

Brief Report



# Short-Term Nitrogen and Phosphorus Release during the Disturbance of Surface Sediments: A Case Study in an Urbanised Estuarine System (Gold Coast Broadwater, Australia)

# Ryan J.K. Dunn <sup>1,\*,†</sup>, Nathan J. Waltham <sup>2,‡</sup>, Peter R. Teasdale <sup>3</sup>, David Robertson <sup>3,§</sup> and David T. Welsh <sup>3</sup>

- <sup>1</sup> Griffith School of Engineering, Gold Coast campus, Griffith University, Gold Coast QLD 4222, Australia
- <sup>2</sup> Catchment Management Unit, Gold Coast City Council, PMB 5042 Gold Coast Mail Centre, Gold Coast QLD 9729, Australia
- <sup>3</sup> Environmental Futures Centre, Gold Coast campus, Griffith University, Gold Coast QLD 4222, Australia; p.teasdale@griffith.edu.au (P.R.T.); d.welsh@griffith.edu.au (D.T.W.)
- \* Correspondence: rdunn@apasa.com.au; Tel.: +61-(0)7-55-741-112
- + Present address: RPS APASA Pty Ltd., P.O. Box 5692 Gold Coast Mail Centre, Gold Coast QLD 9726, Australia; rdunn@apasa.com.au
- <sup>‡</sup> Present address: Centre for Tropical Water and Aquatic Ecosystem Research (TropWATER), James Cook University, Townsville QLD 4811, Australia; nathan.waltham@jcu.edu.au
- § Present address: Monash Sustainability Institute, Monash University, Melbourne VIC 3168, Australia; davidr.sci@gmail.com

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Abstract: Understanding the effects of sediment disturbances on nutrient loadings is important for the management of estuarine settings. This study investigated the initial influence of sediment disturbance on water column nutrient concentrations in a shallow estuarine setting within the Gold Coast Broadwater, using a laboratory-based approach. Undisturbed sediment cores (200 mm  $\emptyset$  × 330 mm length, plexiglass) were incubated before and after being subjected to a disturbance event, to investigate the effect on the immediate and subsequent short-term water column nutrient concentrations. Sediment  $NH_4^+_{bio}$  and  $PO_4^{3-}_{bio}$  concentrations ranged from 150 to 478 and 1.50 to 8.56 nmol  $g^{-1}$  dry wt, respectively. Water column NH<sub>4</sub><sup>+</sup> concentrations underwent the greatest increase (>1000% or approx. 14 times greater) immediately following disturbance, with mean effluxes increasing by >300%. Thereafter, water column NH<sub>4</sub><sup>+</sup> concentrations and efflux rates declined to near initial pre-disturbance concentrations. Water column NH4<sup>+</sup> concentrations accounted for 0.58%–5.50% of the depth-integrated sediment NH4<sup>+</sup><sub>bio</sub> concentration, indicating mobilization of the sediment bound exchangeable  $NH_4^+$ . The observed changes in  $PO_4^{3-}$  concentrations and fluxes were much lower in comparison to those observed for N-species. Following disturbance, increases in the water column PO<sub>4</sub><sup>3-</sup> concentration accounted for 7.16%–8.22% depth-integrated sediment bioavailable  $PO_4^{3-}$  at +1 and +2 hours, and 5.65% at +7 hours, respectively. These results provide important insight into the potential implications of disturbance events, such as vessel activities and dredging operations, within the case study region, providing information for potential management options and relevant water quality concerns.

Keywords: nutrients; sediment disturbance; water quality

#### 1. Introduction

Estuarine surface sediments typically accumulate high loads of organic matter and are therefore regions of intense microbial activity and nutrient cycling. They are often characterised by high concentrations of soluble nitrogen and phosphorus from the remineralisation of deposited organic matter and anthropogenic sources [1,2], becoming additional nutrient sources to overlying waters [3]. This exchange of nutrients across the sediment water interface (SWI) modifies both sediment nutrient pools (pore water, exchangeable/adsorbed, and strongly bound) and water column conditions [4,5]. Three key processes for the exchange of nutrients across the SWI are: the passive diffusion of solutes, biota-mediated transport (bioirrigation, bioturbation, and bioresuspension [6]), and the physical disturbance and resuspension of sediment [7,8]. Disturbance and resuspension events may occur under natural circumstances (e.g., tidal currents and wave induced; [9]), but also as a result of anthropogenic activities such as vessel movement (e.g., vessel wake and propeller wash) and mooring, and dredging practices [10,11]. These anthropogenic activities can induce negative impacts on the water quality, altering suspended sediment and nutrient loadings [10,12–14], in addition to modifying the biology of fauna present in bottom sediments [15], as well as water column communities [11,16].

