The drivers of biogeochemistry in beach ecosystems: A cross-shore transect from the dunes to the low-water line

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A B S T R A C T

This study addresses key processes in high-energy beach systems using an interdisciplinary approach. We assess spatial variations in subsurface pore water residence times, salinity, organic matter (OM) availability, and redox conditions and their effects on nutrient cycles as well as on microbial community patterns and microphytobenthos growth. At the study site on Spiekeroog Island, southern North Sea, beach hydrology is characterized by the classical zonation with an upper saline plume (USP), a saltwater wedge, and a freshwater discharge tube in between. Sediment and pore water samples were taken along a cross-shore transect from the dunes to the low-water line reaching sediment depths down to 5 m below sediment surface. Spatial variations in pore water residence time, salinity, and organic matter availability lead to steep redox and nutrient gradients. Vertical and horizontal differences in the microbial community indicate the influence of these gradients and salinity on the community structure. Modeled seawater flux through the USP and freshwater flux through the tube are on average 2.8 and 0.75 m³ per day and meter of shoreline, respectively. Furthermore, ridge sediments at the lower beach discharge seawater at rates of 0.5 and 1.0 m³ per day and meter of shoreline towards the runnel and seaside, respectively. Applying seawater and freshwater fluxes and representative nutrient concentrations for the discharge zones, nutrient fluxes to adjacent nearshore waters are 117 mmol NH₄ +, 55 mmol PO₄ ³⁻, and 575 mmol Si(OH)₄ per day and meter of shoreline. We propose that this nutrient efflux triggers growth of microphytobenthos on sediment surfaces of the discharge zone. A first comparison of nutrient discharge rates of the beach site with a nearby sandy backbarrier tidal flat margin indicates that the beach system might be of less importance in supplying recycled nutrients to nearshore waters than the backbarrier tidal flat area.

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1. Introduction

Sandy beaches as coastal ecosystems provide important ecosystem services, such as reduction of storm impacts, nutrient cycling, water purification, nursery and feeding-breeding habitats (Nel et al., 2014 and references therein). However, beach ecosystems are vulnerable and may be affected, for example, by erosion caused by climate-induced sea level rise (Hinkel et al., 2013) or increased nitrogen availability due to human activities (Ibánhez and Rocha, 2014). To preserve the ecosystem services and to assess the vulnerability of beach ecosystems, it is a prerequisite to develop a detailed understanding of the biogeochemical processes and dynamics in these systems. However, there is a lack of understanding compared to other coastal systems such as estuaries, coral reefs, and mangroves (Nel et al., 2014).

Sandy beach systems are an important part of the global land-sea interface because they make up two-thirds of the world’s ice free coastline (Mclachlan and Brown, 2006). Several studies show that sandy beach sediments exert a major effect on coastal carbon, nutrient, and trace element cycling (Anschutz et al., 2016, 2009; Charbonnier et al., 2013; Heiss and Michael, 2014; Rodellas et al., 2014; Santoro, 2010; Santos et al., 2008). Sandy sediments are characterized by a high permeability,
which permits advective pore water flow besides diffusion (Huet et al., 1998; Riedel et al., 2010). Consequently, seawater infiltration continuously supplies oxygen and fresh organic matter to the sediment leading to high levels of organic matter (OM) degradation (Huet et al., 2014) and nutrient cycling (Gao et al., 2012; Marchant et al., 2014). When remineralization products are discharged into the coastal zone via submarine groundwater discharge (SGD), they can stimulate primary production and may even cause harmful algae blooms and eutrophication (Paerl, 1997). The phenomenon of SGD is observed worldwide (Moore, 1996) and the global magnitude of fresh and saline SGD is estimated to be larger than riverine fluxes to the ocean (Kwon et al., 2014; Moore et al., 2008). However, the contribution of beach systems to global SGD budgets and consequently the global importance of beaches for the cycling of nutrients, carbon species, and trace elements need to be studied in more detail.

Beach ecosystems represent major transition zones (land-sea, freshwater-saltwater, oxic-anoxic). Biogeochemical processes are controlled by groundwater flow pathways and residence times, which depend on physical factors such as tidal amplitude, wave height, seasonal changes in water table elevation (Anwar et al., 2014; Gibbes et al., 2008; Michael et al., 2005; Robinson et al., 2007b, 2014), beach morphology, and sediment permeability (Bakhtyar et al., 2011; Butt et al., 2001). Additionally, chemical gradients related to OM flux and redox changes influence the ecosystem (Charette and Sholkovitz, 2006; Loveless and Oldham, 2010; McMullister et al., 2015; Santos et al., 2008). Therefore, the biota in these ecosystems have to cope with strong chemical and physical constraints (Harris et al., 2011; Hubas et al., 2007b; Rodil and Lasra, 2004). Nevertheless, it was shown that despite these steep gradients, beach systems exhibit a high microbial activity (Forehead et al., 2013) and that the microbial community can modulate its growth efficiency when exposed to environmental stress (Hubas et al., 2007a).

Although beach ecosystems might be of importance for global biogeochemical cycles, significant disciplinary and interdisciplinary gaps in understanding beach ecosystem functioning remain because most studies were uni-disciplinary and focused either on ecology, physical beach environment, management, or biology (Dugan et al., 2010; Nel et al., 2014). Therefore, we conducted a study that combines the disciplines sedimentology, hydrology, biogeochemistry, microbiology, and ecology. We assessed the effects of spatial variations in subsurface pore water residence times, salinity, OM availability, and redox conditions on OM and nutrient cycling as well as on microbial community patterns and microphytobenthos growth. Furthermore, we tested the hypothesis that exposed beach systems facing the open North Sea are of greater importance for nutrient recycling and efflux than sandy backbarrier tidal flat sediments.

2. Material and methods

2.1. Study area and sampling strategy

Spiekeroog forms part of the barrier island chain in front of the North-west German coastline (Fig. 1). The island has a length of 9.8 km and a maximum width of 2 km. Barrier islands like Spiekeroog are typically characterized by a freshwater lens, which develops as precipitation continuously infiltrates into the often highly permeable sediments and overlies the saltwater (Röper et al., 2010). In addition to wind and wave driven processes, the island’s beaches are affected by semidiurnal tides with a mean tidal range of 2.7 m (mesotidal). The offshore wave spectrum varies from 0.5–2 m at fair weather up to 11 m during storm conditions (Dobrynin et al., 2010). The beach slope of 2.3° from the backshore to the foreshore, measured during the sampling campaign in May 2014, reflects the transitional stage of beach morphology from the dissipative winter situation to the reflective summer situation, classified after Short and Woodruff (2009) (Fig. 2). A winter berm is well developed directly at the transition from the backshore to the foreshore. The foreshore is dissected by a ridge and runnel system, which is oriented slightly shore-oblique. During falling tide, the runnels drain the emerged parts of the beach towards the East.

