i>Bodotria scorpioides</i>	C	<i>Fusinus rostratus</i>	G
<i>Harmothoe johnstoni</i>	P	<i>Callianassa subterranea</i>	D	<i>Glycymeris insubrica</i>	B
<i>Harmothoe lunulata</i>	P	<i>Cirolana borealis</i>	I	<i>Hiatella rugosa</i>	B
<i>Hydroides norvegicus</i>	P	<i>Corystes cassivelaunus</i>	D	<i>Hyala vitrea</i>	G
<i>Jasmineira elegans</i>	P	<i>Diastylis rugosa</i>	C	<i>Hyalopecten similis</i>	B
<i>Kefersteinia cirrata</i>	P	<i>Ebalia deshayesi</i>	D	<i>Lentidium mediterraneum</i>	B
<i>Lanice conchylega</i>	P	<i>Ethusa mascarone</i>	D	<i>Lepidopleurus cancellatus</i>	Po
<i>Laonice cirrata</i>	P	<i>Galathea intermedia</i>	D	<i>Loripes lacteus</i>	B
<i>Lepidonotus clava</i>	P	<i>Gammaridae</i> n.c.	A	<i>Lucinella divaricata</i>	B
<i>Lumbrineris latreilli</i>	P	<i>Gnathia phallonajopsis</i>	I	<i>Mactra stultorum</i>	B
<i>Lumbrineris tetraura</i>	P	<i>Inachus dorsettensis</i>	D	<i>Melanella polita</i>	G
<i>Magelona papillicornis</i>	P	<i>Iphinoe tenella</i>	C	<i>Modiolarca subpicta</i>	B
<i>Malacoceros girardi</i>	P	<i>Leptognathia brevimanu</i>	T	<i>Modiolula phaseolina</i>	B
<i>Maldane sarsi</i>	P	<i>Liocarcinus arcuatus</i>	D	<i>Myrtea spinifera</i>	B
<i>Marphysa bellii</i>	P	<i>Nannastacidae</i> n.d.	C	<i>Mysella bidentata</i>	B
<i>Melinna palmata</i>	P	<i>Ostracoda</i> n.c.	O	<i>Nucula nitidosa</i>	B
<i>Micronephthys sphaerocirrata</i>	P	<i>Pagurus anachoretus</i>	D	<i>Nuculana pella</i>	B
<i>Monticellina dorsobranchialis</i>	P	<i>Philocheilus sculptus</i>	D	<i>Paphia aurea</i>	B
<i>Neanthes succinea</i>	P	<i>Phtisiça marina</i>	A	<i>Parvicardium exiguum</i>	B
<i>Nematonereis unicornis</i>	P	<i>Pilumnus hirtellus</i>	D	<i>Phaxas adriaticus</i>	B
<i>Nephtys hombergii</i>	P	<i>Platysquilla eusebia</i>	S	<i>Philine scabra</i>	G
<i>Nothria conchylega</i>	P	<i>Processa canaliculata</i>	D	<i>Pitar rudis</i>	B
<i>Notomastus latericeus</i>	P	<i>Pycnogonida</i> n.d.	Py	<i>Polinices nitida</i>	G
<i>Ophiodromus agilis</i>	P	<i>Rissoides desmaresti</i>	D	<i>Prochaetoderma raduliferum</i>	Ap
<i>Orbinia cuvieri</i>	P	<i>Sicyonia carinata</i>	D	<i>Psammobia fervensis</i>	B
<i>Owenia fusiformis</i>	P	<i>Solenocera mebranacea</i>	D	<i>Solemya togata</i>	B
<i>Paraonides neapolitana</i>	P	<i>Upogebia pusilla</i>	D	<i>Tellimya ferruginosa</i>	B
<i>Pectinaria auricoma</i>	P	<i>Ascidia involuta</i>	As	<i>Tellina distorta</i>	B
<i>Pectinaria koreni</i>	P	<i>Alcyonium</i> sp.	An	<i>Thracia papyracea</i>	B
<i>Pherusa plumosa</i>	P	<i>Anthozoa</i> n.c.	An	<i>Thyasira flexuosa</i>	B
<i>Phyllodoce macrophthalma</i>	P	<i>Condylactis aurantiaca</i>	An	<i>Nemertea</i> n.c. sp1	N
<i>Pilargis verrucosa</i>	P	<i>Hydrozoa</i> n.d.	H	<i>Phoronis muelleri</i>	Ph
<i>Pista cristata</i>	P	<i>Amphipholis squamata</i>	Op	<i>Sycon</i> sp.	Sy
<i>Poecilochaetus fauchaldi</i>	P	<i>Amphiura chiajei</i>	Op	<i>Timeidae</i> n.d.	D
<i>Polydora flava</i>	P	<i>Amphiura filiformis</i>	Op	<i>Aspidosiphon muelleri</i>	Si
<i>Pomatoceros lamarckii</i>	P	<i>Astropecten irregularis</i>	St	<i>Phascolion strombi</i>	Si
<i>Pomatoceros triqueter</i>	P	<i>Echinocyamus pusillus</i>	E	<i>Sipunculus nudus</i>	Si

Appendix 2. Average abundance (ind m⁻²), % contribution and cumulative contribution of taxa which most contribute to within-group similarity and between-group dissimilarity between impacted (Imp) stations at before extraction (B Ex) and after 30 month extraction (A Ex 30) surveys (SIMPER).

