



Controls on soil organic carbon stocks in tidal marshes along an estuarine salinity gradient

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Abstract. Tidal marshes are sedimentary environments and are among the most productive ecosystems on Earth. As a consequence they have the potential to reduce atmospheric greenhouse gas concentrations by sequestering organic carbon (OC). In the past decades, most research on soil organic carbon (SOC) storage in marsh environments has focused on salt marshes, leaving carbon dynamics in brackish and freshwater marshes largely understudied and neglecting the diversity among tidal marshes. We therefore conducted an extensive sampling campaign to quantify and characterize SOC stock in marshes along a salinity gradient in the Scheldt estuary (Belgium and the Netherlands). We find that SOC stocks vary significantly along the estuary, from 46 in freshwater marshes to 10 kg OC m⁻² in salt marshes. Our data also show that most existing studies underestimate total SOC stocks due to shallow soil sampling, which also influences reported patterns in OC storage along estuaries. In all sampled tidal marsh sediments the SOC concentration is more or less constant from a certain depth downward. However, this concentration decreases with increasing salinity, indicating that the amount of stable SOC decreases from the upper estuary towards the coast. Although the net primary production of macrophytes differs along the estuary, our data suggest that the differences in OC storage are caused mainly by variations in suspended sediment concentration and stable particulate OC (POC) content in the water along the estuary. The fraction of terrestrial suspended sediments and POC that is transported downstream of the maximum turbidity zone is very limited, contributing to smaller amounts of long-term OC sequestration in brackish and salt marsh sediments. In addition, high rates of sediment deposition on freshwater tidal marshes

in the maximum turbidity zone promote efficient burial of OC in these marsh sediments.

1 Introduction

As a consequence of increasing atmospheric greenhouse gas concentrations and the recognition that soils have the potential to store vast amounts of organic carbon (OC), there is large interest in the OC storage potential of soils in different ecosystems (Duarte et al., 2013; Govers et al., 2013; Scharlemann et al., 2014). Although coastal vegetated habitats occupy only 0.2 % of the ocean surface, it has been estimated that they account for approximately 50 % of carbon burial in marine sediments, referred to as blue carbon (Donato et al., 2011; Duarte et al., 2013; Mcleod et al., 2011; Nelleman et al., 2009). It has recently been shown that the OC sequestration potential of these ecosystems will depend to a large extent on future climatic changes and sea level rise (Cherry et al., 2009; Kirwan and Blum, 2011; Kirwan and Mudd, 2012; Weston et al., 2011). Moreover, changing carbon storage in these ecosystems can potentially cause important feedbacks to atmospheric concentrations of carbon dioxide (CO₂) and methane (CH₄; Duarte et al., 2013; Pendleton et al., 2012; Poffenbarger et al., 2011). Constraining the amount of OC that is sequestered in these ecosystems and understanding the processes controlling the size of this pool are of major importance for understanding the global carbon cycle.

A significant fraction of coastal wetlands is occupied by tidal marshes. These are vegetated intertidal areas located along coastlines and estuaries in sub-Arctic to tropical climates (although they occur mostly in temperate zones) and

are among the most productive ecosystems on Earth (Rocha and Goulden, 2009; Whigham, 2009). Their elevation increases as a consequence of the deposition of both mineral sediments and allochthonous organic matter (OM) during flooding events on the one hand and the incorporation of in situ produced biomass (both above and below ground) on the other hand (Fagherazzi et al., 2012; Neubauer, 2008). Recently formed young tidal marshes, with a low elevation, receive more mineral sediments than their higher counterparts, with sedimentation rates decreasing through time until the marsh platform elevation is in equilibrium with the local mean high water level (Temmerman et al., 2003).

Despite the fact that the importance of vegetated coastal ecosystems and tidal marshes is now widely recognized, estimates of the total amount of OC stored in tidal marshes are subject to a large uncertainty. Estimates of soil organic carbon (SOC) stocks in salt marshes (i.e. tidal marshes bordering saltwater bodies) range between 0.4 and 6.5 Pg (Bridgman et al., 2006; Chmura et al., 2003; Duarte et al., 2013). To the best of our knowledge, no global estimates are available for brackish and freshwater marshes.