In line with the gradients in hydrodynamic energy and topography, we have chosen a land-sea transect on the exposed seaward-facing side of Spiekeroog Island for the investigations (Fig. 1). During a joint sampling campaign in May 2014, sediment, pore water, and water column samples were collected. The main beach transect consisted of five sites extending from close the dunes (site 0 and 1, both behind a backshore berm), to the high tide line (site 2), to a ridge (site 3), and finally to the low-water line (site 4). Sedimentological studies extended to the shoreface as well including the two subtidal sites 5 and 6 (Fig. 2). Furthermore, phytobenthos samples were collected along three transects (with 50 m distance in between), stretching from the high to the low-water line (Phytobenthos 1–11; Fig. 2). Hydrological data were recorded close to sites 0, 2, and 3 (GWM 0, 2, 3; Fig. 2).
Fig. 2. Beach topography of the sampling transect stretching the back and foreshore from the rise of the dunes to the low-water line (sites 0–4) and finally reaching the shoreface (sites 5 and 6). The depth profiles indicate the mean grain size distribution at each site, the gray shaded area marks the grain size range of fine sand. At sites 0–4 sediment and pore water were jointly sampled and analyzed with respect to sedimentological, biogeochemical, and microbiological aspects. Hydrological data were recorded close to sites 0, 2, and 3 (GWM 0, 2, 3). Phytobenthos samples were collected along three transects covering approximately the distance between sites 2 to 4 of the main transect (Phytobenthos 1–11). At sites 5 and 6 only the sedimentology was studied.
2.2. Sample and data collection

2.2.1. Measurement of beach topography

The beach topography and study sites were mapped with a Trimble 5800 GPS using a data correction service, which minimizes the accuracy error of the geographic position (WGS 84) to 12 mm both vertically and horizontally.

2.2.2. Hydraulic investigations

Three groundwater monitoring wells (GWM) were installed along the transect. (Figs. 2, 3). At these wells, groundwater level measurements were continuously recorded with pressure transducers (Schlumberger Mini-Diver) for a duration of 3 days. All measured groundwater levels were corrected for the groundwater specific density and air pressure.

2.2.3. Sediment sampling

Sediment cores were taken at seven sites (five at the backshore/foreshore and two offshore ones; Fig. 2). The offshore cores at sites 5 and 6 were taken on board the research vessel "Senckenberg" using an electric underwater vibrocorer. Onshore cores were taken by vibrocoring as described by Wilms et al. (2006). Sediment was recovered in aluminum tubes (onshore, 8 cm inner diameter) or PVC-liners (offshore, 10 cm inner diameter). For biogeochemical and molecular analyses, sediment cores were transported to the lab and immediately processed. Cores were cut transversely at respective depth intervals to take subcores (inner diameter). For sedimentological analyses cores were cut lengthwise.

For analyzing particulate nutrient and chlorophyll a concentrations of the benthic microalgae community, sediment corers made of hard plastic material with a length of 20 cm and a diameter of 1.9 cm were used. All cores were filled by putting 3 cm of each corer into the sediment and closing them with rubber stoppers at both sides.

2.2.4. Water column sampling

Seawater samples for dissolved organic matter (DOM) and rare earth element (REE) analyses were taken during low and high tide at -0.5 m water depth by submerging 2 l polycarbonate bottles (Nalgene, acid-rinsed). Samples were processed as described in the subsequent paragraph. Filtered REE samples were filled into acid-cleaned 500 ml high-density polyethylen bottles and acidified with ultra-clean HCl to a pH of ~2.

2.2.5. Pore water sampling

Pore water sampling down to 2 m below sediment surface (mbss) was done using stainless steel push point samplers as described in Reckhardt et al. (2015). Sediment depths exceeding 2 mbss were sampled by elongating a stainless steel tip stepwise with 90 cm long steel extension rods (outer diameter 15 mm), which were pushed into the ground by percussion drilling (Charette and Allen, 2006). Pore water samples were collected through a 6 m long Teflon tube (inner diameter 5 mm) using a vacuum hand pump. Samples from the island's freshwater lens were obtained as described in Röper et al. (2012).

Aliquots for metal (Mn, Fe), total sulfur, silica and phosphate analyses via ICP-OES and samples for photometric nutrient analyses (NO\textsubscript{x}, NO\textsubscript{2}, NH\textsubscript{4}, total alkalinity) were treated as specified in Reckhardt et al. (2015). Aliquots for REE were filtered through AcroPak500 cartridges with 0.8/0.2 μm Supor filter membrane and acidified with ultrapure 6 M HCl to a pH of ~2. Metal and nutrient samples were stored dark and at 4 °C until analysis. For hydrogen sulfide analysis, 1 ml of sample was pipetted immediately into an Eppendorf vial, which contained 1 ml ZnCl\textsubscript{2} solution (2%). Salinity was determined directly in the field with a Multi 3430 (WTW) conductivity sensor. Samples for pore water (500 ml) and seawater (2 L) DOM analysis were immediately filtered through a sequence of pre-combusted (400 °C, 4 h) glass microfiber filters (GMF, 2 μm) and glass fiber filters (GF/F, 0.7 μm) using polyethylene syringes. Filtered samples were acidified to pH 2 with HCl (p.a.) and stored in polycarbonate bottles (acid-rinsed) at 4 °C in the dark. Samples for colored dissolved organic matter (CDOM) analysis in pore waters were
immediately filtered using a glass syringe and pre-combusted (400 °C, 4 h) glass fiber filters (GF/F, 0.7 μm) as pre-filter and 0.2 μm Nuclepore track-etched membrane filters. Sample bottles, acid-rinsed prior to use and then rinsed with filtered sample, were stored at 4 °C in the dark until analysis.

2.3. Numerical modeling

Numerical density-dependent flow and transport modeling was carried out for a two-dimensional vertical cross-section of the studied transect, and aimed to identify the principle flow patterns and residence times. The observed groundwater levels thereby served as calibration constraints. Modeling was carried out with the USGS groundwater modeling code SEAWAT (Langenhan et al., 2007). The horizontal extent of the model domain was 700 m, i.e., from the dunes to 400 m offshore (Fig. S1). The spatial discretization in horizontal direction ranged from 4 m near the southern and northern boundaries of the model to 1 m in the intertidal zone. The aquifer base was set to 44 m below mean sea level (mbsl), consistent with the presence of a more or less continuous clay layer at 44–55 mbsl that acts as an aquitard (Röper et al., 2012). The spatial discretization in vertical direction ranged from 2 m in the deeper parts of the aquifer to 0.25 m in the shallow subsurface. Fresh groundwater entered from the South with a flow rate of 0.5 m³ per day and meter of shoreline, estimated from the approximate distance to the groundwater divide (500 m inland from the southern model boundary) and an approximate recharge rate of 350 mm year⁻¹ (Röper et al., 2012). Seasonal variations in the groundwater recharge were not considered. Because the groundwater flow and solute transport time scales in the subterranean estuary are orders of magnitude larger than those of the tidal seawater level changes, flow and transport in the beach subsurface are mainly driven by the phase-averaged ideals of average over a tidal cycle (Vandenbohede and Lebbe, 2006). Therefore, when focusing on the principle groundwater flow and transport patterns, it is sufficient to calibrate the hydraulic model against the phase-averaged measured groundwater levels, rather than the propagation of the tidal wave into the aquifer. An idealized semi-diurnal tide with a 14-day neap-spring tidal cycle was simulated with the SEAWAT-periodic boundary condition package (Mulligan et al., 2011; Post, 2011). Storm floods and wave action were not accounted for.

During model calibration only the horizontal and vertical hydraulic conductivities, as well as the specific yield (unconfined storage coefficient), were adjusted. The transport parameters, i.e., effective porosity and longitudinal and transverse dispersivity were set fixed consistent with literature values for beach zones (e.g., Robinson et al., 2006). In agreement with the results of the grain size analysis (Fig. 2), we assumed homogeneity in the entire model domain. The simulation time was 700 days and ensured the simulation results being at a dynamic equilibrium and independent from the initial conditions. With a root-mean-square error of RMS = 0.11 m and a Nash-Sutcliffe-Parameter of NS = 0.95 we obtained a model sufficiently calibrated for the purpose of the study. However, the largest deviation was found for GWM3, since the de-saturation of a ridge located 100 m afar from the mean sea level mark during ebb tide was not captured well by the model. Details of model parameters and their estimated and applied values, flow and boundary conditions, and calibration results are given in the supporting information (Table S1, Figs. S1, S2).

