We wish to thank Anonymous Referee #1 for his thorough and useful comments. We have now revised our manuscript in accordance with the comments raised by the referee. We believe that this revision has substantially improved the quality of the manuscript. Please find below how we have addressed each comment, point by point.

**Comment 1**  
*Page 3, line 22: I understand that a deposit of fine, cohesive sediment will decrease the supply of dissolved oxygen to the deposit-underlying sediment and so decrease the decomposition of organic matter in this sediment with oxygen as electron acceptor. If so, the contribution of anaerobic pathways to the overall decomposition will increase and the upwards diffusing reduced soluble end-products of this decomposition will likely be oxidised with oxygen at the oxic–anoxic boundary somewhere inside the deposit or in the deposit overlying seawater. That is, the re-oxidation of reduced substances (line 24) is not inhibited but simply relocated. Of course, this would not apply for reduced solid phases, but this perhaps needs to be clarified.*

Indeed, this is correct and we have therefore rephrased this part to clarify better how a physical barrier alters the contribution of anaerobic pathways:

Page 3, line 22: “Firstly, the formation of a physical barrier increases the contribution of anaerobic pathways to the overall decomposition and relocates the re-oxidation of reduced solutes upwards (Colden and Lipcius 2015; Hohaia et al. 2014). Under these circumstances, reduced solid phases would only oxidise when sediment reworking or irrigation of large burrows by macrofauna brings them to the oxic layer.”

**Comment 2**  
*Page 4, line 5: In my book, bioturbation includes the displacement of particles and the irrigation of burrows. In line 5, it reads ‘bioturbation or bio-irrigation’, so I assume that the authors do not consider burrow irrigation as a form of bioturbation. Perhaps this needs to be clarified as well.*

Kristensen et al. (2012) proposed to use bioturbation as an umbrella term, incorporating both burrow ventilation and particle reworking. Indeed, burrow ventilation is a mechanism evolved by infauna to enable a constant supply of fresh nutrients and oxygen by pumping overlying water into their burrows, and as a transport process clearly associated with bioturbation. However, since we aimed at disentangling the mechanisms of deposition-induced alteration of SCOC (burrow ventilation, macrofauna respiration or particle mixing into oxic layers) we preferred to distinguish between bioturbation (i.e. particle reworking) and bio-irrigation (i.e. burrow ventilation). We incorporated this rationale in the manuscript:

Page 3, line 27: “Though both processes are interrelated and sometimes grouped under the umbrella term ‘bioturbation’ (Kristensen et al., 2012), we opted to use them as separate concepts, in order to clearly distinguish between particle reworking and solute transfer. Bioturbation and bio-irrigation can be significantly altered under...”

**Comment 3**  
*Page 4, line 27. The authors state that their control (T0) did not receive a layer of pre-treated sediment. In line 30, however, they explain that the control did receive a 0.5 cm frozen mud cake, which consisted of pre-treated sediment and luminophores.*

Indeed, this is correct.
How did this layer affect the mud–seawater solute exchange and the behaviour of macroinfauna? I feel the authors should discuss this.

Our objective to disentangle the different mechanisms of altered oxygen consumption necessitated the application of a luminophore-spiked mud cake on all treatments (including the control sediments). Without such thin cake on the control the importance of particle mixing and disturbance of the sediment matrix at the sediment-water interface for deposition-altered functioning would have been impossible to investigate. Moreover, though luminophores are in essence inert particles, the absence of such a luminophore mud cake on the natural sediment in the control could potentially have introduced bias between treatments due to species specific responses to e.g. small modifications in physico-chemistry of the sediment matrix, hence creating an experimental artefact. The high survival and appearance of clear bioturbation signs at the sediment surface, already the day after application of the mud cake in the control (photos are included in Supplementary material Annex 2), indicate that the application of the thin deposit evoked fast migration to the sediment-water interface in the control. However, we do not believe that this thin deposition and subsequent fast disturbance related to benthos migration significantly altered functioning at the longer term, i.e. at the end of the experiment 14 days after addition of the mud cakes. This hypothesis is supported by the high survival but lower bioturbation and bio-irrigation in the control as compared to the T1 treatment. Collectively, this suggests a fast recovery of the sediment-water solute exchange following the deposition of the thin mud cake in the control. Indeed the measured oxygen penetration depth and SCOC in the control are comparable in magnitude to the diffusive and sediment community oxygen fluxes measured in the same habitat and season in previous studies (Van Colen et al. 2012; see manuscript for full reference). We have added this rationale in the revised manuscript:

Page 4, line 31: “… on top of the natural sediment surface. The addition of this mud cake ensured the quantification of particle mixing in these treatments and avoided potential bias between treatments due to species specific responses to the physico-chemical environment created by the mud cake. The addition of a luminophore mud cake on top of the sediment surface in the control treatment did not profoundly affect the natural oxygen fluxes or oxygen penetration depth. Our measured values were comparable in magnitude to those of previous studies in the same habitat and season (Van Colen et al. 2012; Annex 1), and clear bioturbation signs on the sediment surface soon after deposition indicate fast migration to the sediment-water interface (Annex 2).”

Comment 4 Page 5, line 3. The deposit was free of organic matter, so its oxygen demand must have been low increasing the penetration of oxygen into the layer. How do the authors know that this deposit ‘prohibited (passive) exchange of dissolved oxygen between the sampled community and the water column’? Did you measure the penetration of oxygen into the freshly deposited layers with microelectrodes and did you find the oxic-anoxic boundary somewhere inside the layer? If so, how did the four different deposits (0.5, 1, 2, 5 mm) perform in regard to this penetration?

The oxygen penetration depth varied from shallower in the control to deeper below the sediment-water interface in the more extreme deposits (that were largely depleted in organic matter as compared to the control). However, oxygen penetration
depth remained restricted to the deposited layer for all treatments. Thus, oxygen did not diffuse below the deposited layers into the natural community. The vertical profiles of oxygen penetration are submitted as supplementary material to the manuscript, to which we now refer in the text (Page 5, line 3; Page 9, line 32; See comment 3).

Comment 5  
*Page 5, line 33. Here, BMU is defined as ‘biological-mediated oxygen uptake’. I found this misleading because biological mediated oxygen consumption is also included in estimates of DOU, that is, the consumption by bacterial processes, micro- and meiofauna. I believe that this contribution to the overall sediment oxygen consumption should be termed ‘macrofauna mediated oxygen uptake’.*

We agree that the terminology we used was potentially confusing and have therefore followed the suggestion by this referee to change this term to ‘macrofauna-mediated oxygen uptake’ in the revised version of the manuscript.

Comment 6  
*Page 8. Please consider moving numbers in parentheses to a table; this would improve the readability of your text.*

We have accepted this comment. We now refer to Tables 2 and 4 in the revised manuscript which contain the results of the statistical test.

Comment 7  
*Page 8, line 37. ‘biotic-mediated oxygen consumption’. See comment above and please use terms consistently.*

This inconsistency apparently remained unnoticed by me and the co-authors, and we have now corrected this throughout the revised manuscript.

Comment 8  
*Page 9, lines 14–28. I recommend moving this section to the introduction, so the discussion starts with your results.*

We have adopted this comment.

Comment 9  
*Page 9, line 31. Please show the oxygen penetration data in the Results section.*

See also reply to comment 4. Oxygen penetration depths are now provided as supplementary material to the manuscript.

Comment 10  
*Page 23, line 6. ‘benthic-mediated oxygen uptake (BMU)’. See comment above and please use terms consistently.*

Complied with this comment; see also reply to comment 7.
We are thankful for the useful comments of Anonymous Referee #2. We have revised the figure and table captions accordingly:

The current table legend is more like discussion, and a mere descriptive of data is preferred. It would be more appropriate that the authors simply provide the objective data, and let the readers to judge whether your conclusion or interpretation are reasonable or not.