Sediment resuspension often leads to increases in water column nutrient concentrations through the mixing of formerly interstitial waters with the overlying water column and the desorption of nutrients from the resuspended particles or release from dissolving minerals [17,18]. Furthermore, resuspension potentially increases the rates of remineralisation by the bacteria attached to the benthic particles that are released into the water column upon exposure to higher oxygen concentrations [19]. Although it is difficult to estimate the proportion of nutrients that will be transferred between the sediment, sourced from a system of interest, and overlying water during sediment disturbance and resuspension events, estimates of the exchangeable concentrations in sediment slurries in laboratory holding tanks can be estimated. Given the potential for adverse effects of disturbed and resuspended sediments on nutrient loadings within estuarine settings, an understanding of nutrient release processes, influencing factors, and resultant nutrient loads, is important for the sustainable management of an ecologically and economically important estuarine seascape. To date, studies relating to the effects of vessel- and dredge-induced resuspension on suspended sediment and nutrient dynamics include both laboratory-based (closed system; e.g., [11,14] and field-based approaches (open system; e.g., [10,12]). In both cases, these experiments report increases in suspended sediment concentrations and/or water column nutrient concentrations, which vary in magnitude and duration according to influencing factors such as disturbance depth, frequency, intensity, and pre- and post-disturbance sediment and hydrodynamic conditions, e.g., [10–14,20–22].

The objective of this study was to investigate the initial influence of the disturbance and resuspension of estuarine sediment on water column nutrient concentrations using laboratory-based sediment incubations. This case study focuses on sediments and waters directly collected from a shallow urbanised estuarine waterbody within the Gold Coast Broadwater (Australia), Saltwater Creek, a system that experiences, and is likely to be subjected to increasing future, anthropogenic sediment disturbance events [23]. This data contributes to the body of works used to evaluate future sediment disturbances in shallow estuarine settings.

# 2. Materials and Methods

# 2.1. Study Location

Saltwater Creek is an urbanised stream and estuary located on the Gold Coast (Australia). The creek is approximately 17 km long, flowing from its headwaters in the Nerang State Forest to the Gold Coast Broadwater [24] (Figure 1). The average width of the creek is 30 m, with the depth of the first 3 km typically 3 m (below mean sea level), with the remaining upstream region of the creek typically 1 m in depth [25]. Freshwater flow from the catchment is only significant after periods of rainfall. Thunderstorms, common during the summer period (November–January), often result in intense



**Figure 1.** Map of: (a) Australia, showing (b) the Gold Coast Broadwater, and (c,d) the sediment sample collection site within Saltwater Creek (▲) and water collection site nearby the creek entrance (■) (satellite imagery source: Google Earth).

The estuarine zone of the creek is turbid [23,28], although it varies from a relatively clean mouth to having intermittently highly turbid upper reaches [26]. Catchment runoff and biological activity

are known sources of increased sediment loadings, whilst natural sediment resuspension has been identified as playing a lesser role [26]. The suspended sediment concentrations along the estuarine zone range from approx. 20–30 mg L<sup>-1</sup> and 80–300 mg L<sup>-1</sup> during periods without and with rainfall (high tide period), respectively [27]. Furthermore, seasonal NO<sub>x</sub> and NH<sub>4</sub><sup>+</sup> water column concentrations range between 0.5–34 µmol L<sup>-1</sup> and 0.4–7.4 µmol L<sup>-1</sup>, respectively, with rainfall events increasing the nutrient concentrations [27].

Bottom surface sediments along the estuarine zone of the creek are characterised by <63  $\mu$ m and 63–180  $\mu$ m size fractions, accounting for a combined contribution of up to 90% of the total particle size distribution [27]. The mineralogy of the surrounding intertidal and subtidal surface sediments of the area is dominated by halite, quartz, anorthite, and kaolinite [29]. Additionally, Dunn et al. [27] report sediment NH<sub>4</sub>+<sup>bio</sup> concentrations ranging from approx. 200–240 nmol g dry wt<sup>-1</sup> and surface sediments to typically be net sources of dissolved inorganic nitrogen to the overlying water [27].

Bioturbating fauna (e.g., worm (*Simplisetia aequisetis*), amphipod (*Victoriopisa australiensis*), and crab (*Heloecius cordiformis*)) occur within the surface sediments, with *V. australiensis* accounting for up to 45% of the bioturbating macrofaunal density at the sediment collection site [27].

Saltwater Creek and its adjoining waterways have high recreation and amenity values, serving as a major access route for residents of adjoining residential canal estates and providing an entry point for vessels into the Coomera River (Figure 1). A dredging program has been previously conducted in Saltwater Creek to restore navigation passage depths; however, concerns have been raised regarding ongoing access and use of the creek without routine maintenance dredging to maintain channel depths.

Due to considerable urban development/expansion in the catchment, combined with the utility of Saltwater Creek and adjoining waterways as an important fish habitat, the system has been the focus of recent scientific investigation and management intervention (e.g., [23,25–28,30]).