2.4. Sample analyses

2.4.1. Grain size analysis

Sediments were sampled in regular intervals (9–12 cm) of each lithological unit, i.e., sediment layers defined by sharp boundaries and a specific particle composition, color, and grain size. Sediment particle sizes were analyzed with a HORIBA LA-950 laser diffraction particle size analyzer following the method applied in Reckhardt et al. (2015).

2.4.2. Quantitative and qualitative analysis of dissolved and particulate organic matter

Dissolved organic carbon (DOC) was quantified on triplicate subsamples of filtered and acidified water samples using high temperature catalytic oxidation on a Shimadzu TOC-VCPH instrument with an analytical accuracy and precision better than 5%. The stable carbon isotope composition was determined on solid phase extracted DOC (SPE-DOC) as outlined in Seidel et al. (2014).

Ultra-high resolution mass spectrometry was applied to characterize the molecular DOM composition. DOM extracts were diluted with ultrapure water and methanol (MS grade) to yield a DOC concentration of 20 mg C l⁻¹ and a methanol to water ratio of 1:1 (v/v) for analysis on a solarix FT-ICR-MS (Bruker Daltonik GmbH, Bremen, Germany) connected to a 15 T superconducting magnet (Bruker Biospin, Wissembourg, France). Instrument settings and raw data processing were described in Seidel et al. (2014). For this study, we considered peaks with a signal-to-noise (S/N) ratio of 2±4 and detection in >3 samples. Peak intensities were normalized to the sum of peak intensities in each sample.

Molecular DOM composition was statistically analyzed using the free software PAST (Hammer et al., 2001). Cluster analysis was based on Bray Curtis dissimilarity of presence and abundance distribution of individual detected masses in each sample.

Absorbance of CDOM was measured using quartz cuvettes with a path length of 0.05 m (at site 1 a 0.01 m cell was used if the sample volume was very small) and an UV-VIS-spectrophotometer (UV-2700, Shimadzu). Samples were scanned at slow scan speed with an increment of 0.5 nm in the spectral range between 200 and 800 nm. Ultrapure water was used as reference. Absorbance values A(λ) were baseline corrected and then converted to the absorption coefficient αCDOM(λ) according to

\[ \alpha_{CDOM}(\lambda) = 2.303 \times A(\lambda)/\lambda \]

where λ is the wavelength of the cuvette in meters (Kirk, 2011). Based on this spectral information the CDOM absorption coefficient at 275 nm and 440 nm as well as the spectral slope (SCDOM) in the spectral range of 275–295 nm were deduced (Garaba et al. 2014). Additionally the slope ratio (SP) defined by Helms et al. (2008) as the ratio

\[ \text{SP} = \frac{\text{SCDOM}(275–295 \text{ nm})}{\text{SCDOM}(350–400 \text{ nm})} \]

was derived. Precision and accuracy of the analyses were <5%.

Sediment samples were freeze-dried and homogenized in an agate mill. Total carbon was determined using a carbon analyzer (vario El cube, Elementar Analysensysteme GmbH), and the total inorganic carbon was analyzed coulometrically (UIC, Joliet, USA). Total organic carbon was calculated as the difference between total carbon and total inorganic carbon. The precision and accuracy of the analyses were <3%.

2.4.3. Oxygen concentrations and consumption rates

Oxygen (O₂) concentrations were measured using optode microsensors (Pyroscience, sensor type OXr 430 HS) that were introduced into the sediment cores immediately after retrieval by drilling holes into the aluminum liner. In addition, high resolution vertical O₂ profiles were measured at site 3 to resolve the shallow O₂ penetration depth. Furthermore, pore water was retrieved using a push point sampler at sites 2 and 4 and O₂ was measured using an optical O₂ flow sensor (Pyroscience, sensor type OXFTC).

O₂ consumption rates of surface sediments (upper 5 cm) from sites 3 and 2 were determined after Marchant et al. (2014) using flow through columns of 10 cm diameter and 20 cm length. The flow through columns were filled with sand while completely submerged in seawater to prevent the trapping of air bubbles. Using a peristaltic pump air saturated surface water was pumped through the flow through columns with flow rates adjusted to 8.2 and 9.1 ml min⁻¹. This corresponds to a pore water flow of 470–520 cm d⁻¹, which is comparable to the modeled flow velocity in the USP. O₂ concentrations were measured...
at in- and outlet using optical O2 flow through sensors (Pyroscience, sensor type OXFTC). O2 consumption rates were calculated by dividing the O2 difference between in- and outlet by the retention time, which was on average 1 h.

2.4.4. Analyses of dissolved nutrients, sulfate, hydrogen sulfide, and trace elements

Dissolved Fe, Mn, Si, and P were analyzed using inductively coupled plasma optical emission spectrometry (ICP-OES, Thermo Scientific). Samples with low Fe and Mn concentrations were additionally measured by inductively coupled plasma mass spectrometry (ICP-MS, Thermo Finnigan MAT). The SO4\(^{-2}\) concentration was determined by ion chromatography (IC, Metrohm) and sulfide was measured after Cline (1969). Nutrients (NH\(_4\), NO\(_3\), NO\(_2\), and total alkalinity) were analyzed as described in Reckhardt et al. (2015). Accuracy and precision was <10% for all measurements.

Rare earth element (REE) samples were spiked with a multi-element REE isotope spike (DKM, prepared by Gilbert N. Hanson of SUNY Stony Brook) and allowed to equilibrate for at least 48 h. A UV/H\(_2\)O\(_2\)-oxidation was applied for 24 h to destroy organic matter in the samples. The REEs were then preconcentrated from seawater using an automated seaFAST system (Elemental Scientific) following a method modified from Hathorne et al. (2012). The REEs were analyzed and quantified by isotope dilution ICP-MS using a Thermo Element 2 and a Cetac Aridus II system (Elemental Scientific). REEs were spiked with a REE isotope spike (DKM, prepared by Gilbert N. Hanson of SUNY Stony Brook) and allowed to equilibrate for at least 48 h. A UV/H\(_2\)O\(_2\)-oxidation was applied for 24 h to destroy organic matter in the samples. The REEs were then preconcentrated from seawater using an automated seaFAST system (Elemental Scientific) following a method modified from Hathorne et al. (2012). The REEs were analyzed and quantified by isotope dilution ICP-MS using a Thermo Element 2 and a Cetac Aridus II system (Elemental Scientific). After German et al. (1995) using PAAS-normalized values (Post-Archaean Australian Shale, Taylor and McLennan, 1985).

To present salinity, DOC, nutrient, SO\(_4\)\(^{-2}\), and trace element data, concentrations below the limit of quantification were set to zero, which is appropriate considering the wide concentration range we present here for each element. On the basis of these data sets contour plots were created using the interpolation mode v4 (biharmonic spline interpolation) of MATLAB R2013b (The MathWorks Inc., Natick, MA). Additionally, the concentration at each pore water extraction location was plotted color-coded onto the interpolated area to emphasize the exact value.

2.4.5. Particulate nutrient and chlorophyll a analyses

For analyzing particulate nutrient and chlorophyll a concentrations of the benthic microalgae community, microalgae were separated from the sediment. Microscopic investigations ensured that the samples contained mostly benthic algae and not phytoplankton, which might have settled on the sediment. Seawater (0.2 μm filtered) was added to the sediment samples and these samples were stirred and placed in an ultrasonic bath (20 min). Afterwards the overlaying suspension was decanted. This was done three times and the overall suspension of each sample was filtered onto pre-combusted, acid-washed glass fiber filters (Whatman GF/F). These filters were stored at −20 °C until further analysis.