SIMPER (cut-off 50%) no trasformation										
Group		B Ex Imp		Groups		B Ex Imp vs A Ex 30 Imp				
Average similarity		71.32		Average dissimilarity		54.28				
Taxa		Av.Abund	Cont.%	Cum.%	Group		B Ex Imp	A Ex 30 Imp		
<i>Spiophanes kroyeri kroyeri</i>		261.1	17.2	17.2	Taxa		Av.Abund	Av.Abund	Cont.%	Cum.%
<i>Apseudes latreillii</i>		207.4	13.1	30.3	<i>Apseudes latreillii</i>		207.4	614.8	18.0	18.0
<i>Gammaridae n.c.</i>		188.9	12.8	43.1	<i>Spiophanes kroyeri kroyeri</i>		261.1	7.4	11.0	29.0
<i>Ariceida claudiae</i>		238.0	9.8	52.9	<i>Ariceida claudiae</i>		238.0	129.6	6.3	35.3
					<i>Gammaridae n.c.</i>		188.9	75.9	4.9	40.1
					<i>Aphelochaeta marioni</i>		43.5	119.4	4.1	44.3
					<i>Magelona papillicornis</i>		3.7	85.2	3.6	47.9
					<i>Jasmineira elegans</i>		38.9	106.5	3.0	50.9
Group		A Ex 30 Imp								
Average similarity		69.53								
Taxa		Av.Abund	Cont.%	Cum.%						
<i>Apseudes latreillii</i>		614.8	34.8	34.8						
<i>Jasmineira elegans</i>		106.5	5.3	40.1						
<i>Paraonides neapolitana</i>		100.9	4.6	44.7						
<i>Magelona papillicornis</i>		85.2	4.1	48.7						
<i>Ariceida claudiae</i>		129.6	4.0	52.7						
SIMPER (cut-off 50%) 4rt root trasformation										
Group		B Ex Imp		Groups		B Ex Imp vs A Ex 30 Imp				
Average similarity		75.45		Average dissimilarity		40.27				
Taxa		Av.Abund	Cont.%	Cum.%	Group		B Ex Imp	A Ex 30 Imp		
<i>Spiophanes kroyeri kroyeri</i>		261.1	4.1	4.1	Taxa		Av.Abund	Av.Abund	Cont.%	Cum.%
<i>Apseudes latreillii</i>		207.4	3.8	7.8	<i>Magelona papillicornis</i>		3.7	85.2	2.3	2.3
<i>Ariceida claudiae</i>		238.0	3.5	11.4	<i>Spiophanes kroyeri kroyeri</i>		261.1	7.4	2.2	4.5
<i>Gammaridae n.c.</i>		188.9	3.5	14.9	<i>Prionospio malmgreni</i>		0.0	32.4	2.0	6.5
<i>Aponuphis brementi</i>		77.8	3.0	17.9	<i>Callianassa subterranea</i>		0.0	24.1	2.0	8.5
<i>Paraonides neapolitana</i>		55.6	2.8	20.7	<i>Malacoceros girardi</i>		15.8	0.0	1.8	10.3
<i>Ampelisca diadema</i>		58.3	2.7	23.4	<i>Eunice pennata</i>		16.7	0.0	1.8	12.1
<i>Lumbrineris latreilli</i>		49.1	2.7	26.1	<i>Ampharete acutifrons</i>		0.0	15.8	1.8	13.9
<i>Aphelochaeta multibranchiis</i>		54.6	2.7	28.8	<i>Syllis cornuta</i>		0.0	12.0	1.7	15.6
<i>Jasmineira elegans</i>		38.9	2.4	31.2	<i>Pectinaria koreni</i>		0.0	10.2	1.6	17.2