#### 2.2. Sample Collection

Sediment and water was collected in May 2008, 2 km from the mouth of Saltwater Creek (Figure 1), corresponding to site 1 in Dunn et al. [27]. The site sampled represents a section of Saltwater Creek that receives recreational vessel traffic and has been dredged in the past. Three intact cores (70 mm  $\emptyset \times 800$  mm length, PVC) were collected within a 10  $\times$  10 m area from an intertidal-subtidal location, to determine the wet-bulk density, porosity, organic matter determined as loss-on-ignition (LOI<sub>550</sub>), grain size distribution, bioavailable ammonium concentrations (NH<sub>4</sub><sup>+</sup><sub>bio</sub>; porewater + exchangeable NH<sub>4</sub><sup>+</sup>), and sediment bioavailable phosphate concentrations (PO<sub>3</sub><sup>4–</sup><sub>bio</sub>; porewater + exchangeable PO<sub>4</sub><sup>3–</sup>). Upon collection, cores were immediately sliced into eight depth horizons (0–1, 1–2, 2–4, 4–6, 6–10, 10–20, 20–40, >40 cm) and stored in the dark (<4 °C) before freezing within two hours of collection.

To investigate the effects of sediment disturbance and resuspension on nutrient mobilization and water column NO<sub>x</sub>, NH<sub>4</sub><sup>+</sup>, and PO<sub>4</sub><sup>3-</sup> concentrations, six undisturbed sediment cores (200 mm  $\emptyset \times 330$  mm length, plexiglass) were collected from the above mentioned intertidal-subtidal location at low tide, to facilitate ease of core collection, and were relocated to the laboratory. The depth of sediment contained within each core was 160 ± 10 mm. The sediment surface within each core was generally smooth, uniformly horizontal, and void of visible debris. The sediment collection area was considered typical of sediments which would undergo disturbance and resuspension events, such as those created by vessel propeller wash, or dredging operations, due to the close proximity to vessel mooring structures adjacent to private residencies and the downstream positioning from the canal estates.

Additionally, water samples were collected using acid washed (10% v/v HCl) low density polyethylene (LDPE) bottles (Nalgene), to determine dissolved inorganic and total dissolved nitrogen and phosphorus concentrations, and total suspended sediment (TSS) concentrations. Nutrient samples were immediately filtered through pre-washed, pre-ashed 25 mm Ø GF/F (pore size  $\equiv 0.7 \mu$ m) filters and were frozen until analysed. Suspended solids were collected by filtration onto pre-washed,

pre-ashed, and pre-weighed 47 mm Ø GF/F filters. Nutrient samples were frozen (-20 °C) whilst awaiting analysis. In situ physicochemical water column parameters were measured using a calibrated multi-probe unit (TPS 90-FLMV, TPS, Brisbane, Australia).

In addition, a large volume (>500 L) of site water from Saltwater Creek was collected during the flood tide phase on the day of sediment collection to facilitate maintenance and incubation of sediment cores before and during the sediment disturbance and resuspension event, and all subsequent incubation sampling events.

#### 2.3. Sediment Resuspension Flux Measurements

Following collection, cores were returned to the laboratory within two hours of collection, carefully filled with creek water, and submerged in holding tubs for equilibration, according to the methods described in Dunn et al. [27]. Following the equilibration period, water in the holding tanks was replaced with fresh site water and the cores were re-equilibrated for two hours before incubations commenced. To initiate incubations, water levels were lowered in the holding tubs, giving an overlying water depth of ~120 mm (equivalent to ~3.7 L) within each replicate core, resulting in a large volume ratio of sediment to water of approximately 1.4:1. Sediments were initially incubated in dark conditions without disturbance to the SWI, before being subjected to a simulated sediment disturbance and resuspension event, to investigate the change in short-term water column nutrient concentrations and fluxes. Nutrient fluxes were determined following water sample collection -2 and -0.5 hours prior to sediment disturbance (incubation 1), immediately following sediment disturbance (0 hours), +1 hour following disturbance (incubation 2), +2 and +3 hours following disturbance (incubation 3), and +6 and +7 hours following disturbance (incubation 4), respectively. All flux incubations were conducted in dark conditions. Cores were closed during incubations using a floating plastic lid and the water was maintained at a temperature of  $22 \pm 1$  °C following the protocols described in [27]. In between each time series incubation event, additional creek water was carefully added to the pre-existing incubation core waters in order to fully submerge the cores within the holding tank.

Following the initial flux incubation (representing typical conditions prior to sediment disturbance), the entire sediment within each core was mechanically mixed (i.e., sediments overturned with the overlying water) using a manual stirring rod over a 2–3-minute period, simulating a sediment disturbance and resuspension event. Water samples were collected from each core immediately following the sediment disturbance and resuspension to initiate the initial flux determination, and thereafter at the beginning and end of the subsequent flux incubations.