Particulate organic phosphorus (POP) was measured after muffle firing the filters at 550 °C for 24 h by molybdate reaction after sulfuric acid digestion (Wetzel and Likens, 2003). Particulate organic carbon (POC) was measured using a CN elemental analyzer (Thermo Flash EA 1112). Chlorophyll a (Chl a) concentrations were measured photometrically after extraction with acetone (Wetzel and Likens, 2003). To be able to compare concentrations of POP, POC, and Chl a among transects and stations we normalized these concentrations, based on the mean value of each parameter for each transect, and log transformed them.

2.4.6. Total bacterial cell counts and microbial community analysis

For total cell counts, 0.5 cm\(^3\) sediment was fixed with 2% formaldehyde overnight. The sediment was centrifuged and washed three times with 1× phosphate-buffered saline (PBS; 145 mM NaCl, 1.4 mM NaH\(_2\)PO\(_4\), 8 mM Na\(_2\)HPO\(_4\), pH 7.4). Samples were stored in PBS/ethanol (v/v, 2:3) at −20 °C. For SybrGreen staining, the samples were sonicated three times each for 1 min. After settling of the sand particles, the supernatant containing the cells was transferred to another vial. In this way, the sediment was washed six times with PBS and the supernatants combined. Cells from the supernatant were filtered onto 0.1 μm Nuclepore polycarbonate filters and stained with SybrGreen solution directly on the object slide. For each sample, at least 400 cells were counted by epifluorescence microscopy.

DNA was extracted from 0.5 cm\(^{-3}\) frozen sediment with PowerSoil DNA Isolation Kit (MoBio) according to the manufacturer’s instructions. Polymerase chain reaction (PCR) for denaturing gradient gel electrophoresis (DGGE) was performed with 31–41 cycles (depending on the final DNA concentration of the PCR products) with primers 357F with GC clamp and 907R (Muyzer et al., 1995) for bacterial 16S rRNA genes, primers Arch-amoA-F and Arch-amoA-R (Francis et al., 2005) for archaeal amoA genes, primers amoA-1F and amoA-2R (Rotthauwe et al., 1997) for betaproteobacterial amoA genes, primers aprA-1-FW-GC and aprA-5-RV with GC clamp (Meyer and Kuever, 2007) for aprA genes. For archaeal 16S rRNA genes, a nested PCR was performed according to Nicol et al. (2003). For the first PCR primers Ar3F and Ar9R were used with 25 cycles, PCR products were purified with QiQuick PCR purification kit (Qiagen) and used for the second PCR with primers S4F and PARCH519r and 37–44 cycles. The specificity of the primer sets was tested by excision of single DGGE bands, rearrangement and sequence analysis.

DGGE was performed with PCR products loaded onto 6% polyacrylamide gels in 1× TAE buffer (40 mmol l\(^{-1}\)) with a denaturing gradient from 50 to 70% (with 100% denaturant corresponding to 7 M urea and 40% formamide) at 100 V and 60 °C for 20 h for bacterial 16S rRNA genes, 15–55% for 15 h for archaeal amoA genes, 30–60% for 20 h for betaproteobacterial amoA genes, and 40–70% for 20 h for aprA genes. For PCR products of archaeal 16S rRNA genes, the gel consisted of 8% polyacrylamide with a denaturing gradient from 35 to 80% and ran for 20 h. Gels were stained in 1× SybrGold solution in 1× TAE buffer and washed for 30 min in distilled water. The band patterns were analyzed by cluster analysis using the software package GelComparII (Applied Maths) as described by Wilms et al. (2006).

3. Results

3.1. Sedimentology

Beach sediments consist of quartzitic fine to medium grained sand (mean grain sizes 170–320 μm; Fig. 2). Shells or shell debris are found in all cores and they occur as distinct layers or are finely dispersed. Stratification is best visible where layers of coarser-grained sand, heavy minerals, or organic-rich mud are intercalated in the fine sand. The mud pebbles originate from erosion of subfossil semi-consolidated mud beds (Hellwig and Stock, 2014). Internal sedimentary structures are restricted to even lamination.

3.2. Modeling of groundwater flow and residence times

The model results (Fig. 3) reflect the typical hydrological zonation of beaches with an upper saline plume (USP), a terrestrial groundwater flow and freshwater discharge tube, and a saltwater wedge as proposed by Robinson et al. (2007a, 2007b). At our study site, the modeled USP reaches a depth of about 20 m. The subsurface residence times in the USP are in the order of weeks to months, depending on the depth of the respective flow-paths (Fig. 3), and the phase-averaged pore water velocities at the entry points reach approximately 0.3–0.7 m d\(^{-1}\). In contrast, the residence times in the saline wedge are in the order of...
years to decades. The residence time of the fresh groundwater from the dune base to the discharge point is approximately 6–13 years, whereby the last section of the flow path, i.e., the freshwater discharge tube is passed within less than a year. Fresh groundwater originating from the island’s freshwater lens displays a typical age stratification, with a maximum groundwater age of ~40 years in the deepest part of the freshwater lens according to tritium-helium dating (Röper et al., 2012). Below the beach, this old freshwater and the much younger
freshwater from shallower parts become well mixed in the freshwater discharge tube, where they also mix with the very young saltwater of the USP and the older water from the lower salt water wedge.

The position of the freshwater discharge tube requires further investigations because the model proposes its position at the beach runnel, whereas less saline water was determined by pore water analysis approximately 50 m seaward from the model-predicted discharge point (Figs. 3, 4). The deviation between model and measurements may be attributed to the simplification of the seaward and landward flow boundary conditions in the model, i.e., neglect of seasonal groundwater recharge and storm surges. It is also possible that the freshwater discharge point varied with time due to changes in beach morphology on longer time scales, leaving residual brackish water at the location of the earlier freshwater discharge point.

A groundwater flow divide exists at the ridge of the beach transect. This leads to varying residence times of <6 years and up to 25 years on the landward and seaward sides of the divide, respectively. The simulated residence time distribution below the ridge has to be regarded as rough estimate due to the overestimation of the phase-averaged hydraulic head at GW3 (Fig. 3).

The analysis of the model’s water budget reveals that the saltwater flux through the USP is approximately 2.8 m³ per day and meter of shoreline. From the ridge the total saltwater flux of 1.5 m³ per day and meter of shoreline is diverted to the runnel and to the seaside, with 0.5 and 1.0 m³ per day and meter of shoreline, respectively. The total fresh water discharge is 0.75 m³ per day and meter of shoreline. The overall error in the model-based estimation of SGD fluxes as a result of model assumptions, i.e., estimation of fresh groundwater flux, neglecting waves and storm floods, as well as the calibration performance, can be considered as being roughly 0.5 to 2 times the derived values. This error estimation is based on findings by Xin et al. (2010) who found in their modeling analysis that the total SGD rate increased by 33% when waves and tides were considered compared to tides alone. Furthermore, the saline SGD rate through the upper saline plume doubled.

3.3. Pore water chemistry

3.3.1. Salinity and dissolved organic matter

Salinities are <10 in vicinity of the dunes and reach 31 in the intertidal zone (Fig. 4). At site 3, salinity decreases distinctly to about 15 below 1 m sediment depth. Dissolved organic carbon (DOC) concentrations range between 80 and 175 μM and are highest close to the mean low water level (Fig. 4). CDOM absorption at 275 nm ranges between 4.2 m⁻¹ and 9.5 m⁻¹ and is related to DOC concentrations and salinity (Fig. 5), while absorption at 440 nm stays nearly constant (0.6 ± 0.4 m⁻¹) for all samples. The slope of the CDOM absorption spectra at 275 to 295 nm ranges from 0.011 nm⁻¹ at low DOC and salinity levels to 0.019 nm⁻¹ for higher DOC concentrations and salinities (Fig. 5). The slope ratio S₀ is below 1.0 for salinities <17 and above 1.0 for higher salinities.