There are multiple reasons for the large uncertainty in estimates of the global OC storage in tidal marsh soils. Firstly, the total area of global salt marshes currently used to estimate global stocks is poorly constrained, with estimates between 22 000 and 400 000 km² (Chmura et al., 2003; Woodwell et al., 1973), while a global inventory for freshwater marshes is lacking (Barendregt and Swarth, 2013). Secondly, the available data set on SOC stocks is limited, both in terms of the number of samples analysed and the geographical scope: hitherto, most studies were carried out in a limited number of estuaries, mostly located on the southern and eastern coasts of North America. Differences in sampling procedure and depth beneath the soil surface also contribute to uncertainty. Very often only topsoil samples are analysed, with a limited number of studies considering carbon storage in deeper horizons, although it has been recognized that these also store a significant amount of OC (Elschot et al., 2015; Wang et al., 2011). An additional factor complicating the extrapolation of data to tidal marshes, for which no data are available, is the wide range of reported OC contents for tidal marsh soils (Bouillon and Boschker, 2006; Middelburg et al., 1997). A third important reason for the uncertainties mentioned above is that tidal marshes in estuaries are characterized by steep gradients of multiple environmental and ecological factors (Craft, 2007). A sharp increase in salinity towards the coast is present, resulting in a longitudinal estuarine gradient from salt marshes in the most seaward part over brackish marshes to freshwater tidal marshes. As a consequence of this salinity gradient a vegetation gradient develops, with macrophyte biomass generally being higher on freshwater and brackish marshes compared to salt marshes (Dausse et al., 2012; Hansen et al., 2016; Weston et al., 2014; Wieski et al., 2010). In addition, the OC input in tidal marsh soils is a mixture of upland, riverine, estuarine and marine

sources and the relative contribution of these sources to the total OC pool varies significantly along the estuary (Middelburg and Nieuwenhuize, 1998).

Currently available data suggest that these environmental gradients along estuaries generally result in decreasing SOC stocks with increasing salinity (Craft, 2007; Hansen et al., 2016; Hatton et al., 1983; Wieski et al., 2010). However, our knowledge on how location along the estuary affects total SOC stocks and which processes control the magnitude of these stocks is, at present, still very limited, mainly because most studies only consider SOC storage down to a depth of 0.3 m and generally the reasons for the observed variability are not identified. Nevertheless, understanding the effect of environmental gradients on SOC dynamics in tidal marshes is important. Such understanding may not only help to improve our estimates of current SOC storage but will also be of great help in assessing the effects of sea level rise on these SOC stocks (Morrissey et al., 2014; Poffenbarger et al., 2011; Weston et al., 2011). Here, we study the variation in SOC inventories in tidal marshes along a salinity gradient in the Scheldt estuary, located in Belgium and the Netherlands. This estuary is characterized by strong gradients in salinity and sediment concentration, making it a suitable location to investigate the impact of these gradients on OC stocks in tidal marsh soils. The aims of this study are therefore (1) to determine the SOC stocks of tidal marsh soils along the salinity gradient of a temperate estuary, (2) to determine the main controls on SOC stocks along this gradient and (3) to use this knowledge to assess how future environmental changes may influence SOC stocks in estuarine tidal marsh soils.

2 Materials and methods

2.1 Study sites

The Scheldt river is located in western Europe and flows into the North Sea in the southern Netherlands (Fig. 1). The estuary of the river extends from its mouth up to 160 km upstream where the tide is stopped by sluices near the city of Ghent (Belgium). The estuary is influenced by a semi-diurnal meso- to macrotidal regime, with mean tidal ranges between 3.8 m at the mouth and 5.2 m in the inner estuary (Meire et al., 2005). The estuary has a total length of about 235 km (including tributary tidal rivers) and comprises a salt or polyhaline zone (salinity > 18), brackish or mesohaline zone (salinity 5–18) and freshwater/oligohaline zone (salinity 0–5; Fig. 1). The Scheldt estuary is described in detail in Van Damme et al. (2005) and Meire et al. (2005).