Comment 1 For example, in Table 1 (page 16) (1) Table legend. The top 3 species with the cumulative contribution (>50%) to the total dissimilarity between treatments. The SIMPER analysis, and the cumulative contribution can be described as footnote. (2) Treatment. Please explain what these treatment mean. For example. What does T0-1 refer to. Please simply state what these data represent. In addition, the cumulative contribution column is redundant and can be deleted, or the contribution column can be deleted because these two column are essentially the same.

We deleted the ‘Contribution’ column from the table and rewrote the caption as follows, according to the referee’s suggestions:

“The three species with highest cumulative contribution (> 50 %) to the total dissimilarity between treatments. The first column shows the treatments being compared (e.g. T0-1: a comparison between treatments T0 and T1).”

We added a footnote to the table:

“Results from a SIMPER analysis.”

Comment 2 As for Table 2 (page 17) (1) In table legend, Please do not start with “Results ...”. Please go straight forward what you want to present. For example, statistical factors (2) Is it necessary to show all these factors?

We agree that a mere representation of F- and p-values could be sufficient for a decent understanding of our results. All unnecessary columns (i.e. df, SS and MS) were therefore deleted in the revised manuscript. We suggest the following caption, based on the referee’s comment:

“Statistical factors from 2-factor blocked ANOVA tests with ‘Treatment’ (4 levels) and ‘Tank’ as factors. M1 till M4 stand for motility classes, as defined by Solan et al. (2004) (M1: living fixed in a tube, M2: sessile, but not fixed in a tube, M3: slow movement through the sediment, M4: free movement in a burrow system). Significant pair-wise differences between treatments are given in the table. All results for species and functional groups are given for densities.”

Comment 3 As for Table 3 (page 19). Please consider whether all these equations need to be shown in a table. Maybe the equations could be placed in the supplementary materials.

We opted to present the results of the linear regressions entirely in the main manuscript, as it provides a complete understanding of the strength and direction of the relationship between the response and predictor variables. Therefore, we would prefer to keep the results as they are currently presented in the table.

In accordance to Comment 2, we opted to rewrite the caption of this table:
“Linear regressions of sediment community oxygen consumption (SCOC) against sets of species (or functional group) densities, and ecosystem processes (bio-irrigation - Q - and bioturbation - D\text{bi}^V), and of bio-irrigation against the densities of species. Only significant models (P (slope) < 0.05) were considered. M2 and M3 are motility classes as defined by Solan et al. (2004) – M2: sessile, but not fixed in a tube, M3: slow movement through the sediment.”

Comment 4  
**Figure 1. please explain the x axis, i.e., what the treatment of 0, 1, 2 and 5 mean.**

We added the next sentence to the figure caption of this and the next two figures:

“The four treatments represent the thickness of the applied sediment layer (in cm).“
Functional trait responses to sediment deposition reduce macrofauna-mediated ecosystem functioning in an estuarine mudflat

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Abstract. Human activities, among which dredging and land use change in river basins, are altering estuarine ecosystems. These activities may result in changes in sedimentary processes, affecting biodiversity of sediment macrofauna. As macrofauna control sediment chemistry and fluxes of energy and matter between water column and sediment, changes in the structure of macrobenthic communities could affect the functioning of an entire ecosystem. We assessed the impact of sediment deposition on intertidal macrobenthic communities and on rates of an important ecosystem function, i.e. sediment community oxygen consumption (SCOC). An experiment was performed with undisturbed sediment samples from the Scheldt river estuary (SW Netherlands). The samples were subjected to four sedimentation regimes: one control and three with a deposited sediment layer of 1, 2 or 5 cm. Oxygen consumption was measured during incubation at ambient temperature. Luminophores applied at the surface, and a seawater-bromide mixture, served as tracers for bioturbation and bio-irrigation, respectively. After incubation, the macrofauna was extracted, identified and counted, and classified into functional groups based on motility and sediment reworking capacity. Total macrofaunal densities dropped already under the thinnest deposits. The most affected fauna were surficial and low-motile animals, occurring at high densities in the control. Their mortality resulted in a drop in SCOC, which decreased steadily with increasing deposit thickness, while bio-irrigation and bioturbation activity showed increases in the lower sediment deposition regimes, but decreases in the more extreme treatments. The initial increased activity likely counteracted the effects of the drop in low-motile, surficial fauna densities, resulting in a steady rather than sudden fall in oxygen consumption. We conclude that the functional identity in terms of motility and sediment reworking can be crucial in our understanding of the regulation of ecosystem functioning and the impact of habitat alterations such as sediment deposition.

Key words: biogeochemical cycling, bio-irrigation, bioturbation, ecosystem functioning, functional traits, macrobenthos, SCOC, sediment deposition
1 Introduction

It is widely accepted that biodiversity plays an important role in ecosystem functioning. A higher biodiversity can convey a higher resilience and a more efficient functioning of ecosystems in terms of, among others, nutrient cycling and primary productivity (Cardinale et al., 2012; Hooper et al., 2005). Since biodiversity-mediated ecosystem functioning depends on the functional identities of the species present in the community and their densities (Braeckman et al., 2010; Van Colen et al., 2013), functional community descriptors often predict functioning better than taxonomic diversity (Wong and Dowd, 2015). Functional traits, e.g. in terms of motility or sediment reworking rate, can be an indication for a species’ behaviour. By being able to rework more or less sediment, species can differentially influence biogeochemical cycling (Wrede et al., 2017). Furthermore, variations in population densities of individual species can influence the ecosystem functioning as well (e.g. Braeckman et al., 2010). Habitat changes that alter densities and/or induce behavioural change of specific functional groups of organisms, e.g. top predators or key players in biogeochemical cycling (Allen and Clarke, 2007; Villnäs et al., 2012), are therefore likely to change the functioning of ecosystems. Natural disturbances occur frequently in coastal and estuarine ecosystems, and recent intense anthropogenic activities often significantly reduce ecosystem resilience (Alestra and Schiel, 2015). An important example of such a human-induced change in coastal and estuarine habitats is sediment deposition. Natural sedimentation is caused by surface runoff from the catchment area or by tidal movements; the former can be intensified by land use change (Thrush et al., 2004). Furthermore, dredging and dumping activities also contribute to sediment deposition, either directly or by creating sediment plumes that subsequently settle down on the seabed (Van Lancker and Baeye, 2015). Such deposition events are expected to alter the productivity of coastal soft-sediment habitats via direct and indirect mechanisms that affect biogeochemical cycling. Firstly, the formation of a physical barrier increases the contribution of anaerobic pathways to the overall decomposition and relocates re-oxidation of reduced solutes deposition of fine sediments reduces aerobic mineralisation through the formation of a physical barrier at the sediment-water interface that inhibits re-oxidation of reduced substances in the sediment (Colden and Lipcius, 2015; Hohaia et al., 2014). Under these circumstances, reduced solid phases would only oxidise when sediment reworking or irrigation of large burrows by macrofauna brings them to the oxic layer. Macrofauna plays an important role in the biogeochemical cycling of soft sediments through sediment particle mixing (i.e. bioturbation) and the assisted transfer of solutes through the sediment (i.e. bio-irrigation) (Braeckmann et al., 2010, 2014; Van Colen et al., 2012; Thrush et al., 2006). Both processes are interrelated and sometimes grouped under the umbrella term ‘bioturbation’ (Kristensen et al., 2012), we opted to use them as separate concepts, in order to clearly distinguish between particle reworking and solute transfer. Bioturbation and bio-irrigation can be significantly altered under increased sediment deposition through changes in macrobenthic densities (Alves et al., 2017) or behaviour (Rodil et al., 2011). For example, sessile organisms that live attached to the substratum or in tubes, often have a limited capacity to escape burial, and suspension feeders risk clogging of their feeding apparatus (Ellis et al., 2002; Lohrer et al., 2004). Secondly, macrofauna activities can interfere with the deposition induced physical barrier at the sediment-water interface. Sediment deposition induced loss of macrofauna species density and change of behaviour therefore represents a second, more indirect pathway of how deposition events can alter ecosystem functioning.
Tidal flats are dynamic, sedimentary environments that naturally undergo processes of erosion and deposition. Per tidal cycle, different elevation changes have been observed, e.g. from decreases of 3.3 mm in the Yangtze estuary (China) to increases of 6 mm in the estuary of the Seine (France) (Deloffre et al., 2007; Shi et al., 2012). Our study was performed on a mudflat in the estuary of the river Scheldt (Belgium, the Netherlands), which is characterised by its meso- to macro-tidal regime and well-mixed water column. Sediment input from the river basin is relatively low and sand extraction and sea level rise lead to a net export of sediment from the estuary (De Vriend et al., 2011). Sediment accretion on the estuary’s tidal flats can amount to about 2 cm yr\(^{-1}\) (Weerman et al., 2011; Widdows et al., 2004), which suggests that natural sedimentation on the intertidal mudflats is unlikely to exceed even a few millimetres per tidal cycle. More extreme changes in the bed level of mudflats can however happen during storm events, either by erosion of the top centimetres of the sediment or by deposition of new sediment (Hu et al., 2015; Marion et al., 2009). Besides natural processes, anthropogenic factors influencing sedimentation are prominent in the estuary, among which dredging in the main channels to ensure access to the port of Antwerp, and dumping of the dredged material to retain sediment within the estuary, are the most important (Jeuken and Wang, 2010; Meire et al., 2005). Most of this dredged sediment is disposed of near shoals and tidal flats, and can as such affect the intertidal ecosystem (Bolam and Whomersley, 2005; De Vriend et al., 2011; Zheng, 2015).