Flux rates were estimated from the difference in individual solute concentrations in the initial and final water samples:

$$F_x = \left( \left( (C_f - C_i) \times V \right) / A \times t \right) \times 10000 \tag{1}$$

where  $F_x$  is the flux of the dissolved species x (µmol m<sup>-2</sup> h<sup>-1</sup>);  $C_f$  and  $C_i$  are the final and initial concentrations of x (µmol L<sup>-1</sup>), respectively; V is the volume of water in the core (L); A is the sediment surface area within the core (cm<sup>2</sup>); and t is the incubation time (h).

Additionally, the percentage change ( $\Delta$ ) in water column nutrient and TSS concentrations was calculated as:

% change = 
$$(([X]_t - [X]_{ti})/[X]_{ti}) \times 100$$
 (2)

where  $[X]_t$  is the water column nutrient or TSS concentration at time *t* and  $[X]_{ti}$  is the water column nutrient or TSS concentration at time *i*.

#### 2.4. Sample Handling and Analytical Techniques

The sediment wet-bulk density, porosity, and  $\text{LOI}_{550}$  were determined following the techniques described in Dunn et al. [31]. The proportion of clay and silt (<63 µm), sand (180 < x > 63 µm), and gravel (>180 µm) fractions were determined by dry sieving and are expressed as the percent of dry weight. Sediment NH<sub>4</sub><sup>+</sup><sub>bio</sub> and PO<sub>4</sub><sup>3-</sup><sub>bio</sub> concentrations were determined following the extraction of

1 cm<sup>3</sup> of homogenised sediment for 24 hours in 9 mL 2 M KCl and 1 M MgCl<sub>2</sub>, respectively, as outlined in [32] and [33], respectively.

During sediment incubations, water samples were collected using acid washed and Milli-Q element water rinsed 60 mL plastic syringes and tubing. Nutrient samples were filtered through 25 mm  $\emptyset$  GF/F (pore size  $\equiv 0.7 \mu$ m) filters and were frozen (-20 °C) until analysis.

Concentrations of water column  $NO_2^- NO_3^-$ ,  $NH_4^+$ ,  $PO_4^{3-}$ , and extracted bioavailable  $NH_4^+$ and  $PO_4^{3-}$  concentrations were determined using an automated nutrient analyser (Easychem Plus Random Access Analyzer; Systea Analytical Technologies, Anagni, Italy), incorporating standard colorimetric analyses. Deionised (Milli-Q Element) water and filtered low nutrient seawater were used for all sample preparation and analyses. Natural filtered seawater certified reference materials produced by the National Low Level Nutrient Collaborative Trials, in addition to laboratory standards, were used for quality assurance purposes. The total suspended sediment (mg L<sup>-1</sup>) concentrations were determined gravimetrically.

# 2.5. Data Presentation and Statistical Analysis

Physicochemical sediment data, water column parameters, and nutrient flux rates are reported as mean  $\pm$  standard deviation. Estimates of sediment-water column nutrient fluxes (µmol m<sup>-2</sup> h<sup>-1</sup>) and sediment depth profiles of bioavailable nutrients are utilised to investigate the potential release of nutrients following sediment disturbance within the estuarine sediments of Saltwater Creek. Comparisons of physicochemical parameters of the various depth horizons were performed by ANOVA and the means were compared using Tukey's HSD analysis. Pearson correlations were used to identify relationships between physicochemical sediment parameters of the various depth horizons. The significance of observed differences between TSS and dissolved inorganic nutrient concentrations, and nutrient flux rates before and following the sediment disturbance and resuspension event (n = 6), was calculated with a paired *t*-test. Normality was tested using the Kolmogorov-Smirnov test, and adjusted as necessary to satisfy the ANOVA assumptions. The total suspended sediments, dissolved inorganic nutrient concentrations, and flux rates following the resuspension event, were compared by ANOVA using SPSS for Windows (version 19, SPSS Inc., Chicago, IL, USA).

#### 3. Results and Discussion

# 3.1. Background Sediment and Water Column Conditions

Depth profiles of the surface sediment properties are presented in Table 1. Grain sizes between 63–180  $\mu$ m were predominant, accounting for 51.3% to 63.2% of the total sediment mass at each depth. The contribution of fine grained sediments (<63  $\mu$ m) ranged from 24.1% to 36.7%, being greatest in the upper 1–2 cm depth horizon. No other discernible trends in grain size distributions with depth were evident. The sediment density and porosity values ranged from 1.37 to 1.78 g cm<sup>-3</sup> and 64.2% to 76.1%, respectively. Wet-bulk densities and porosity values demonstrated typical vertical profiles, where the density increases and the porosity decreases with depth [34].