Tidal marshes are present along the entire length of the estuary and tributary tidal rivers, resulting in approximately 518 ha of freshwater marshes, 3035 ha of brackish marshes and 652 ha of salt marshes (Tolman and Pranger, 2012; Van Braeckel et al., 2013). We sampled a salt, brackish and freshwater tidal marsh, and within each marsh we sampled two

Table 1. Main properties of the sampled tidal marshes.

Name	Name in this study	Vegetation	Tidal range (m) ^a	Elevation relative to local MHWL (m)	POC% of suspended sediment ^b	Marsh sediment depth (m) ^c
Notelaar marsh	Freshwater low	<i>Phragmites australis</i>	5.14	+0.24	6–10	1.2
	Freshwater high	<i>Salix</i> sp. + <i>Urtica dioica</i>	5.14	+0.25	6–10	> 1.4
Waarde marsh	Brackish water low	<i>Elymus athericus</i>	4.85	+0.01	4–5	0.8
	Brackish water high	<i>Elymus athericus</i>	4.85	+0.14	4–5	> 1.4
Paulina marsh	Saltwater low	<i>Spartina anglica</i>	4.19	−0.66	3–4	0.2
	Saltwater high	Mixed vegetation ^d	4.19	+0.11	3–4	0.6

^a From Meire et al. (2005). ^b From Abril et al. (2002). ^c Based on depth profiles of grain size and OC concentration. ^d *Atriplex portulacoides*, *Limonium vulgare*, *Triglochin maritima*, *Elymus athericus*, *Puccinellia maritima*.

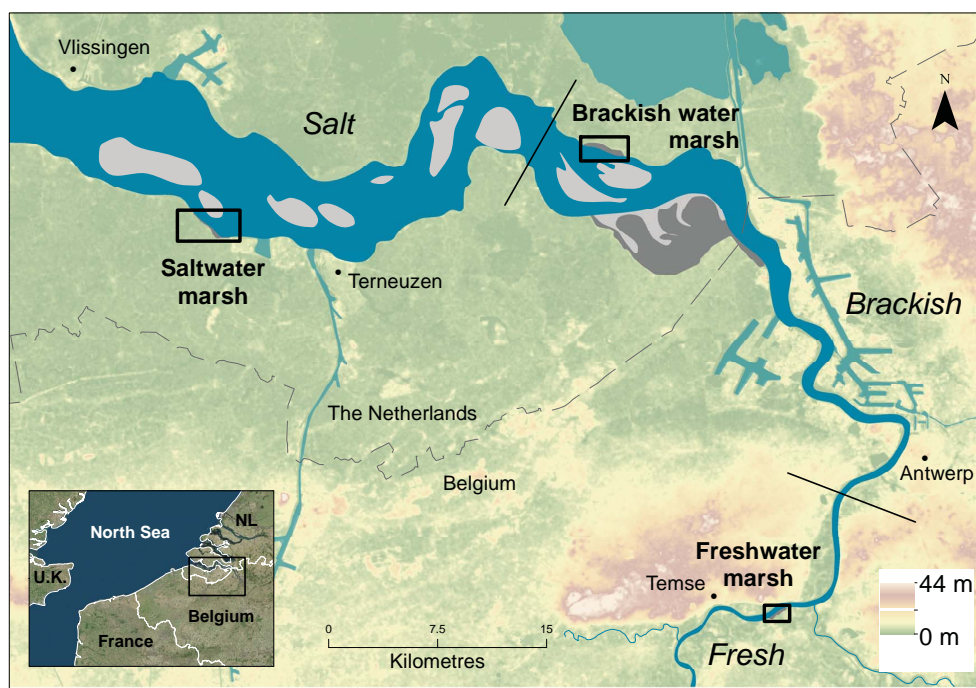


Figure 1. Map of the Scheldt estuary showing the salinity zones and the location of the sampled tidal marshes in a western European context. Intertidal sand flats are depicted in light grey.