The effects of sediment deposition on taxonomic diversity (Thrush et al., 2003), behaviour (Hohaia et al., 2014; Townsend et al., 2014), and ecosystem functioning (Larson and Sundbäck, 2012; Montserrat et al., 2011) have recently received considerable attention. However, to the best of our knowledge, no integrated study of the effect of sediment deposition on the benthic processes that drive biogeochemical cycling (i.e. bioturbation and bio-irrigation) has hitherto been published. This study therefore aims to obtain a mechanistic understanding of sediment deposition effects on ecosystem functioning by experimentally assessing the impacts of deposition events of different magnitude (i.e. thickness of the deposited sediment layer) on benthic community diversity and biological traits (i.e. diversity, densities), benthic processes (i.e. bioturbation and bio-irrigation) and biogeochemical cycling in an intertidal soft-sediment habitat. We hypothesize that sediment deposition reduces oxygen availability to the community underneath, consequently affecting the survival of the macrobenthos and inducing escaping behaviour (Riedel et al., 2008; Villnäs et al., 2012). This may influence biogeochemical cycling, by affecting bioturbation or bio-irrigation (Van Colen et al., 2012; Renz and Forster, 2014).

2 Materials and Methods

2.1 Sample collection and experimental set-up

Samples were collected in March 2015 at the Paulina mudflat (SW Netherlands), which is located along the southern shore of the polyhaline part of the Scheldt estuary (51 ° 21.02 ‘N 3 ° 43.78 ‘E). The Scheldt estuary experiences a number of human-induced processes that can increase sediment deposition on tidal flats, among which dredging, and the local deposition of dredged sediments at the edges of tidal flats, are some of the most important examples (De Vriend et al., 2011; van der Wal et al., 2011). The Paulina mudflat harbours a
functionally rich benthic macrofaunal community that is numerically dominated by polychaetes (Van Colen et al., 2008, 2010).

Twenty-four cylindrical sediment corers (10 cm inner diameter, 29 cm length) were used to randomly collect cores within a 5 x 5 m patch of sediment, consisting of 46 ± 0.9 % mud (<63 µm), 22.9 ± 0.4 % very fine sand (63 – 125 µm), 21.7 ± 0.6 % fine sand (125 – 250 µm) and 9.4 ± 0.2 % medium sand (250 µm – 500 µm).

Additional sediment for the experimental deposition treatments had been collected at the same site a few days before the start of the experiment. This additional sediment was sieved over a 1 mm mesh, dried in the lab at 60 °C, heated in a muffle furnace at 500 °C to remove all organic matter (so that treatment effects could be unambiguously assigned to the physical smothering effect), rinsed with demineralized water, and subsequently sieved again.

All cores were cut to 9 cm, and each core was subsequently subjected to one of four treatments, each with six replicates. Each treatment except the control (T0) consisted of the application of a layer of the pre-treated sediment with a thickness of 1 (T1), 2 (T2) or 5 cm (T5), including a 0.5 cm thick frozen mud cake containing “Magenta” luminophores (Environmental Tracing Systems Ltd., Helensburgh, UK; median grain size 65 µm) and pre-treated sediment in a 1:1 volume:volume ratio to measure bioturbation activity. The control treatment only received a luminophore cake on top of the natural sediment surface. The addition of this mud cake ensured the quantification of particle mixing in these treatments and avoided potential bias between treatments due to species specific responses to the physico-chemical environment created by the mud cake. The addition of a luminophore mud cake on top of the sediment surface in the control treatment did not profoundly affect the natural oxygen fluxes or oxygen penetration depth. Our measured values were comparable in magnitude to those of previous studies in the same habitat and season (Van Colen et al., 2012; Annex 1), and clear bioturbation signs on the sediment surface soon after deposition indicate fast migration to the sediment-water interface (Annex 2).

Seawater from the sampling location (10 °C and a salinity of 20.3, kept still in barrels in the lab for half a day to allow suspended sediment to sink down) was carefully added on top of each core, up to the top edge of the corer. After addition of the water, the added sediment layers compacted to an average of 1.09 ± 0.18 (T1), 1.52 ± 0.10 (T2) and 3.75 ± 0.11 cm (T5), respectively. The cores were incubated in two tanks under ambient temperature and salinity conditions, filled until half the corer height to buffer for small changes in temperature, and provided with a constant air supply through bubbling underneath the water surface in each core. Each tank had a total capacity of 12 corers, and contained three replicates of each treatment. Oxygen did not penetrate deeper than the lower boundary of the deposited sediment layers in the deposition treatments, hence the sediment deposition created a physical barrier at the sediment-water interface prohibiting (passive) exchange of dissolved oxygen between the sampled community and the water column at the onset of the experiment (Annex 1). The experiment ran for 15 days, with different measurements taking place during this period. After letting the cores rest to regain biogeochemical equilibria, sediment oxygen profiles were measured on days 7 and 8, oxygen fluxes on day 12, followed by two days of measuring bio-irrigation and a final day on which the cores were sliced for further analysis.
2.2 Biogeochemical cycling

For the SCOC measurements, all cores were equipped with a magnetic stirring ring and sealed with an air-tight lid, fitted with two luer stopcocks enabling the sampling of the overlying water for the measurement of sediment-water column exchange of oxygen. During five hours (approximately one-hour intervals), 40-ml water samples were collected through one of the stopcocks using a glass syringe. Replacement water was added by opening the second stopcock and allowing tank water to flow in. The water samples were treated with Winkler reagents (Parsons et al., 1984) and stored at 4 °C until Winkler titration (Mettler Toledo G20, DGi 101-Mini oxygen electrode, LabX Light Titration software, Columbus, OH, USA). Sediment community oxygen consumption rates (SCOC) were then calculated from the linear decline in oxygen concentration, according to Eq. (1):

\[
SCOC = -\frac{dC}{dt} \frac{V}{A}
\]

(1)

where \(\frac{dC}{dt}\) is the change in oxygen concentration in the overlying water (in mmol L\(^{-1}\) d\(^{-1}\)), \(V\) is the volume of the overlying water (in L), and \(A\) is the sediment surface area (in m\(^2\)).