3.3.2. Oxygen saturation, consumption, and flux

At site 1, pore water between 1 and 2.2 m depth (sediments were only water saturated below 1 m depth) is O₂ saturated showing no trend with depth (Fig. 4). At site 2, O₂ saturation decreases from 60 to 80% at 0.6 m to 2% at 2 m depth. At site 3, O₂ saturation is 60–100% in surface sediments down to 4.5 cm. Further below, it decreases rapidly to below 5% at 13 cm depth. At site 4, O₂ is below 10% at 10 cm and 0.7% at 50 cm depth. O₂ respiration experiments using flow through columns show that O₂ concentrations at the outlet of the columns are in steady state after 1–2 h of pumping. The O₂ differences between inlet and outlet are 44 μM (site 2) and 133 μM (site 3) and from the pore water retention time of 1 h, potential O₂ consumption rates of 44 μM h⁻¹ and 133 μM h⁻¹ can be calculated. Assuming steady state conditions, the rate can be integrated over the O₂ penetration depth to estimate the O₂ flux. The resulting O₂ fluxes are 9.3 mmol m⁻² h⁻¹ at site 3 and 84 mmol m⁻² h⁻¹ at site 2. However, for the latter site the calculated flux likely overestimates the true flux as we assumed that the O₂ consumption rates measured in the surface sediments (0–5 cm) were the same for the entireoxic sediment layer from 0 to 1.9 m.

3.3.3. Inorganic nutrients

Dissolved inorganic nitrogen consists mainly of NO₃⁻ near the dune and concentrations decrease with increasing salinity towards the low-water line, where inorganic nitrogen is comprised mostly of NH₄⁺ (Fig. 4). At site 2, NO₂⁻ concentrations are highest (Fig. 4). PO₄³⁻, Si(OH)₄, and total alkalinity concentrations increase from the duneward part to the low-water line (Fig. 4). Additionally, Si(OH)₄ concentrations increase significantly at site 3, coincidently with decreasing salinity.

3.3.4. Redox-sensitive trace metals and sulfate

Dissolved Mn and Fe concentrations increase from the backshore towards the low-water line and attain maximum concentrations at the low-water line (Fig. 4). The spatial SO₄²⁻ distribution resembles the salinity distribution (Fig. 4) and sulfide was below the limit of quantification (<2 μM) in all samples (data not shown).
Besides Mn, Fe, and sulfate, rare earth elements can be used to classify the redox regime of pore waters. Ce is oxidized to insoluble Ce(IV) and removed from the dissolved phase under oxic conditions, leading to a depletion relative to its neighbor elements and hence to negative Ce-anomalies (e.g., Elderfield, 1988; Haley et al., 2004). Nd is scavenged by organic and inorganic particles from oxic water due to its high particle reactivity (e.g., Elderfield, 1988). Under suboxic/anoxic conditions, Ce oxides dissolve and Nd is released from particles leading to positive or no Ce-anomalies and elevated Nd concentrations. Along the beach transect, the Ce-anomaly shows a separation between sites 0/1 and sites 3/4 (Fig. 4). In the backshore (sites 0/1), negative Ce-anomalies (<1) are found, whereas at sites 3/4, Ce-anomalies are absent or slightly positive (>1). A similar spatial distribution is apparent for Nd with concentrations almost one order of magnitude higher at sites 3/4 than at the other sampling locations (Fig. 4).

3.4. Benthic chlorophyll a and particulate nutrients

Chlorophyll a, particulate organic phosphorus (POP), and particulate organic carbon (POC) concentrations of phytobenthos (Fig. 6 upper panel) tend to increase from the mean high water level towards the site located in the runnel (phytobenthos site 6) draining the emerged parts of the beach during falling tide and decrease again at sites located closer to the mean low water level. The normalized POP, POC and Chl a concentrations differ significantly between sites (ANOVA, \( p < 0.001 \)). Especially the site located in the middle of the runnel (phytobenthos site 6) is significantly different compared to sites closer to the high and low water lines (TukeyHSD PostHoc Test, \( p < 0.01 \); Fig. 6 lower panel).

3.5. Cell counts and microbial diversity

Total bacterial cell counts range between \( 1.4 \times 10^7 \) and \( 2.6 \times 10^8 \) cells cm\(^{-3}\) sediment. Generally, they are lower at sites 0 and 1 than at the other three sites most likely because of the low water content in these sediments. Molecular analysis by using denaturing gradient gel electrophoresis (DGGE) analysis (Figs. 7, S3) reveals distinct vertical and horizontal compositions of microbial communities in the beach subsurface. This molecular technique is based on the sequence-specific separation of gene fragments amplified from an environmental sample. The resulting banding patterns indicate the presence or absence of a certain species that carry the investigated gene. Similarities and differences between the microbial communities can be visualized by cluster analysis of the DGGE banding patterns. Over all samples, the diversity of bacterial 16S rRNA genes is generally higher compared to the archaeal, indicating a more complex bacterial community composition on the species level. Besides analyzing this general phylogenetic marker, we additionally studied key genes indicative for metabolic processes to
obtain a more detailed view on variations of selected physiologic traits within the microbial communities. The ammonia monoxygenase (amoA) is involved in the aerobic oxidation of ammonia. Ammonia-oxidizing betaproteobacteria and archaea can specifically be distinguished. The dissimilatory adenosine-5′-phosphosulfate reductase (aprA) is present in SO₄²⁻-reducing and sulfur-oxidizing prokaryotes. Interestingly, independent on the genes analyzed, DGGE banding patterns of sites 0 and 1 as well as 3 and 4 cluster together, indicating unique microbial community compositions in the duneward and seaward parts of the transect (Fig. 7). All depths of site 3 and 4 with all genes analyzed form a coherent cluster. In contrast, samples of all depths of site 2 either cluster with the landward or seaward sites.

4. Discussion

4.1. Beach hydrology and salinity gradients influence the beach ecosystem

The groundwater flow and transport field (Fig. 3) reveals the typical hydrological zonation of beaches with an USP, the saltwater wedge, and the freshwater discharge tube in between. This zonation is also reflected in the spatial salinity distribution (Fig. 4). In general, such a groundwater flow field is due to nearshore flow processes such as density-driven circulation, tide-induced circulation, wave-setup induced circulation, local circulation due to wave-bed form interaction, and fresh groundwater discharge (Precht and Huettel, 2003; Robinson et al., 2007b).

In response to tidal forcing and beach morphology, the model predicts that the USP reaches a depth of 15–20 m below the beach surface (Fig. 3). Thus, the USP at Spiekeroog beach may extend down to greater depths than previously shown in other studies using salinity measurements and/or hydrological modeling (Charette and Sholkovitz, 2006; Heiss et al., 2014; McAllister et al., 2015; Santos et al., 2008; Ullman et al., 2003). The majority of these studies were, however, carried out in microtidal environments, which may explain why the USP does not reach depths exceeding 5 m.

4.2. Spatial differences in organic matter sources and turnover result from pore water circulation patterns

The stock of particulate organic carbon in the permeable beach sands is very low (<0.03%). This indicates that there must be an additional source of organic carbon fueling the microbial activity in the sediments, which is likely the infiltration of dissolved and particulate organic matter (OM) by pore water circulation. Residence times are short in the USP, less than eight months or even shorter in the zone at the mean high water level that is saturated and desaturated at tidal rhythm (Fig. 3). This leads to a continuous resupply of the sediments with OM as indicated by elevated DOC concentrations at site 2 (Fig. 4). The zone of elevated DOC concentrations is, however, relatively narrow. This may be due to a fast recirculation of remineralization products to the water column (Chipman et al., 2010; Ehrenhauss et al., 2004; Seidel et al., 2015). Furthermore, sampling took place several weeks after the main spring phytoplankton bloom, which started at the beginning of March, as reflected in sharply decreasing silicate concentrations at a nearby permanently installed time series station (similar measurements are described in Grunwald et al. (2010) and Beck and Brumsack (2012)). Consequently, the OM of the spring phytoplankton bloom had likely already been processed in the upper sediments of the USP. In contrast to the USP, high DOC concentrations at sites 3 and 4 are not only due to continuous OM input, but also due to longer pore water residence times in the order of some years in the ridge sediments (Figs. 3, 4), which allow the enrichment of OM remineralization products like DOC and alkalinity.