**Table 1.** Depth profiles of physicochemical parameters in surface sediments of Saltwater Creek (mean  $\pm$  standard deviation; *n* = 3).

Depth (cm)	Wet-Bulk Density (g cm <sup>-3</sup> )	Porosity (%)	LOI <sub>550</sub> (%)	<63 µm (%)	63–180 μm (%)	>180 μm (%)	NH4 <sup>+</sup> <sub>bio</sub> (nmol g <sup>-</sup>	PO4 <sup>3-</sup> bio <sup>-1</sup> dry wt)
0-1	$1.37\pm0.05$	$76.1\pm6.91$	$9.23\pm3.56$	$24.1\pm4.08$	$60.8\pm2.39$	$15.1\pm5.43$	$478 \pm 189$	$8.56 \pm 1.03$
1–2	$1.38\pm0.07$	$70.1\pm5.22$	$9.16\pm2.88$	$36.7\pm22.17$	$63.2\pm8.04$	$13.1\pm7.64$	$208\pm85.1$	$7.72 \pm 1.95$
2–4	$1.51\pm0.16$	$68.4 \pm 1.47$	$7.68 \pm 2.01$	$26.5\pm3.33$	$61.3\pm6.40$	$12.2\pm3.24$	$181\pm103$	$5.57 \pm 1.01$
4-6	$1.53\pm0.15$	$66.3\pm2.76$	$7.77 \pm 2.50$	$27.9\pm2.70$	$60.6 \pm 4.77$	$11.4\pm6.48$	$212\pm90.7$	$4.27 \pm 1.67$
6-10	$1.66\pm0.07$	$66.1\pm3.78$	$6.72 \pm 1.12$	$29.7\pm5.13$	$59.7\pm3.71$	$10.5\pm7.97$	$186\pm93.1$	$1.50\pm0.56$
10-20	$1.68\pm0.51$	$66.9\pm5.93$	$7.66\pm0.81$	$34.9\pm4.60$	$55.4 \pm 4.63$	$9.32 \pm 2.88$	$161\pm51.9$	$2.53\pm0.32$
20-40	$1.73\pm0.16$	$65.1 \pm 4.95$	$6.51 \pm 1.25$	$30.7\pm3.86$	$51.3\pm5.15$	$16.5\pm5.72$	$150\pm87.6$	$2.11\pm0.71$
>40	$1.78\pm0.32$	$64.2\pm5.87$	$4.12\pm2.11$	$34.8\pm4.47$	$57.7 \pm 4.86$	$6.94 \pm 6.80$	$155\pm60.4$	$2.96 \pm 1.13$

Sediment NH<sub>4</sub><sup>+</sup><sub>bio</sub> and PO<sub>4</sub><sup>3-</sup><sub>bio</sub> concentrations ranged from 150 to 478 and 1.50 to 8.56 nmol g<sup>-1</sup> dry wt, respectively. NH<sub>4</sub><sup>+</sup><sub>bio</sub> concentrations were greater in the upper-most surface sediments (0–1 cm), exhibiting concentrations two-fold greater than those measured at any other depth, with a further decrease in concentrations below a 10 cm depth. Similarly, PO<sub>4</sub><sup>3-</sup><sub>bio</sub> concentrations were significantly greater (*F* = 17.069, *p* = 0.006) in the upper sediments (0–10 cm) compared to the deeper sediments below a 10 cm depth. Concentrations corresponded to those previously observed within the sediments of Saltwater Creek and neighbouring estuarine systems [5,27,28,31]. Sediment organic matter (LOI<sub>550</sub>) was greatest in the surface 0–2 cm depth horizon and decreased with depth thereafter (Table 1). Sediment bioavailable nutrient concentrations were significantly correlated with sediment density (PO<sub>4</sub><sup>3-</sup><sub>bio</sub>; *r* = 0.922, *p* = 0.001), porosity (NH<sub>4</sub><sup>+</sup><sub>bio</sub>; *r* = 0.919, *p* = 0.001, PO<sub>4</sub><sup>3-</sup>; *r* = 0.868, *p* = 0.005), and LOI<sub>550</sub> (PO<sub>4</sub><sup>3-</sup>; *r* = 0.726, *p* = 0.041) (Table 2).

In situ water column dissolved nutrient concentrations measured from water samples obtained during flood tide conditions were:  $0.59 \pm 0.15 \mu mol L^{-1}$  for NO<sub>3</sub><sup>-</sup>,  $0.81 \pm 1.18 \mu mol L^{-1}$  for NO<sub>x</sub>,  $0.54 \pm 0.22 \mu mol L^{-1}$  for NH<sub>4</sub><sup>+</sup>, and  $0.34 \pm 0.14 \mu mol L^{-1}$  for PO<sub>4</sub><sup>3-</sup>, which correspond to the ranges previously reported for Saltwater Creek and adjoining waters [5,27–30]. The mean in situ TSS concentrations during the flood tide conditions were  $24 \pm 5.3 \text{ mg L}^{-1}$ . The snapshot survey of the in situ physicochemical parameters revealed that the values were:  $7.8 \pm 0.2 \text{ pH}$ ,  $28.1\% \pm 0.1\%$  salinity,  $24.3 \pm 0.2 \text{ °C}$  temperature, and  $85.0\% \pm 2.3\%$  saturation for dissolved oxygen. These parameters have been demonstrated to vary considerably within Saltwater Creek according to location and rainfall runoff [25–28].