For the measurement of diffusive oxygen uptake (DOU), vertical sediment oxygen profiles were measured with a Unisense OX100 Clark-type needle electrode (Unisense, Aarhus, Denmark). Three profiles were measured in each core and the result was averaged, to account for spatial variability in the sediment. The DOU could then be calculated by multiplying the negative slope of the initial decrease in oxygen concentration, by its diffusion coefficient (Glud, 2008). The oxygen uptake that could be attributed to macrofaunal respiration was calculated by the formulae described in Mahaut et al. (1995), in which ash-free dry weights (AFDW), calculated from wet weights of the animals (see further) is used to calculate respiration rates:

\[
R = 0.0174 W^{0.0844}
\]

(2)

where \(R\) is the respiration rate in mg C d\(^{-1}\) and \(W\) the mean individual AFDW in mg C. The amount of carbon was estimated to be 50 % for all species (Wijsman et al., 1999). Since this formula is only valid for the temperature range of 15 to 20 °C, a Q\(_{10}\) of 2 was then assumed to correct the bias, and a respiratory quotient of 0.85 was used to calculate the oxygen consumption, here characterised as faunal uptake (FU; Braeckman et al., 2010; Mahaut et al., 1995). The remaining part of SCOC, after subtraction of DOU and FU, is the biologically-macrofauna-mediated oxygen uptake (BMUMU), caused indirectly by stimulation of aerobic remineralisation by macrofaunal bioturbation and irrigation.

2.3 Bio-irrigation and bioturbation

One day after the oxygen flux measurements, water was siphoned off from each core and replaced by a NaBr-seawater mixture to assess bio-irrigation. The NaBr solution had the same density as the seawater; both were mixed to obtain a solution with a final concentration of 0.1 M NaBr. The solution was added with 100 mL syringes on all cores until as close as possible to the edge, which amounted to 700 ml for T0, T1 and T2, and 600 ml for T5. A first sample of 2 ml was taken immediately after adding the mixture and subsequently after 1, 2, 18 and 21
hours. The bromide concentrations were measured with ion-chromatography and used to calculate bio-irrigation rates:

\[ Q = - \frac{V_{OW}}{C_{OW} - C_{PW}} \frac{dC_{OW}}{dt} \]  

(3)

where \( Q \) is the bio-irrigation rate, \( V_{OW} \) is the volume of the overlying water in L, \( C_{OW} \) is the initial concentration of bromide in the overlying water (mol L\(^{-1}\)), \( C_{PW} \) the bromide concentration in the pore water and \( \frac{dC_{OW}}{dt} \) the change of bromide concentration in the overlying water over time (in mol L\(^{-1}\) d\(^{-1}\)). For \( C_{PW} \), an estimation was made by measuring the background concentration in untreated seawater.

On the 14\(^{th} \) day of the experiment, the remaining water was siphoned off the cores, which were subsequently sliced per 5 mm from the top until 2 cm into the natural sediment. Deeper slices were cut at a thickness of 10 mm. The sediment in each slice was thoroughly homogenised, after which 5 to 10 mL was sampled and frozen at -20 °C, awaiting further processing for the quantification of bioturbation.

The samples were subsequently dried for 48 hours at 60 °C; water was then carefully added again, after which the sediment was spread open in a 55 mm inner diameter Petri dish. Each sample was photographed under UV light (365 nm peak wavelength) and luminophores were counted with computer scripts in Matlab v8.1 (MathWorks Inc., 2013) and R (R Development Core Team, 2013). A vertical profile of luminophore pixel counts was constructed for each sediment core and additional R scripts were used to fit the profiles to a non-local bioturbation model from which the biodiffusion coefficient \( D_{NL} \), in cm\(^2\) d\(^{-1}\) was calculated (Wheatcroft et al., 1990). Since luminophores were only applied on the sediment-water interface, the measured profiles represent disturbance of the surface by bioturbating fauna, rather than providing a total picture of the sediment mixing underneath the surface.

### 2.4 Macrofauna

The remaining 85 to 90 % of the sediment was rinsed over a 500 µm mesh-sized sieve to collect the macrofauna. The animals were stained with a Rose Bengal dye in order to facilitate the identification. Organisms were identified to species level, except for Oligochaeta and Spio sp. After identification, all animals were weighed to assess their biomass. The ash-free dry weight (AFDW) was determined by using conversion factors from wet weights (Sistermans et al., 2006). Biomasses were used to calculate the faunal respiration (Mahaut et al., 1995).

### 2.5 Data analysis

Diversity indices (Shannon-Wiener diversity \( H' \) (base e), Pielou’s evenness \( J' \) and species richness \( S \)) were calculated with Primer v6.1 (Clarke and Gorley, 2006). All taxa were assigned to functional groups based on their motility (from M1 – living fixed in a tube – till M4 – free three-dimensional movement through a burrow system) and sediment reworking activity (surficial modifiers, biodiffusors, upward conveyors and downward conveyors),
according to Queiros et al. (2013). All downward conveyors in our study were also classified as upward conveyors, since they can perform both sediment reworking activities.

Differences between the treatments for all biotic and abiotic variables, including all species` densities, were first tested by a 2-way ANOVA, where “Tank” and “Treatment” were used as factors. Since these analyses demonstrated that there were no interaction effects of tank and treatment, a blocked-design ANOVA was applied, with “Tank” as the blocking factor. A Tukey HSD test was used for pairwise comparisons in case of a significant treatment effect. In case the assumptions of normality (tested with a Shapiro-Wilk test) and homogeneity of variances (assessed with Levene’s test) for ANOVA were not met, a fourth-root transformation was performed on the data. Differences in community composition were tested with multivariate two-way permutational analysis of variance (PERMANOVA; Anderson et al., 2008). A Similarity Percentages analysis (SIMPER), based on a Bray-Curtis similarity matrix, was used to determine the species which contributed most to the differences between treatments. When a significant treatment effect was found, pairwise PERMANOVA tests were performed in order to detect differences between the treatments. The PERMANOVA tests were followed by a PERMDISP test to define whether the found effects are influenced by heterogeneity of multivariate dispersions.

Linear regressions were applied to find relationships between the different response variables. Most importantly, relationships were identified between ecosystem functioning (SCOC), benthic processes (bioturbation, bio-irrigation) and the various biotic variables, including densities of all individual species. Further regression tests investigated the contribution of individual species to the density – ecosystem functioning relationship, by using the densities of all taxa as predictor variables. The optimal model was selected via stepwise combined backward and forward selection. The variance inflation factor (VIF) was used to determine multicollinearity of the predictor variables. All assumptions for linear regression were tested on the residuals and met (no outliers and normal distribution).

All statistical analyses were performed with R v3.0.3 (R Development Core Team, 2013), except the PERMANOVA and SIMPER tests, for which Primer v6.1 with PERMANOVA+ add-on was used (Clarke and Gorley, 2006).