To determine whether DOM is of marine or terrestrial origin, carbon isotopic compositions of SPE-DOC were analyzed. They fall in a narrow range of −24.1 to −25.6‰, indicating a predominantly marine source of the DOM (seawater endmember −24.3‰ (n = 4, σ = 0.07), freshwater endmember determined in the freshwater lens of Spiekeroog Island on average −27.4‰ (n = 14, σ = 0.8)). The marine origin of the DOM is further supported by the finding that the molecular composition of pore water DOM is more similar to seawater DOM than to DOM found...
that the DOM composition at a nearby beach site is in et al., 2008; Riedel et al., 2013). For example, Seidel et al. (2015) showed sites, which might be a consequence of changing redox conditions (Jaffé et al., 2008), except for the sample of lowest salinity at site 3. Furthermore, the molecular DOM composition of waters from the freshwater lens (FL) is different from the molecular composition of pore water DOM (sites 0–4). Pore water DOM is more similar to North Sea water (NSW) DOM. North Sea water was sampled close to the molecular masses detected in DOM from pore water, seawater and the island’s freshwater lens. Freshwater lens samples were obtained at seven locations labeled 25–31, as depicted in Röper et al. (2012), from shallow (A) and deep (B) observation wells. The molecular DOM composition of waters from the freshwater lens (FL) is different from the molecular composition of pore water DOM (sites 0–4). Pore water DOM is more similar to North Sea water (NSW) DOM. North Sea water was sampled close to the beach (north, low water (LW), mean water (MW), and high water (HW)), in the beach runnel, and in the harbor of Spiekeroog Island. The molecular composition of pore water DOM extracted at sites 0 and 1 cluster together as well as that at sites 3 and 4, indicating specific DOM compositions. The color scheme of the pore water sampling sites is the same as in Fig. 7.

4.3. Pore water origin and redox state control nutrient and metal cycling

Redox conditions can be categorized into oxic diagenesis (O₂ reduction), suboxic diagenesis (NO₃⁻, Mn and Fe reduction) and anoxic diagenesis (SO₄²⁻ reduction and methanogenesis) (Froelich et al., 1979). Along the studied beach transect, the redox gradient ranges from completely oxic conditions down to deep sediment layers at the landward sites (sites 0 and 1) to oxic only at the surface and anoxic deeper in the sediments at the three seaward sites (Fig. 4).

4.3.1. Oxic conditions control biogeochemistry at sites 0 and 1

Sites 0 and 1, which are only irregularly flooded by seawater, e.g., during storm surges or exceptional high tide levels, exhibit oxic conditions as reflected by high NOₓ concentrations and O₂ saturation (Fig. 4, for O₂ concentrations in μM see Table S2 in the supporting information). Lowest measured Ce (negative Ce-anomalies) and Nd concentrations (Fig. 4) further support the oxic character at these sites, as they indicate Ce removal upon its oxidation to insoluble Ce(IV) and Nd adsorption onto particles (Elderfield, 1988).

The high NOₓ concentrations up to 260 μM do not originate from fertilizers or sewage transported to the beach via groundwater flow because the study site is located in an environmentally protected area, where agriculture and fossil fuel traffic are absent. This is in contrast to several other subterranean estuaries where fertilizers and sewage represent an important nitrogen source (Andersen et al., 2007; Kroeger and Charette, 2008; Wong et al., 2014). Instead, the high NOₓ concentrations may originate either from seawater or rainwater, or from biogeochemical reactions occurring on site. Seawater infiltration can, however, not be regarded as major inorganic N source because seawater concentrations are distinctly lower than those found at sites 0 and 1 (9 μM in May 2014). Instead, rainwater is likely an important source because the average N concentration in rainwater of neighboring islands (Langeoog and Norderney) from monthly measurements over a two year period amounts to 155 μM (85 μM NH₄, 70 μM NO₃) (Armbärtser, 2002). Due to highly permeable dune/beach sediments and only scarce dune vegetation, high N amounts can reach deeper sediment layers. Assuming that pore water at site 0 is simply a mix of rainwater and seawater, 43–67% of the total inorganic N measured in the pore water can be explained by conservative mixing. Even with a maximum seasonal seawater inorganic N concentration of 50 μM (Reckhardt et al., 2015), this proportion only amounts to 47–75%. Consequently, 25–57% (seasonal differences in seawater inorganic N are considered by giving this range) of the inorganic N is produced on-site, for example by organic matter degradation. High O₂ availability favors NH₄ oxidation to NOₓ, which accumulates in the pore water. This conclusion is further supported by the finding that the potential for ammonia oxidation (the first step of nitrification) is shown by DGGE analysis of genes for ammonia oxidation (amoA) of both betaproteobacteria and thaumarchaeota. Both groups are present at all sites and depths and especially the ammonia-oxidizing archaea exhibit the highest diversity in the deeper sediment zones of sites 0, 1, and 2.

4.3.2. Changing redox conditions control biogeochemistry at site 2

At or close to site 2, the redox conditions change from oxic to suboxic in deeper sediment horizons. O₂ saturation decreases with depth and reaches zero at about 2 m. Rate measurements using surface sediments from 0 to 10 cm depth show high potential O₂ consumption rates of 44 μM h⁻¹ indicating that processes such as organic matter turnover and nitrification occur rapidly in the upper layer. Integrating the volumetric rate down to the O₂ penetration depth results in an O₂ flux of
84 mmol m$^{-2}$ h$^{-1}$, which is even higher than the rate determined for comparable shallow subtidal or intertidal sands (Rusch et al., 2006; Werner et al., 2006). The presence of NO$_3^-$ at site 2 is also indicative of changing redox conditions because it is either released when NH$_4^+$ is oxidized via nitrification when encountering aerobic conditions (Kowalchuk et al., 1997) or when NO$_3^-$ is transformed to N$_2$ via denitrification under anaerobic conditions (Evard et al., 2013; Slomp and Van Cappellen, 2004).

### 4.3.3. Anoxic conditions control biogeochemistry at sites 3 and 4 below an oxic line of some centimeters depth

The oxic line stretches only across a few centimeters at sites 3/4 and potential O$_2$ consumption rates are three times higher compared to site 2. This suggests that biogeochemical processes such as OM degradation and NH$_4^+$, Fe or sulfide oxidation occur rapidly in this thin, O$_2$-containing zone. Below the surface sediments, suboxic to anoxic conditions dominate as indicated by increasing concentrations of dissolved Mn/Fe, probably due to on-site reduction of their particulate phases. The intrusion of fresh/brackish waters from below is an unlikely Mn and Fe source because of their low concentrations in the island’s freshwater lens (<3 and 16 μM, respectively, Reckhardt et al., 2015). The classification of the deeper pore waters of sites 3/4 as mainly anoxic is supported by the absence of a negative Ce-anomaly and the increased Nd concentrations (Fig. 4).