Parameter		Porosity (%)	LOI <sub>550</sub> (%)	<63 μm (%)	63–180 μm (%)	>180 µm (%)	NH4 <sup>+</sup> <sub>bio</sub> (nmol g <sup>-</sup>	PO4 <sup>3-</sup> bio <sup>1</sup> dry wt)
$\mathbf{M}_{1} = (1 + 1) + (1 $	r	-0.850 **	-0.891 **	0.365	0.755 *	-0.470	-0.684	0.922 **
wet-buik density (g cm *)	p	0.008	0.003	0.374	0.030	0.240	0.61	0.001
Porosity(0/)	r		0.784 *	0.473	0.498	0.495	0.919 **	0.868 **
1 010sity (78)	р		0.021	0.236	0.209	0.213	0.001	0.005
$I \cap I_{rro}$ (%)	r			-0.301	0.488	0.576	0.581	0.726 *
LO1550 (78)	р			0.468	0.220	0.135	0.131	0.041
<63 um (%)	r				-0.182	-0.455	-0.611	-0.278
<05 µm (78)	р				0.666	0.258	0.108	0.505
63-180 um (%)	r					-0.126	0.381	0.672
03-100 µm (78)	р					0.766	0.352	0.068
>180 um (%)	r						0.432	0.407
>100 µm (70)	р						0.285	0.316
$NH_{+}$ , (pmol $a^{-1}$ dry wt)	r							0.725 *
till bio (tillor g dry wt)	р							0.042

**Table 2.** Pearson correlation coefficients for physicochemical sediment parameters of the various depth horizons (n = 8; \* correlation significant at the 0.05 level (2-tailed); \*\* correlation significant at the 0.01 level (2-tailed)).

# 3.2. Sediment Disturbance Simulation and Sediment Incubations

Following the sediment disturbance and resuspension event, TSS concentrations immediately increased by >1000% or greater than 30 times higher, compared to concentrations measured prior to the simulated event (Table 3). Thereafter, TSS concentrations decreased over the duration of the incubations through both particle settlement on the sediment surface and TSS removal (and dilution) from the mesocosm during the part-replenishment of incubation waters. The reduction of TSS concentrations in the incubation cores demonstrated a significant correlation (r = 0.986, p = 0.002) with time (Figure 2). Concentrations of TSS during incubations following the disturbance and resuspension event represent gross overestimates of the in situ water column concentrations that would exist throughout a small estuary like Saltwater Creek, other than those waters within the immediate vicinity of a disturbance event. The settlement of heavier particles and tidal exchange with "cleaner" waters would likely result in rapidly decreasing TSS concentrations away from the disturbance source.

Time (h)	Initial Concentration									
Time (ii)	TSS (mg L <sup>-1</sup> )		$NO_3^{-}$ (µmol L <sup>-1</sup> )		$NO_x$ (µmol L <sup>-1</sup> )		$\mathrm{NH_4^+}$ (µmol L $^{-1}$ )		PO4 <sup>3-</sup> (µmol L <sup>-1</sup> )	
-2	$29\pm1$		$0.50\pm0.08$		$0.90\pm0.16$		$0.21\pm0.05$		$0.22\pm0.05$	
Time (h)	$\Delta$ TSS		$\Delta NO_3^-$		$\Delta NO_x$		$\Delta NH_4^+$		$\Delta PO_4^{3-}$	
	$(mg L^{-1})$	%	$(\mu mol \ L^{-1})$	%	$(\mu mol \ L^{-1})$	%	$(\mu mol \ L^{-1})$	%	$(\mu mol \ L^{-1})$	%
-0.5	-1	-3	0.12	24.7	0.14	15.8	0.27	125.2	-0.01	-3.5
0	932	>1000	0.28	44.8	0.30	28.6	6.26	>1000	0.03	13.5
+1	-189	-20	0.38	42.5	0.49	36.5	5.67	84.1	0.08	33.8
+2	-56	-7	-0.29	-22.4	-0.39	-21.6	-11.10	-89.5	0.05	14.8
+3	-39	-5	0.21	21.3	0.19	13.5	0.29	22.2	-0.01	-3.7
+6	-159	-23	-0.85	-70.1	-0.84	-51.5	-0.27	-17.3	-0.13	-34.9
+7	-46	-9	0.03	7.1	0.07	8.4	0.20	15.0	0.02	9.6

**Table 3.** Time-series of mean changes (n = 6) in total suspended sediment and dissolved inorganic nutrient concentrations according to mesocosms sampling timeline. Negative times represent before, time 0 represents immediately prior to, and positive times represent hours after the sediment disturbance and resuspension event, respectively.