3 Results

3.1 Macrofauna

Sediment deposition affected community structure with the community present in T5 differing significantly from the control (2-factor Permanova pseudo-F = 2.457, \( P = 0.013 \); pair-wise comparisons T0-5: \( P = 0.010 \)). The PERMDISP test was not significant for either the main test or the pair-wise comparison (main test \( F = 0.858, P = 0.5795 \); T0-T5: \( P = 0.6282 \)). Species that contributed most to the dissimilarity in community structure between these treatments were *Aphelochaeta marioni* and *Oligochaeta* spp. (Table 1). Densities of *Polydora cornuta* (T0: 381.97 ± 131.50 ind m\(^{-2}\), T1: 169.77 ± 53.68 ind m\(^{-2}\), T2: 42.44 ± 26.84 ind m\(^{-2}\), T5: 0 ± 0 ind m\(^{-2}\)) and *Scrobicularia plana* (T0: 403.19 ± 60.77 ind m\(^{-2}\), T1: 381.97 ± 80.53 ind m\(^{-2}\), T2: 106.10 ± 51.11 ind m\(^{-2}\), T5: 106.10 ± 83.28 ind m\(^{-2}\)) (Table 2) were significantly lower in T5. (*P. cornuta* T0-T5: \( P = 0.003 \), T1-T5: \( P = 0.014 \); *S. plana* T0-T5: \( P = 0.007 \).
The control community had significantly higher total densities than the other communities (T0-T1: \( P = 0.011 \), T0-T2: \( P = 0.013 \), T0-T5: \( P = 0.001 \)), while lowest Shannon-Wiener diversity and species richness were found for the T5 community (Fig. 1, Table 23,4). Community evenness did not differ significantly among treatments.

In general, changes in macrobenthic community composition mirrored differential responses of specific motility and sediment reworking traits (Fig. 2, Table 23). Densities of the two groups of organisms with lowest motility were negatively affected by the applied treatments while densities of more motile species were not significantly different among treatments (Fig. 2a). The density of tube-building organisms (M1) decreased gradually with the thickness of the deposited sediment, whereas densities of species with limited movement (M2) were impaired by all sediment deposition treatments, irrespective of their magnitude (Fig. 2a).

All sediment reworking groups were affected by the deposition (Fig. 2b). For surficial modifiers, all treatments showed lower densities compared to the control, and for upward conveyors T5 was significantly lower than all other treatments (Surf. Mod. T0-T1: \( P = 0.033 \), T0-T2: \( P = 0.013 \), T0-T5: \( P = 0.006 \); Upw. Conv. T0-T5: \( P < 0.001 \), T1-T5: \( P = 0.009 \), T2-T5: \( P = 0.006 \) (Table 3,4). The density of biodiffusors was only significantly reduced in T5 compared to the control (\( P = 0.024 \) ) (Fig. 2b).

Activity of the macrofauna (bioturbation and bio-irrigation) was significantly affected by the deposition treatments (Table 24). Bioturbation activity was significantly higher in T1 than in all other treatments (T0-T1: \( P = 0.016 \), T1-T2: \( P = 0.048 \), T1-T5: \( P = 0.032 \) ) (Table 3,4), and was lowest in T5. While the biodiffusion coefficient \( D_b^{NL} \) reached average values in the control treatment, it rose significantly in T1 and dropped again in T2 and T5 (Fig. 3a). A similar pattern was observed for bio-irrigation, but here we only found a significant difference between T1 and T5 (\( P = 0.019 \) ) (Fig. 3b).

### 3.2 Ecosystem functioning

Sediment community oxygen consumption (SCOC) decreased with increasing thickness of the applied sediment layer, ranging from 54.68 ± 5.35 mmol m\(^{-2}\) d\(^{-1}\) in the control, over 46.79 ± 3.53 mmol m\(^{-2}\) d\(^{-1}\) in T1 and 44.37 ± 3.52 mmol m\(^{-2}\) d\(^{-1}\) in T2, to 40.68 ± 3.60 mmol m\(^{-2}\) d\(^{-1}\) in T5. Only T5 differed significantly from the control (\( P = 0.030 \) ) (Fig. 3c, Table 4). Faunal respiration (FU) accounted for 2.67 ± 1.01 % of the total SCOC in T0, 3.64 ± 1.64 % in T1, 1.75 ± 0.30 % in T2 and 1.99 ± 0.41 % in T5, while the DOU amounted for 18.55 ± 2.64 mmol m\(^{-2}\) d\(^{-1}\) in T0, 13.71 ± 1.80 mmol m\(^{-2}\) d\(^{-1}\) in T1, 11.56 ± 1.79 mmol m\(^{-2}\) d\(^{-1}\) in T2, and 16.37 ± 1.84 mmol m\(^{-2}\) d\(^{-1}\) in T5. Neither DOU nor FU showed any significant changes between treatments (Table 4), demonstrating the importance of biotiemacrofauna-mediated oxygen uptake (BMU-MMU) in the patterns of total SCOC.

Multiple linear regression showed that the variability in SCOC was significantly related to total macrofaunal density and \( D_b^{NL} \), explaining together 54.4% of the variability in SCOC (\( P < 0.001 \)). When total density was divided over the functional groups, we found significant relationships with \( D_b^{NL} \) and motility groups M2 and M3 (\( P = 0.001 \); \( R^2 = 0.53 \)), and with surficial modifiers and biodiffusors (\( P < 0.001 \); \( R^2 = 0.56 \)). Other variables of community diversity (Shannon-Wiener diversity, species richness, and Pielou’s evenness) were not significant
predictors of ecosystem functioning. While no single species was found to contribute significantly to $D_{ob}$, a combination of several species contributed significantly to the variability in SCOC ($P < 0.001; R^2 = 0.56$). The taxa with a significant contribution were *A. marioni* and *Cyathura carinata* (Table 5). The statistically optimal model for bio-irrigation included *Hediste diversicolor* and *P. cornuta* as positive contributors to this process ($P < 0.001; R^2 = 0.73$) (Table 5).

4 Discussion

Tidal flats are dynamic, sedimentary environments that naturally undergo processes of erosion and deposition. Per tidal cycle, different elevation changes have been observed, e.g. from decreases of 3.3 mm in the Yangzte estuary (China) to increases of 6 mm in the estuary of the Seine (France) (Deloffre et al., 2007; Shi et al., 2012). Our study was performed on a mudflat in the estuary of the river Scheldt (Belgium, the Netherlands), which is characterized by its meso- to macro-tidal regime and well-mixed water column. Sediment input from the river basin is relatively low and sand extraction and sea level rise lead to a net export of sediment from the estuary (De Vriend et al., 2011). Sediment accretion on the estuary’s tidal flat can amount to about 2 cm yr$^{-1}$ (Weerman et al., 2011; Widdows et al., 2001), which suggests that natural sedimentation on the intertidal mudflat is unlikely to exceed even a few millimetres per tidal cycle. More extreme changes in the bed level of mudflat can however happen during storm events, either by erosion of the top centimetres of the sediment or by deposition of new sediment (Hu et al., 2015; Marion et al., 2009). Besides natural processes, anthropogenic factors influencing sedimentation are prominent in the estuary, among which dredging in the main channels to ensure access to the port of Antwerp, and dumping of the dredged material to retain sediment within the estuary, are the most important (Jeuken and Wang, 2010; Meire et al., 2005). Most of this dredged sediment is disposed of near shoals and tidal flats, and can as such affect the intertidal ecosystem (Bolam and Whomersley, 2005; De Vriend et al., 2011; Zheng, 2015).

Our results show that even thin sediment deposits can cause a drop in total macrofaunal density, mainly by impacting the highly abundant surface-dwelling animals with low motility (Figs 1-2a,b). These animals, which belong to reworking and motility class 2 due to their sessile lifestyle (Solan et al., 2004), lack the capacity to escape the deposited sediment and are not adapted to living in deeper sediment layers (Essink, 1999). Since the oxygen penetration depth never exceeded the thickness of the deposited sediment layer (Annex 1), we can assume that oxygen stress was a major driver for the observed decrease in faunal densities. In treatments T1 and T2, oxygen stress was possibly reduced by the increased activity of the macrofauna, due to the animals still being able to disturb the surface and oxygenate the underlying sediment. Hypoxia can induce escaping behaviour in benthic fauna, as observed in our intermediate treatments, and increase mortality when more severe (Riedel et al., 2008; Villnäs et al., 2012).