locations with different but known rates of historical sediment accretion (Fig. 2 and Table 1). The first location was at the high marsh with an elevation of 0.1 to 0.3 m above mean high water level (MHWL), which has been accreting during the past decades at a rate that is in equilibrium with the rise of MHWL. At the second location, marsh formation started during the past decades at heights well below MHWL. Average accretion rates at these locations were therefore significantly higher than sea level rise (Fig. 2). For the period 1955–2002, the average accretion rates at low and high marshes were 0.5 and 0.75 cm yr^{−1} for the salt marshes 1–2 and 0.5–1 cm yr^{−1} for the brackish marshes and 1–2 and 1 cm yr^{−1} for the freshwater marshes (Temmerman et al., 2004). The vegetation history for the different sites is shown in Fig. 3. The locations of

the sampled tidal marshes are indicated in Fig. 1 and GPS coordinates of the sampling locations are provided in Table S1 in the Supplement.

2.2 Sample collection

Depth profiles were collected in November 2014 using a manual gouge auger (0.06 m diameter) down to a maximum depth of 1.4 m. Three replicate soil cores were collected with a maximum distance of 3 m in between the coring locations. The cores were divided into 0.03 m sections and every soil sample was stored in a resealable bag in the field before transport to the lab. Samples for soil bulk density and root density measurements were collected using a Kopecky ring sampler

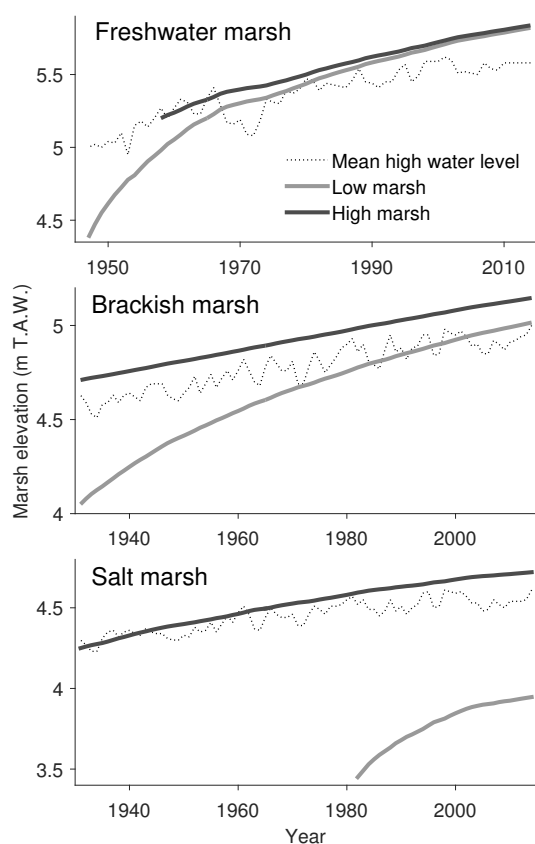


Figure 2. Evolution of marsh surface elevation and mean high water level (relative to Belgian ordnance level, m T.A.W.) at the sampled locations (based on Temmerman et al., 2004).

or with the gouge auger if soil wetness prevented the use of Kopecky rings. These samples were collected at the soil surface and at depth increments of 0.1 up to 0.5 m, then in 0.2 m increments down to the maximum depth of 1.4 m. Above-ground biomass was collected on a surface area of 0.25 m² (five replicates) at the end of August 2015 at each coring location. The difference in timing between soil and biomass collections is due to the fact that standing biomass is maximum in August in western European tidal marshes (Groenendijk, 1984; De Leeuw et al., 1990).