Although anaerobic OM degradation via SO$_4^{2-}$ reduction is not directly shown by our SO$_4^{2-}$ and sulfide measurements, the grayish color of the sediments indicates SO$_4^{2-}$ reduction and sulfide formation leading to the formation of Fe sulfides. Furthermore, changes in the O isotope composition of the pore water SO$_4^{2-}$ indicate that microbial conversion of SO$_4^{2-}$ must have occurred and sulfide reoxidation is likely an important part of sulfur cycling (Reckhardt et al., 2017). The relative importance of Mn, Fe, and SO$_4^{2-}$ reduction can, however, not be estimated by our data set because of missing rate measurements. Snyder et al. (2004) showed, for example, that Fe reduction accounts for most of OM degradation at the salinity transitions of an island freshwater lens of Sapelo Island, Georgia, with SO$_4^{2-}$ reduction occurring simultaneously. They concluded that metal reducing bacteria can co-occur with SO$_4^{2-}$ reducing bacteria even at high SO$_4^{2-}$ concentrations if O$_2$ is episodically re-introduced and a dynamic cycle between dissolved Fe and Fe oxides is maintained.

### 4.3.4. Summary biogeochemical zones

Different biogeochemical zones can be identified in the Spiekeroog beach system (Fig. 9). At sites 0 and 1, oxic conditions and aerobic organic matter degradation dominate. Site 2, which is located close to the mean high water level (MHWL), exhibits oxic to suboxic conditions and the biogeochemical zones aerobic degradation, nitrification, and denitrification can be identified. In contrast, at sites 3 and 4 the oxic sediment layer stretches only some centimeters, whereas below indications of anoxic conditions and reduction of particulate Mn/Fe and SO$_4^{2-}$ are found.

The biogeochemical zones, in turn, are related to OM and nutrient availability as well as pore water residence time. For example, the freshwater diluting the seawater close to the dunes is older (up to 2 years) than the saline water circulating through the USP (some months), where the recharge is higher (Fig. 3). Consequently, residence time does not serve as an explanation for oxic redox conditions in brackish water near the dunes and decreasing oxygen concentrations within the USP (Fig. 4). Here, OM availability may determine redox conditions, either by the amount being introduced or by the lability of the available OM. At our study site, the mainly marine OM origin suggests that the OM amount is of higher importance for controlling redox conditions than the lability. Close to the low-water line, however, older pore water from greater depth reaches shallow sediments (Fig. 3). Consequently, DOC accumulation and Mn/Fe/SO$_4^{2-}$ reduction conditions may be mainly a function of residence time. Similarly, the importance of redox conditions and pore water residence time for controlling oxygen and nutrient concentrations were identified previously for other systems affected by submarine groundwater discharge (Charbonnier et al., 2016; Santos et al., 2008; Slomp and Van Cappellen, 2004).

Previous studies further showed that saltwater-freshwater interfaces may be reaction hotspots (Charette and Sholkovitz, 2002; Roy et al., 2010; Spiteri et al., 2008a). This is not supported by our data. Redox conditions are oxic along the saltwater-freshwater transition zone. Decreases in pore water nutrient concentrations (NO$_3^-$, alkalinity) are only due to dilution with seawater of lower nutrient concentrations.

### 4.4. Biogeochemical zones influence microbial community patterns

In our study area, salinity and redox conditions as well as DOM availability vary considerably (Figs. 4, 8). The diversity of bacteria and archaea, analyzed as 16S rRNA genes and functional genes for aerobic
ammonia oxidation, SO\(_4^{2-}\) reduction and sulfide oxidation, is apparently influenced by these variations in environmental settings (Figs. 6, S3). The clustering of landward and seaward sites, independent on the genes analyzed, indicates that the microbial community was structured by the presence/absence of O\(_2\) and the salinity. At the seaward sites, redox conditions may be the main driving force for changing diversity, where the dominant metabolic pathways (Mn, Fe, SO\(_4^{2-}\) reduction) vary with depths and between sites. Additionally, the non-specific clustering of the different sediment layers of site 2 may indicate that environmental conditions fluctuate with time. We suggest that changes in the redox environment, which can occur at this site due to short pore water residence times and saturation/desaturation cycles in surface sediments (Jansen et al., 2009; Wilson and Gardner, 2006), are the main driver for the microbial community structures.

Interestingly, the same clustering as of microbial communities is found for the molecular DOM composition, with site 0 and 1 as well as 3 and 4 clustering together while site 2 is in between (Figs. 7, 8). From our data, we cannot infer any causal relationships of DOM and microbial community composition. It is possible that different microbial communities influence the DOM composition in a different way, and vice versa. However, it is more likely that the observed co-variance of DOM and microbial communities reflects prevailing environmental conditions, e.g., salinity and/or redox state.

4.5. Pore water discharge influences phytobenthos and nearshore waters

According to the hydrological model, the seawater flux through the USP is on average 2.8 m\(^3\) per day and meter of shoreline. This seawater flux is in the same order of magnitude as the fluxes determined by Hays and Ullman (2007) and Mulligan and Charette (2006) for sandy microtidal settings. In contrast, Anschutz et al. (2009) report a groundwater flux about one order of magnitude higher for a macrotidal sandy environment. At our study site, the ridge sediments further discharge 0.5 and 1.0 m\(^3\) per day and meter of shoreline to the runnel and the seaside, respectively.

We assume that the pore water concentrations measured at the shallowest sampling depth (0.5 mbss) of the site located 50 m seaward of site 2 are representative for the discharge region of the USP. To calculate the nutrient flux from the ridge towards the runnel, we used concentrations of site 3 (0.5 mbss). For discharge from the ridge towards the seaside, we considered the average concentrations of site 4 and the site located some meters seaward (0.5 mbss) were considered. Using these concentrations (Fig. 4) and seawater endmember concentrations of 0.4 μM NH\(_4^+\), 0.3 μM PO\(_4^{3-}\), and 3 μM Si(OH)\(_4\), as measured in May 2014, calculated net nutrient fluxes of saline beach pore water to adjacent seawater per day and meter of shore line are 103 mmol NH\(_4^+\), 49 mmol PO\(_4^{3-}\), and 304 mmol Si(OH)\(_4\).

In contrast to the seawater flux through USP and ridge sediments, the freshwater flux is lower, with 0.75 m\(^3\) per day and meter of shoreline. We used average nutrient concentrations of 14 samples of the freshwater lens from depths between 6 and 43 mbsl to determine freshwater endmembers (19 μM NH\(_4^+\), 7.4 μM PO\(_4^{3-}\), and 361 μM Si(OH)\(_4\)). Including freshwater and seawater flux, total nutrient discharge per day and meter shore line amounts to 117 mmol NH\(_4^+\), 55 mmol PO\(_4^{3-}\), and 575 mmol Si(OH)\(_4\). We used NH\(_4^+\), not NO\(_3^-\), for these calculations, because NH\(_4^+\) is the N species present in the sub-anoxic discharge region of the beach and in the freshwater lens samples.

Nutrient efflux of beach sediments can significantly sustain productive benthic communities that inhabit the seepage zone (Hays and Ullman, 2007; Ullman et al., 2003). In our study area, increasing normalized Chl a, POC, and POP concentrations in the seepage zone, especially the beach runnel (Phytobenthos site 6, Fig. 6), indicate a stimulation of the phytobenthos growth by groundwater controlled nutrient effluxes.