Being identified as significant contributors to changes in SCOC, surface-dwelling and low-motile animals are expected to show density patterns similar to those of SCOC itself. However, SCOC only gradually declined with increasing thickness of the deposited sediment, and this decrease became significant only in the most extreme treatment (T5). Since DOU proved to be constant over all treatments and macrofaunal respiration was negligible compared to the total oxygen consumption, the observed changes in SCOC could be attributed to oxygen uptake.
caused indirectly by activity of the benthos (i.e. bioturbation and/or bio-irrigation). However, both bio-irrigation and bioturbation, the latter of which was linearly related to SCOC, showed that activity increased in treatments T1 and T2. This activity was likely caused by animals for which we found a linear relationship with bioturbation or bio-irrigation, like *H. diversicolor*, that are highly mobile and can bury upwards towards the surface, thereby partly irrigating the sediment. *Hediste diversicolor* is a ‘gallery-diffusor’, which combines biodiffusion in a dense gallery system with biotransport to the bottoms of the tubes (François et al., 2002; Hedman et al., 2011), as well as a well-known bioirrigator (Kristensen and Hansen, 1999; Riisgaard and Larsen, 2005). Its activity can be expected to result in the oxygenation of deeper sediment layers, but this effect was probably not sufficient to counteract the loss of less mobile, surface-dwelling fauna. Consequently, we observed a gradual and significant decline in SCOC, caused by the disappearance of an abundant group of organisms. Upon addition of the thick sediment layer in treatment T5, species richness dropped significantly and the densities of upward conveyors decreased considerably, hence preventing the transport of organically rich deep sediment to the surface, through the deposited layer. As a result, the deposited sediment essentially functioned as a barrier, preventing contact between sediment organic matter and oxygen in the water column, and therefore reducing microbial degradation and respiration.

Through alterations in functional trait abundances and community composition, natural and anthropogenic disturbances can affect the entire ecosystem functioning (Bolam et al., 2002; Rodil et al., 2011). In the case of burial by sediment deposition, our experiment revealed that SCOC can be affected by causing mortality among surface-dwelling and low motile animals, forming the most abundant functional groups of macrobenthos in our system. Macrobenthic diversity and abundance have been shown to exert some control on the magnitude of solute fluxes across the sediment-water interface (Herman et al., 1999; Thrush et al., 2006). Furthermore, previous studies have shown that functional traits of species can be of great importance to explain ecosystem functioning, rather than or additional to taxonomic diversity (Braeckman et al., 2010; Hooper et al., 2005). Our results highlight the importance of both macrofaunal densities, and the functional identity of species. It is clear that taxonomic diversity alone was not sufficient to explain the changes in ecosystem functioning in our experiment, whereas closer inspection of the functional identities provided more realistic insights.

It should be noted that the sediment we used for deposition was completely defaunated and did not contain organic matter. Whereas the aim of using defaunated sediment was to allow a better mechanistic understanding of the consequences of sediment deposition, it does not reflect natural conditions. Dredged material from the bottom of the estuary is much richer in organic material and might lead to different results in a similar experiment. Cottrell et al. (2016) showed that benthic species can have a variable tolerance for changes in the enrichment of the sediment, with higher mortalities under high organic loading (and hence likely stronger impacts on macrofauna-mediated biogeochemical cycling).

## 5 Conclusion

Our experiment revealed new insights into the effects of sediment deposition on the intertidal benthic ecosystem. We found a negative effect on ecosystem functioning, with alterations in macrofauna community structure and activity as the underlying mechanisms. With increasing thickness of the deposited sediment layer, a shift to lower
densities of low-motile and surface-dwelling animals resulted in decreased functioning, even though this was initially dampened by an increased activity of more motile and deeper-living fauna. The latter were responsible for a sustained oxygen penetration through the deposited layer under intermediate treatments, but failed to efficiently do so under more extreme circumstances. It was clear that taxonomic diversity did not suffice to explain changes in functioning, while the functional identity of species did give us important additional insights.

Data availability

All data will be deposited in the VLIZ Marine Data Archive (http://mda.vliz.be/introduction.php).

Author contributions

SM, LB and CVC devised the experiments. SM and LB carried out the experimental work and collected all data. SM and CVC led the writing of the manuscript, to which all authors contributed. All authors declare that they do not have any conflict of interest.

Acknowledgements

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References


TABLE 1: The three species with highest cumulative contribution (> 50 %) to the total dissimilarity between treatments*. The first column shows the treatments being compared (e.g. T0-1: a comparison between treatments T0 and T1).

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Average dissimilarity</th>
<th>Species</th>
<th>Cumulative contribution</th>
</tr>
</thead>
<tbody>
<tr>
<td>T0-1</td>
<td>42.14</td>
<td><em>Aphelochaeta marioni</em></td>
<td>37.61 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Oligochaeta spp.</td>
<td>59.97 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td><em>Polydora cornuta</em></td>
<td>65.83 %</td>
</tr>
<tr>
<td>T0-2</td>
<td>36.49</td>
<td><em>Aphelochaeta marioni</em></td>
<td>37.86 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Oligochaeta spp.</td>
<td>54.76 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td><em>Polydora cornuta</em></td>
<td>62.00 %</td>
</tr>
<tr>
<td>T0-5</td>
<td>48.60</td>
<td><em>Aphelochaeta marioni</em></td>
<td>35.25 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Oligochaeta spp.</td>
<td>57.60 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td><em>Polydora cornuta</em></td>
<td>64.39 %</td>
</tr>
<tr>
<td>T1-2</td>
<td>38.74</td>
<td>Oligochaeta spp.</td>
<td>26.49 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td><em>Aphelochaeta marioni</em></td>
<td>52.01 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td><em>Hediste diversicolor</em></td>
<td>60.03 %</td>
</tr>
<tr>
<td>T1-5</td>
<td>42.42</td>
<td><em>Aphelochaeta marioni</em></td>
<td>24.20 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Oligochaeta spp.</td>
<td>46.10 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td><em>Scrobicularia plana</em></td>
<td>56.55 %</td>
</tr>
<tr>
<td>T2-5</td>
<td>41.15</td>
<td>Oligochaeta spp.</td>
<td>31.12 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td><em>Aphelochaeta marioni</em></td>
<td>56.73 %</td>
</tr>
<tr>
<td></td>
<td></td>
<td><em>Hediste diversicolor</em></td>
<td>65.37 %</td>
</tr>
</tbody>
</table>

* Results from a SIMPER analysis
TABLE 2: Densities (in ind m$^{-2}$) of all identified taxa in the macrobenthic communities. All values are means ± standard errors.