**Figure 2.** Total suspended solid concentrations during core incubations. The occurrence of the sediment disturbance and resuspension event is represented by  $\bigtriangledown$  and shading.

Nitrogen fluxes determined prior to the sediment disturbance and resuspension event were typical of the effluxes previously observed for sediments under quiescent conditions, not accounting for sediment disturbance events, within Saltwater Creek [27], a nearby adjoining system [5], and regional southern Moreton Bay waters [4]. Time-series dissolved inorganic nutrient concentrations and estimated flux rates are presented in Table 3 and Figure 3, respectively. Concentrations of  $NO_3^-$  and  $NO_x$  demonstrated effluxes from the bulk sediment prior to disturbance and an increase in water column concentrations after the simulated disturbance and resuspension event (Table 3, Figure 3a,b). Following resuspension, water column NO<sub>3</sub><sup>-</sup> and NO<sub>x</sub> concentrations and NO<sub>3</sub><sup>-</sup> and  $NO_x$  effluxes decreased below those measured prior to the resuspension event. Water column  $NH_4^+$ concentrations underwent the greatest increase (>1000% or approx. 14 times greater) immediately following the disturbance and resuspension event (Table 3), with mean effluxes increasing by >300% following sediment disturbance and resuspension (Figure 3c). Nutrient fluxes associated with recently resuspended sediments have been shown to significantly increase due to both natural circumstances, e.g., [35–39], and anthropogenic disturbance and resuspension events, e.g., [14,40,41]. Increases in NH4<sup>+</sup> effluxes due to the simulated anthropogenic sediment disturbance and resuspension event exceeded those reported under the natural resuspension circumstances of two-six times reported by Corbett [35].



**Figure 3.** Calculated fluxes for (**a**) NO<sub>3</sub><sup>-</sup>, (**b**) NO<sub>*x*</sub>, (**c**) NH<sub>4</sub><sup>+</sup>, and (**d**) PO<sub>4</sub><sup>3-</sup> during core incubations. The occurrence of the sediment disturbance and resuspension event is represented by  $\bigtriangledown$  and shading.

Following disturbance and resuspension,  $NH_4^+$  concentrations during this study declined to near initial concentrations, as did efflux rates. Estuarine sediments typically contain significant pools of  $NH_4^+$  and when resuspended,  $NH_4^+$  enriched porewater is immediately mixed into the overlying receiving water. It was therefore expected that, following the disturbance of the sediments within the mesocosms, increases in the  $NH_4^+$  concentration would be observed in the overlying water column, which was observed during this study. These increased concentrations represent the mobilisation of pore water  $NH_4^+$ , and potentially the adsorption of  $NH_4^+$  onto solid phase surfaces, when sediment particles are resuspended in the overlying water. Following the disturbance and resuspension event, the increase in water column  $NH_4^+$  concentrations accounted for 0.58%–5.50% of the depth-integrated sediment  $NH_4^+_{bio}$  concentration (Table 4).

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**Table 4.** Mean water column nutrient concentration as a percentage of depth-integrated (0–15 cm) sediment  $NH_4^+_{bio}$  and  $PO_4^{3-}_{bio}$  concentrations during incubations. Negative times represent before, time 0 represents immediately prior to, and positive times represent hours after the sediment disturbance and resuspension event, respectively.

Nutrient	Time (hours)										
TVUITEIL	-2	-0.5	0	+1	+2	+3	+6	+7			
$NH_4^+$ PO $t^{3-}$	0.09	0.21	2.99 5.36	5.50 7.16	0.58	0.71	0.58	0.67			
104	4.09	4.72	5.50	7.10	0.22	7.92	5.15	5.05			

Upon sediment disturbance and resuspension events, porewater  $NH_4$ + and exchangeable  $NH_4^+$  within surface sediments are usually released into the receiving waters, as a new equilibrium establishes between the dissolved and exchangeable pools [42].

Due to the typical dominant contribution of  $NO_3^-$  to  $NO_x$  within Saltwater Creek waters,  $NO_3^-$  and  $NO_x$  water column concentrations demonstrated similar responses to the disturbance and resuspension of the sediment column within the mesocosms (Table 3). Increased water column concentrations are presumably attributable to the release of  $NO_3^-$  already present in the sediment porewater, as a result of nitrification within oxic sediment zones. In Saltwater Creek, nitrification within the sediment has been shown to be the primary source of nitrate fuelling nitrate reduction, with  $NO_3^-$  reduction rates coupled to nitrification accounting for ~75% of total nitrate reduction in dark conditions [27]. The removal of dissolved  $NO_x$  from the water column is likely due to denitrification and dissimilatory nitrate reduction to ammonium in the surface sediment and anoxic zones within suspended solids or even in the water column. However, the burial of nitrifiers in anoxic zones as suspended solids resettle would presumably limit nitrification rates and consequently nitrate reduction rates coupled to this process after disturbance events.