<i>Aphelochaeta marioni</i>		43.5	2.3	33.5	<i>Eunice vittata</i>		0.0	9.3	1.5	18.8
<i>Pista cristata</i>		19.4	2.2	35.6	<i>Solemya togata</i>		7.4	0.0	1.5	20.3
<i>Eteone flava</i>		23.1	2.2	37.8	<i>Hydroides norvegicus</i>		0.0	7.4	1.5	21.8
<i>Melinna palmata</i>		24.1	2.2	39.9	<i>Sabella pavonina</i>		0.0	6.5	1.5	23.2
<i>Marphysa bellii</i>		25.0	2.1	42.0	<i>Nematonereis unicornis</i>		17.6	0.9	1.5	24.7
<i>Notomastus latericeus</i>		15.8	2.1	44.1	<i>Marphysa bellii</i>		25.0	5.6	1.5	26.1
<i>Sigambra tentaculata</i>		18.5	2.0	46.1	<i>Liocarcinus arcuatus</i>		0.0	5.6	1.4	27.5
<i>Poecilochaetus fauchaldi</i>		22.2	2.0	48.1	<i>Ophiura albida</i>		6.5	50.9	1.4	28.9
<i>Malacoceros girardi</i>		15.8	1.9	50.0	<i>Lumbrineris tetraura</i>		17.6	2.8	1.4	30.3
Group		A Ex 30 Imp								
Average similarity		75.17								
Taxa		Av.Abund	Cont.%	Cum.%						
<i>Apseudes latreillii</i>		614.8	4.6	4.6						
<i>Jasmineira elegans</i>		106.5	2.9	7.5						
<i>Paraonides neapolitana</i>		100.9	2.8	10.3						
<i>Magelona papillicornis</i>		85.2	2.7	13.0						
<i>Ariceida claudiae</i>		129.6	2.7	15.7						
<i>Ampelisca diadema</i>		83.3	2.6	18.3						
<i>Euchone sp.</i>		79.6	2.6	20.9						
<i>Aphelochaeta multibranchiis</i>		79.6	2.5	23.3						
<i>Poecilochaetus fauchaldi</i>		48.1	2.4	25.7						
<i>Ophiura albida</i>		50.9	2.3	28.0						
<i>Aphelochaeta marioni</i>		119.4	2.3	30.3						
<i>Gammaridae n.c.</i>		75.9	2.5	32.7						
<i>Sigambra tentaculata</i>		39.8	2.2	34.9						
<i>Nemertea n.c.</i>		34.3	2.1	37.0						
<i>Lumbrineris latreilli</i>		32.4	2.1	39.1						
<i>Goniada maculata</i>		31.5	2.1	41.2						
<i>Cirolana borealis</i>		25.9	2.0	43.1						
<i>Syllis amica</i>		50.9	1.9	45.1						
<i>Phtisica marina</i>		22.2	1.9	47.0						
<i>Amage adspersa</i>		24.1	1.9	48.9						
<i>Leptognathia brevimanu</i>		15.8	1.8	50.6						
Group		A Ex 30 Imp		Groups		B Ex Imp vs A Ex 30 Imp				
Average similarity		75.17		Average dissimilarity		40.27				
Taxa		Av.Abund	Cont.%	Cum.%	Group		B Ex Imp	A Ex 30 Imp		
<i>Apseudes latreillii</i>		614.8	4.6	4.6	Taxa		Av.Abund	Av.Abund	Cont.%	Cum.%
<i>Jasmineira elegans</i>		106.5	2.9	7.5	<i>Magelona papillicornis</i>		3.7	85.2	2.3	2.3
<i>Paraonides neapolitana</i>		100.9	2.8	10.3	<i>Spiophanes kroyeri kroyeri</i>		261.1	7.4	2.2	4.5