<table>
<thead>
<tr>
<th>Species</th>
<th>T0</th>
<th>T1</th>
<th>T2</th>
<th>T5</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Polychaeta</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Aphelochaeta marioni</td>
<td>3225.54 ± 724.49</td>
<td>1379.34 ± 388.17</td>
<td>1570.33 ± 358.12</td>
<td>1167.14 ± 267.92</td>
</tr>
<tr>
<td>Eteone longa</td>
<td>21.11 ± 21.22</td>
<td>84.88 ± 42.44</td>
<td>63.66 ± 28.47</td>
<td>21.11 ± 21.22</td>
</tr>
<tr>
<td>Hediste diversicolor</td>
<td>594.18 ± 107.37</td>
<td>551.74 ± 121.53</td>
<td>530.52 ± 129.08</td>
<td>233.43 ± 60.77</td>
</tr>
<tr>
<td>Heteromastus filiformis</td>
<td>254.65 ± 73.51</td>
<td>127.32 ± 46.49</td>
<td>254.65 ± 131.50</td>
<td>84.88 ± 26.84</td>
</tr>
<tr>
<td>Polydora cornuta</td>
<td>381.97 ± 131.50</td>
<td>169.77 ± 53.68</td>
<td>42.44 ± 26.84</td>
<td>0.00 ± 0.00</td>
</tr>
<tr>
<td>Pygospio elegans</td>
<td>297.09 ± 102.21</td>
<td>148.54 ± 76.51</td>
<td>169.77 ± 42.44</td>
<td>0.00 ± 0.00</td>
</tr>
<tr>
<td>Spio sp.</td>
<td>21.22 ± 21.22</td>
<td>0.00 ± 0.00</td>
<td>0.00 ± 0.00</td>
<td>0.00 ± 0.00</td>
</tr>
<tr>
<td>Streblospio benedicti</td>
<td>63.66 ± 43.49</td>
<td>0.00 ± 0.00</td>
<td>42.44 ± 26.84</td>
<td>0.00 ± 0.00</td>
</tr>
<tr>
<td><strong>Oligochaeta spp.</strong></td>
<td>2058.40 ± 343.88</td>
<td>997.37 ± 271.92</td>
<td>1846.20 ± 251.98</td>
<td>933.71 ± 295.26</td>
</tr>
<tr>
<td><strong>Bivalvia</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cerastoderma edule</td>
<td>42.44 ± 26.84</td>
<td>42.44 ± 26.84</td>
<td>0.00 ± 0.00</td>
<td>0.00 ± 0.00</td>
</tr>
<tr>
<td>Macoma balthica</td>
<td>63.66 ± 43.49</td>
<td>233.43 ± 76.51</td>
<td>127.32 ± 32.87</td>
<td>148.54 ± 51.11</td>
</tr>
<tr>
<td>Scrobicularia plana</td>
<td>403.19 ± 60.77</td>
<td>381.97 ± 80.53</td>
<td>106.10 ± 51.11</td>
<td>106.10 ± 83.28</td>
</tr>
<tr>
<td><strong>Gastropoda</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Hydrobia ulvae</td>
<td>106.10 ± 51.11</td>
<td>169.77 ± 53.68</td>
<td>148.54 ± 60.77</td>
<td>212.21 ± 117.00</td>
</tr>
<tr>
<td><strong>Crustacea</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bathyporeia pilosa</td>
<td>0.00 ± 0.00</td>
<td>21.22 ± 21.22</td>
<td>0.00 ± 0.00</td>
<td>21.22 ± 21.22</td>
</tr>
<tr>
<td>Cyathura carinata</td>
<td>636.62 ± 103.96</td>
<td>424.41 ± 78.26</td>
<td>445.63 ± 107.79</td>
<td>509.30 ± 65.75</td>
</tr>
</tbody>
</table>
TABLE 3: Statistical factors from 2-factor blocked ANOVA tests with ‘Treatment’ (4 levels) and ‘Tank’ (2 levels) as factors. M1 till M4 stand for motility classes, as defined by Solan et al. (2004) (M1: living fixed in a tube, M2: sessile, but not fixed in a tube, M3: slow movement through the sediment, M4: free movement in a burrow system). Significant pair-wise differences between treatments are given in the table. All results for species and functional groups are given for densities. Results for the factor ‘Treatment’ from a 2-factor blocked ANOVA tests with ‘Treatment’ (4 levels) and ‘Tank’ (2 levels) as factors. M1 till M4 stand for motility classes, as defined by Solan et al. (2004) (M1: living fixed in a tube, M2: sessile, but not fixed in a tube, M3: slow movement through the sediment, M4: free movement in a burrow system). Significant pair-wise differences between treatments are given in the table. In case of heterogeneity of the variances, a fourth root transformation was applied on the data.

<table>
<thead>
<tr>
<th>Source</th>
<th>F value</th>
<th>P</th>
<th>Pair-wise significance</th>
<th>Transformation</th>
</tr>
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<tr>
<td>M1</td>
<td>12.221</td>
<td>&lt;0.001*</td>
<td>0-5, 1-5, 2-5</td>
<td>Fourth root</td>
</tr>
<tr>
<td>M2</td>
<td>7.013</td>
<td>0.002*</td>
<td>0-1, 0-2, 0-5</td>
<td></td>
</tr>
<tr>
<td>M3</td>
<td>3.05</td>
<td>0.054</td>
<td></td>
<td></td>
</tr>
<tr>
<td>M4</td>
<td>2.284</td>
<td>0.112</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Surficial modifiers</td>
<td>6.087</td>
<td>0.004*</td>
<td>0-1, 0-2, 0-5</td>
<td></td>
</tr>
<tr>
<td>Biodiffusors</td>
<td>4.336</td>
<td>0.017*</td>
<td>0-5</td>
<td></td>
</tr>
<tr>
<td>Upward conveyors</td>
<td>10.112</td>
<td>&lt;0.001*</td>
<td>0-1, 0-2, 0-5</td>
<td></td>
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<tr>
<td>Downward conveyors</td>
<td>24.371</td>
<td>&lt;0.001*</td>
<td>0-5, 1-5, 2-5</td>
<td>Fourth root</td>
</tr>
</tbody>
</table>

**Polychaeta**

- *Aphelochaeta marioni* 4.648 0.013* 0-1, 0-5
- *Eteone longa* 1.103 0.372
- *Hediste diversicolor* 2.284 0.112
- *Heteromastus filiformis* 1.154 0.353
- *Polydora cornuta* 7.254 0.002* 0-2, 0-5, 1-5 Fourth root
- *Pygospio elegans* 5.155 0.009* 0-5, 2-5 Fourth root
- *Spio sp.* 7.254 0.002* 0-2, 0-5, 1-5 Fourth root
- *Streblospio benedicti* 1.879 0.167

**Oligochaeta spp.** 3.873 0.026* None

**Bivalvia**

- *Cerastoderma edule* 1.583 0.226
- *Limecola balthica* 1.939 0.158
- *Scrobicularia plana* 5.337 0.008* 0-2, 0-5

**Gastropoda**

- *Peringia ulvae* 0.329 0.804

**Crustacea**

- *Bathyporeia pilosa* 0.704 0.561
- *Cyathura carinata* 1.055 0.391
- $D_b^{NL}$ 4.826 0.012* 0-1, 1-2, 1-5 Fourth root
- Q 4.177 0.020* 1-5
<table>
<thead>
<tr>
<th></th>
<th>Value</th>
<th>Standard Error</th>
<th>Significance</th>
<th>Depth Range</th>
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<tr>
<td>SCOC</td>
<td>3.358</td>
<td>0.041*</td>
<td></td>
<td>0-5</td>
</tr>
<tr>
<td>DOU</td>
<td>2.178</td>
<td>0.124</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FU</td>
<td>0.869</td>
<td>0.475</td>
<td></td>
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</tr>
<tr>
<td>Total density</td>
<td>8.346</td>
<td>0.001*</td>
<td></td>
<td>0-1, 0-2, 0-5</td>
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<tr>
<td>H’</td>
<td>4.983</td>
<td>0.010*</td>
<td></td>
<td>1-5</td>
</tr>
<tr>
<td>J’</td>
<td>2.594</td>
<td>0.083</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Species richness</td>
<td>6.697</td>
<td>0.003*</td>
<td></td>
<td>0-5, 1-5, 2-5</td>
</tr>
</tbody>
</table>
TABLE 4: Overview of the p-values for all pair-wise tests (Tukey post-hoc test), performed when the main test provided significant results. All results for species and functional groups represent densities.