Prior to resuspension,  $PO_4^{3-}$  fluxes were directed into the sediments (i.e., mean negative flux), whilst immediately following sediment resuspension, increased water column  $PO_4^{3-}$  concentrations resulted from effluxes (Figure 3d). The observed changes in  $PO_4^{3-}$  concentrations and fluxes were much lower in comparison to those observed for inorganic N-species. Subsequent to the resuspension event,  $PO_4^{3-}$  fluxes decreased and were variable in direction (Table 3 and Figure 3d). Water column  $PO_4^{3-}$  concentrations measured during the last sample events (+6 hours and +7 hours) were consistent with the initial water column concentrations.

Phosphorus-liberation from sediments to the overlying water depends on the process of diffusion, based on the difference in phosphate concentration between overlying water and interstitial water [43,44]. Whilst the source of phosphate in sediments is the microbial remineralisation of deposited organic matter, an important factor in the cycling of  $PO_4^{3-}$  in aquatic systems is its interaction with iron (III) oxyhydroxides (FeOOH), where  $PO_4^{3-}$  binds to precipitated FeOOH within surface sediments, decreasing its bioavailability [45].

During the disturbance and resuspension event, porewater  $PO_4^{3-}$  in the deeper anoxic sediment zones of the cores would have been released into the water column, initially increasing  $PO_4^{3-}$ concentrations. However, as co-released porewater dissolved Fe(II) auto-oxidised and precipitated as FeOOH in the oxic water column, much of the mobilised  $PO_4^{3-}$  was presumably adsorbed onto the surfaces of the newly formed FeOOH minerals in the water column, resulting in the decrease in the  $PO_4^{3-}$  concentrations in the water column. This would ultimately lead to  $PO_4^{3-}$  once again being sequestered in the sediment when these iron minerals settled out of suspension. Following the disturbance and resuspension event, the increase in the water column  $PO_4^{3-}$  concentration accounted for 7.16%–8.22% of depth-integrated sediment bioavailable  $PO_4^{3-}$  at +1 hours and +2 hours following sediment disturbance and resuspension, and 5.65% +7 hours following disturbance and resuspension (Table 4).

# 4. Conclusions

The results from this study provide an approximation of resultant water column nutrient conditions within Saltwater Creek (grain size dominated by <63 mm and 63–180 mm size fractions), providing insight for potential management options and relevant water quality concerns. The exchangeable  $NH_4^+$  measured for a particular volume of sediment can be used as a theoretical maximum for  $NH_4^+$  that could potentially be released during sediment disturbance and resuspension events. In practice, however, this maximum is unlikely to be realised, during which an intermediate amount of the nutrient is liberated, as observed in this study, where only 0.58%–5.50% of the bioavailable  $NH_4^+$  present in the sediment contributed to the water column concentration. Conversely, disturbance and resuspension events are unlikely to have a significant effect on water column  $PO_4^{3-}$  concentrations, other than locally, where in the short term, there may be a temporary "spike" in concentrations until phosphates mobilised from sediment porewaters reassociate with suspended sediment particles and newly formed FeOOH minerals, before settling out.

This manipulative study presents a case that during disturbance and resuspension events, there is a net release of nutrients to the water column, which is then followed by a decrease in the sediment NH<sub>4</sub><sup>+</sup> pool. Increases of NO<sub>3</sub><sup>-</sup> and NH<sub>4</sub><sup>+</sup> within the water column will occur in conjunction with a reduction in deeper sediment NH<sub>4</sub><sup>+</sup> pools along the disturbed creek sections. Due to the strong binding of phosphate to FeOOH and other sediment particles, disturbance events are unlikely to liberate substantial amounts of PO<sub>4</sub><sup>3-</sup> contained in the sediment, other than in the short-term, with a short spike in local concentrations followed by a decrease as PO<sub>4</sub><sup>3-</sup> adsorbs or binds to the suspended sediment (demonstrated by the low fluxes across the SWI), with a depletion of the PO<sub>4</sub><sup>3-</sup> water column upon re-settling. Overall, the results here provide an important insight into the implications of sediment disturbances, resulting from vessel activities and dredging operations, of urbanised coastal sediments of the region. It is apparent that even the short term disturbance of estuarine sediments (dominated by <63 mm and 63–180 mm size fractions) would contribute to degraded water quality conditions through an initial increase in TSS and nutrient loadings, which requires full consideration when assessing the risks to local aquatic values.

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