<i>Magelona papillicornis</i>		85.2	2.7	13.0	<i>Prionospio malmgreni</i>		0.0	32.4	2.0	6.5
<i>Ariceida claudiae</i>		129.6	2.7	15.7	<i>Callianassa subterranea</i>		0.0	24.1	2.0	8.5
<i>Ampelisca diadema</i>		83.3	2.6	18.3	<i>Malacoceros girardi</i>		15.8	0.0	1.8	10.3
<i>Euchone sp.</i>		79.6	2.6	20.9	<i>Eunice pennata</i>		16.7	0.0	1.8	12.1
<i>Aphelochaeta multibranchiis</i>		79.6	2.5	23.3	<i>Ampharete acutifrons</i>		0.0	15.8	1.8	13.9
<i>Poecilochaetus fauchaldi</i>		48.1	2.4	25.7	<i>Syllis cornuta</i>		0.0	12.0	1.7	15.6
<i>Ophiura albida</i>		50.9	2.3	28.0	<i>Pectinaria koreni</i>		0.0	10.2	1.6	17.2
<i>Aphelochaeta marioni</i>		119.4	2.3	30.3	<i>Eunice vittata</i>		0.0	9.3	1.5	18.8
<i>Gammaridae n.c.</i>		75.9	2.5	32.7	<i>Solemya togata</i>		7.4	0.0	1.5	20.3
<i>Sigambra tentaculata</i>		39.8	2.2	34.9	<i>Hydroides norvegicus</i>		0.0	7.4	1.5	21.8
<i>Nemertea n.c.</i>		34.3	2.1	37.0	<i>Sabella pavonina</i>		0.0	6.5	1.5	23.2
<i>Lumbrineris latreilli</i>		32.4	2.1	39.1	<i>Nematonereis unicornis</i>		17.6	0.9	1.5	24.7
<i>Goniada maculata</i>		31.5	2.1	41.2	<i>Marphysa bellii</i>		25.0	5.6	1.5	26.1
<i>Cirolana borealis</i>		25.9	2.0	43.1	<i>Liocarcinus arcuatus</i>		0.0	5.6	1.4	27.5
<i>Syllis amica</i>		50.9	1.9	45.1	<i>Ophiura albida</i>		6.5	50.9	1.4	28.9
<i>Phtisica marina</i>		22.2	1.9	47.0	<i>Lumbrineris tetraura</i>		17.6	2.8	1.4	30.3
<i>Amage adspersa</i>		24.1	1.9	48.9	<i>Nemertea n.c.</i>		4.6	34.3	1.3	31.6
<i>Leptognathia brevimanu</i>		15.8	1.8	50.6	<i>Pagurus sp.</i>		0.9	10.2	1.3	32.9
					<i>Nephtys hombergii</i>		11.1	6.5	1.2	34.1
					<i>Glycera capitata</i>		8.3	0.9	1.2	35.2
					<i>Aponuphis brementi</i>		77.8	55.6	1.2	36.4
					<i>Phoronis muelleri</i>		1.9	10.2	1.2	37.6
					<i>Nannastacidae n.d.</i>		0.9	8.3	1.2	38.8
					<i>Monticellina dorsobranchialis</i>		32.4	8.3	1.2	39.9
					<i>Schizaster canaliferus</i>		0.0	9.3	1.1	41.1
					<i>Apseudes latreillii</i>		207.4	614.8	1.1	42.2
					<i>Hydrozoa n.d.</i>		0.9	7.4	1.1	43.3
					<i>Pomatoceros triqueter</i>		0.0	6.5	1.1	44.4
					<i>Amage adspersa</i>		4.6	24.1	1.1	45.5
					<i>Phtisica marina</i>		3.7	22.2	1.1	46.6
					<i>Phaxas adriaticus</i>		0.9	8.3	1.1	47.7
					<i>Diastylis rugosa</i>		0.0	5.6	1.1	48.8
					<i>Micronephthys sphaerocirrata</i>		0.0	4.6	1.0	49.8
					<i>Orbinia cuvieri</i>		0.0	5.6	1.0	50.8