<table>
<thead>
<tr>
<th>Source</th>
<th>T0-T1</th>
<th>T0-T2</th>
<th>T0-T5</th>
<th>T1-T2</th>
<th>T1-T5</th>
<th>T2-T5</th>
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<tbody>
<tr>
<td>M1</td>
<td>0.466</td>
<td>0.312</td>
<td>&lt; 0.001*</td>
<td>0.990</td>
<td>0.0028</td>
<td>0.004*</td>
</tr>
<tr>
<td>M2</td>
<td>0.017*</td>
<td>0.015*</td>
<td>0.002*</td>
<td>1.000</td>
<td>0.805</td>
<td>0.838</td>
</tr>
<tr>
<td>Surficial modifiers</td>
<td>0.033*</td>
<td>0.013*</td>
<td>0.006*</td>
<td>0.974</td>
<td>0.850</td>
<td>0.980</td>
</tr>
<tr>
<td>Upward conveyors</td>
<td>0.016*</td>
<td>0.036*</td>
<td>&lt; 0.001*</td>
<td>0.982</td>
<td>0.186</td>
<td>0.095</td>
</tr>
<tr>
<td>Downward conveyors</td>
<td>0.102</td>
<td>0.289</td>
<td>&lt; 0.001*</td>
<td>0.927</td>
<td>&lt; 0.001*</td>
<td>&lt; 0.001*</td>
</tr>
<tr>
<td>Biodiffusors</td>
<td>0.156</td>
<td>0.959</td>
<td>0.024*</td>
<td>0.344</td>
<td>0.780</td>
<td>0.067</td>
</tr>
<tr>
<td>Aphelochaeta marioni</td>
<td>0.035*</td>
<td>0.065</td>
<td>0.017*</td>
<td>0.989</td>
<td>0.986</td>
<td>0.913</td>
</tr>
<tr>
<td>Polydora cornuta</td>
<td>0.896</td>
<td>0.044*</td>
<td>0.003*</td>
<td>0.167</td>
<td>0.014*</td>
<td>0.611</td>
</tr>
<tr>
<td>Pygospio elegans</td>
<td>0.463</td>
<td>0.981</td>
<td>0.010*</td>
<td>0.687</td>
<td>0.194</td>
<td>0.023*</td>
</tr>
<tr>
<td>Scrobicularia plana</td>
<td>0.997</td>
<td>0.039*</td>
<td>0.039*</td>
<td>0.060</td>
<td>0.060</td>
<td>1.000</td>
</tr>
<tr>
<td>$D_b^{NL}$</td>
<td>0.016*</td>
<td>0.949</td>
<td>0.087</td>
<td>0.048*</td>
<td>0.032*</td>
<td>0.997</td>
</tr>
<tr>
<td>Q</td>
<td>0.104</td>
<td>0.705</td>
<td>0.794</td>
<td>0.541</td>
<td>0.016*</td>
<td>0.222</td>
</tr>
<tr>
<td>SCOC</td>
<td>0.338</td>
<td>0.145</td>
<td>0.030*</td>
<td>0.951</td>
<td>0.552</td>
<td>0.850</td>
</tr>
<tr>
<td>Total density</td>
<td>0.011*</td>
<td>0.043*</td>
<td>0.001*</td>
<td>0.921</td>
<td>0.560</td>
<td>0.240</td>
</tr>
<tr>
<td>H'</td>
<td>0.430</td>
<td>0.721</td>
<td>0.171</td>
<td>0.076</td>
<td>0.007*</td>
<td>0.691</td>
</tr>
<tr>
<td>Species richness</td>
<td>0.973</td>
<td>0.918</td>
<td>0.009*</td>
<td>0.714</td>
<td>0.003*</td>
<td>0.035*</td>
</tr>
</tbody>
</table>

Significant P-values (P < 0.05) are indicated with *
TABLE 5: Linear regressions of sediment community oxygen consumption (SCOC) against sets of species (or functional group) densities, and ecosystem processes (bio-irrigation - Q - and bioturbation - $D_b^{NL}$), and of bio-irrigation against the densities of species. Only significant models (P (slope) < 0.05) were considered. M2 and M3 are motility classes as defined by Solan et al. (2004) – M2: sessile, but not fixed in a tube, M3: slow movement through the sediment.

<table>
<thead>
<tr>
<th>Response/predictor</th>
<th>Regression equation</th>
<th>$R^2$</th>
<th>P</th>
</tr>
</thead>
</table>
| SCOC               | $x_1$: Total density  
$y = 3.35 \times 10^{-3} x_1 + 1.03 \times 10^2 x_2 + 25.6$  
$x_2$: $D_b^{NL}$ | 0.544 | 0.0001 |
|                    | $x_1$: M2  
$y = 3.16 \times 10^{-3} x_1 + 5.43 \times 10^{-3} x_2 + 1.02 \times 10^2 x_3$  
$x_2$: M3  
$x_3$: $D_b^{NL}$ | 0.529 | 0.0176 |
| SCOC               | $x_1$: Surficial modifiers  
$y = 2.92 \times 10^{-3} x_1 + 5.63 \times 10^{-3} x_2 + 1.05 \times 10^2 x_3$  
$x_2$: Biodiffusors  
$x_3$: $D_b^{NL}$ | 0.557 | 0.0135 |
| SCOC               | $x_1$: A. marioni  
$y = 4.53 \times 10^{-3} x_1 + 2.52 \times 10^{-2} x_2 + 25.9$  
$x_2$: C. carinata | 0.556 | 0.0008 |
| Q                  | $x_1$: A. marioni  
$y = -5.76 \times 10^{-4} x_1 + 5.00 \times 10^{-3} x_2 + 3.81 \times 10^{-5} x_3$  
$x_2$: H. diversicolor  
$x_3$: P. cornuta  
$x_4$: P. elegans  
$x_5$: S. benedicti | 0.730 | 0.0002 |
|                    |  | 0.0330 | 0.0030 |
|                    |  | 0.0002 | 0.0306 |
|                    |  | 0.0068 | 0.0030 |
Figure 1: Bar charts representing total macrofaunal densities (ind m$^{-2}$), species richness, Shannon-Wiener diversity, and Pielou's evenness per treatment. Error bars represent mean ± standard error, letters above the error bars indicate pair-wise significant differences. The four treatments represent the thickness of the applied sediment layer (in cm).
Figure 2: (a) Bar chart showing the densities of the four motility classes per treatment, in ind m\(^{-2}\). M1: organisms living fixed in a tube, M2: sessile, but not fixed in a tube, M3: slowly moving organisms, M4: free movement through a burrow system. (b) Bar chart showing the densities in, ind m\(^{-2}\), of the four main functional groups, based on sediment reworking activity. S: Surficial modifiers, B: biodiffusors, UC: upward conveyors, DC: downward conveyors. Error bars represent mean ± standard error, letters above the error bars indicate pair-wise significant differences. The four treatments represent the thickness of the applied sediment layer (in cm).
Figure 3: (a) Bar chart representing the mean bioturbation activity (by means of the biodiffusion coefficient $D^N_L$, in cm² d⁻¹) per treatment ± standard error. (b) Bar chart representing the mean bio-irrigation (in mL min⁻¹) per treatment ± standard error. (c) Bar chart representing the mean oxygen consumption (in mmol m⁻² d⁻¹) per treatment ± standard error. The different components of total sediment community oxygen consumption (SCOC) are represented in the chart: diffusive oxygen uptake (DOU), with error bars, faunal uptake (FU), with error bars, and the remaining benthicmacrofauna-mediated oxygen uptake (BMUMU). The topmost error bars represent the mean ± standard error of the total SCOC (= DOU + FU + BMU). Letters above the error bars indicate pair-wise significant differences. The four treatments represent the thickness of the applied sediment layer (in cm).