4.6. Nutrient discharge from tidal flat margins is more important than efflux from barrier island beaches for nearshore North Sea waters

Overall, this first SGD and nutrient export estimation for the Spiekeroog beach system indicates that groundwater discharge is an important pathway to supply nutrients to nearshore waters of the southern North Sea. However, the present study so far does not support the hypothesis that beach systems may be of greater importance for nutrient recycling and efflux than sandy tidal flat sediments. The latter were shown to be of great importance for carbon, nutrient, and trace metal cycles in the large tidal flat area called Wadden Sea stretching from the Netherlands to Denmark, and the frequent water exchange between tidal flat areas and the North Sea transport carbon, nutrient, and trace metals to coastal waters of the North Sea (e.g. Beck et al., 2008; Billerbeck et al., 2006; Grunwald et al., 2010; Marchant et al., 2014; Moore et al., 2011; Riedel et al., 2011; Roy et al., 2008; Santos et al., 2015). Pore water discharge rates calculated for nearby sandy Wadden Sea sediments vary depending on the method applied and the sediment depth considered (Billerbeck et al., 2006; Moore et al., 2011; Riedel et al., 2010), among which the method most comparable to the model approach applied in our study resulted in lower flux rates (0.97 m\(^3\) per tide and meter of shoreline, which equals ~2 m\(^3\) per day and meter of shoreline; Riedel et al., 2010) compared to our study. Due to the high nutrient enrichment in the tidal flat pore waters compared to Spiekeroog beach sediments (Reckhardt et al., 2015), the nutrient efflux from these sediments can, however, be higher compared to the beach site. For example, the NH\(_4^+\) efflux was estimated to reach up to 430 mmol per tide and meter of shoreline (Riedel et al., 2011), which is about seven times higher than the sum of all NH\(_4^+\) discharge pathways from the beach sediments.

To estimate the importance of beaches in terms of nutrient contribution to the North Sea, the nutrient efflux determined at Spiekeroog beach was extrapolated to the entire East-Frisian barrier island chain from Borkum to Wangerooge (without Kachelotplate and Memmert). This estimation is based on the assumption that the nutrient efflux determined in our study is representative for the beach zones of the barrier islands, which are facing the open North Sea and are in contact with a freshwater lens. This zone was estimated to be 56 km long. Based on nutrient discharge rates described previously, the total fluxes (including seawater and freshwater fluxes) from beaches to adjacent nearshore waters are 24 × 10\(^5\), 11 × 10\(^5\), and 118 × 10\(^5\) mol a\(^-1\) for NH\(_4^+\), PO\(_4^{3-}\), and Si(OH)\(_4\), respectively. Similarly, the nutrient export of the Spiekeroog backbarrier tidal flats was extrapolated to the entire Wadden Sea (from the Netherlands to Denmark) resulting in an export of 70 × 10\(^5\) and 30 × 10\(^5\) mol a\(^-1\) for PO\(_4^{3-}\), and Si(OH)\(_4\), respectively, from the backbarrier areas to the southern North Sea (Grunwald et al., 2010). Although the extrapolations do not consider the same areas (East-Frisian versus total Wadden Sea), a first comparison shows that the beach zone seems to export less (~10%) phosphate and silicate than the backbarrier tidal flat areas.

5. Perspectives

The studies carried out in the beach system of Spiekeroog Island showed that in mesotidal settings sediments are continuously flushed by sea water down to several meters depth in the zone of the USP (Fig. 3). Furthermore, sea water may circulate through ridge sediments. Redox conditions and pore water residence time control biogeochemical zones (Figs. 4, 9). Furthermore, biogeochemistry is influenced by OM availability, especially in the freshwater-dominated part of the beach, which is only irregularly flooded by sea water and fresh OM input is limited.

In summary, the beach consists of three zones: 1) the freshwater-dominated, oxic zone, which is located behind a backshore berm and only flooded irregularly, and which exhibits limited OM availability, 2) the seawater-influenced, oxic to suboxic USP zone where pore water...
circulation may lead to regular oxygen, nutrient, and OM availability changes, and 3) the seawater-influenced, mainly anoxic zone in the ridge sediments. This zonation influences the microbial community patterns as reflected in the clustering of landward and seaward sites (Fig. 7).

Pore water discharge of Spiekeroog beach sediments supplies nutrients to nearshore waters. Nutrient fluxes of saline pore water are the major N and P source to nearshore waters, whereas fresh and saline pore waters transport equal amounts of Si. The importance of Spiekeroog beach sediments in supplying nutrients to nearshore waters is, however, reduced due to its special geographical location next to the large tidal flat area called Wadden Sea. In this area, similar pore water discharge rates, but much higher nutrient concentrations lead to higher nutrient effluxes compared to the beach sediments.

An essential next step while studying the beach system of Spiekeroog Island, but also other systems worldwide, will be to extend the knowledge to greater sediment depths, at least up to where the freshwater discharge tube is intersected and the underlying saltwater wedge is met. This will lead to a refined groundwater flow model, to a better understanding of the key processes in biogeochemistry, of the role of organisms for OM degradation, and of the fluxes at the sediment-water interface. To date, most studies on hydrological and biogeochemical processes in subtidal estuaries of beach systems were carried out in microtidal and/or low-energy environments such as semi-protected bay situations (Charette and Sholkovitz, 2006; Heiss and Michael, 2014; McAllister et al., 2015; Santos et al., 2008; Ullman et al., 2003). Therefore, a maximum sampling depth of about 5 m was sufficient to resolve the USP at these study sites. A mesotidal beach system was, for example, sampled by Robinson et al. (2006). Their model results show an USP stretching to about 8 m depth, but their measurements of the salinity distribution in the subtidal estuary covered only the upper 2 m of the sediments. A meso- to macrotidal setting was studied at the western coast of France by Anschutz et al. (2009, 2016), Buquet et al. (2016), and Charbonnier et al. (2013). However, their biogeochemical sampling strategy did not involve spatial distribution patterns down to several meters depths, but focused on pore water samples of the top of the water-saturated zone. Consequently, it will be a technical challenge in future studies to monitor processes at meso- and macrotidal settings with USPs extending down to depths of about 25 m as indicated by the hydrological model in our study.

Furthermore, only a few comprehensive reactive transport modeling studies have been conducted to date in the context of submarine groundwater discharge (Cardenas et al., 2008; Porubsky et al., 2011; Spiteri et al., 2008a, 2008b). However, reactive transport modeling will allow to gain an in-depth understanding of biogeochemical processes and will help to verify the conclusions drawn using, for example, pore water concentrations or biogeochemical rate measurements alone. Reactive transport modeling will further help to verify pore water discharge rates estimated using seepage meters or radon isotopes. This will be an essential step to address the question how the pore water discharge of beach systems influences biogeochemistry in nearshore waters.

6. Conclusions

The offshore exposed beach of Spiekeroog Island displays a typical beach at the transition from a dissipative winter morphology to a reflective summer situation. The cross-shore beach transect exhibitsoxic conditions within the dunewdewatitarian freshwater influenced section, changing redox conditions with depth close to the mean high water level, and the dominance of Mn(II)/Fe and SO$_4^{2-}$ reduction below a thin oxycline of some centimeters in the seaward part of the transect. The assumption that saltwater-freshwater interfaces are reaction hotspots is not supported by our data. Instead, biogeochemical processes are influenced by the mixing of water masses of different origin (freshwater/seawater) as well as by changing redox conditions within these two water types. The DOM in the studied pore waters down to 5 m depth is mainly of marine origin as indicated by both CDOM and molecular DOM composition analyses. Only in the dunewdewatian section, CDOM parameters reflect some influence of terrestrial OM. Microbial community patterns seem to be mainly influenced by changing redox conditions and to a lesser extent by salinity variations.

Modeled pore water discharge rates are similar to those proposed for other micro- and mesotidal beach settings. Nutrient effluxes seem to stimulate growth of microphytobenthos on beach sediments of the seepage zone. The beach site shows higher pore water discharge rates compared with a nearby sandy backbarrier tidal flat margin. However, lower nutrient concentrations in the pore waters at the beach site indicate that the backbarrier tidal flat area might be of greater importance for supplying recycled nutrients to nearshore waters of the southern North Sea than the beach systems of the barrier islands.

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Appendix A. Supplementary data

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References