2.3 Soil and biomass analysis

Before analysis of the soil samples, macroscopic vegetation residues were removed manually using tweezers. The soil samples were oven-dried at 35 °C for 48 h and crushed until they passed through a 2 mm sieve. After carbonates were removed with a 10 % HCl solution, the samples were analysed for OC, $\delta^{13}\text{C}$ and C : N ratio using an Elemental Analyser (FlashEA 1112 HT, Thermo Scientific). Soil texture was determined using a laser diffraction particle size analyser (LSTM 13 320, Beckman Coulter) and grain size was classified into clay (< 2 μm), silt (2–63 μm) and sand (> 63 μm)

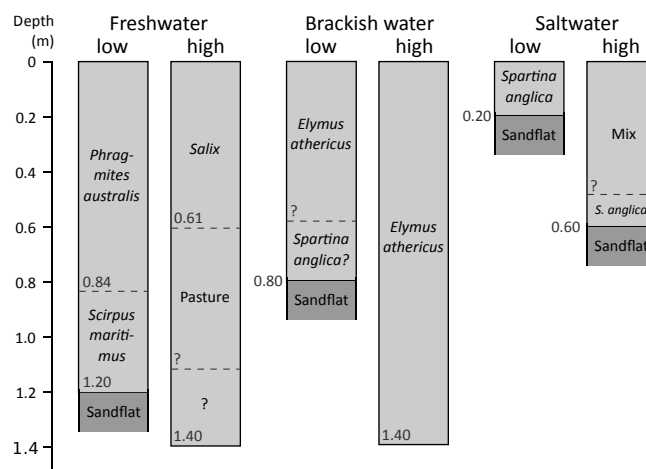


Figure 3. Depth profiles of the sampled tidal marshes showing the vegetation history at each location. At locations where the sand flat was reached this is indicated, at the other locations the marsh sediments extended below 1.4 m depth. The vegetation history is based on Temmerman et al. (2003) and information from the $\delta^{13}\text{C}$ profiles of this study, in combination with information from Boschker et al. (1999) and Middelburg et al. (1997). Mix denotes mixed vegetation which includes *Atriplex portulacoides*, *Limonium vulgare*, *Triglochin maritima*, *Elymus athericus* and *Puccinellia maritima*. A “?” near a dashed line indicates that the exact depth of this line is uncertain, a “?” after species names indicates that the presence of this species was hypothesized.

fractions. Soil pH was determined after diluting 5 g of soil in 25 mL of a 0.01 M CaCl₂ solution and electrical conductivity was measured after diluting 5 g of soil in 25 mL of deionized water. The collected biomass was dried at 60 °C for 48 h after sediments were removed and weighed in order to calculate the total dry weight of the biomass. The total above-ground biomass of one of the replicates collected on a 0.25 m² surface area was shredded and repeatedly divided into equal parts until only a small portion was left. This was further grinded and analysed for OC content, $\delta^{13}\text{C}$ and C : N ratio using the Elemental Analyser (FlashEA 1112 HT, Thermo Scientific). Soil bulk density samples were dried at 105 °C for 24 h. After soil bulk densities were calculated, the samples were washed over a 0.5 mm sieve using deionized water and all roots were collected. The roots were cleaned using deionized water, dried at 60 °C and weighed.

2.4 Data analysis

At every location one soil profile was analysed in detail (every other depth sample, i.e. 0–0.03, 0.06–0.09 m, ...). For all three replicate profiles every third sample was analysed (i.e. 0–0.03, 0.09–0.12 m, ...) to a depth of 0.72 m. Below this depth, samples were analysed every 0.18 m. Continuous depth profiles of OC percentage for layers of 0.01 m were obtained by linear interpolation, using the average OC percentage at the depths at which three replicates were anal-

Table 2. General characteristics of the soil profiles at the studied sites. Bulk density values are averages for the upper metre of soil, while soil pH and electrical conductivity were measured in the topsoil only.

	Bulk density (g cm ⁻³)	Soil pH	Electrical conductivity (dS cm ⁻¹)
Freshwater low	0.40 ± 0.07	7.47 ± 0.02	0.0271 ± 0.0009
Freshwater high	0.54 ± 0.04	7.35 ± 0.10	0.0262 ± 0.0007
Brackish water low	0.89 ± 0.06 ^a	7.70 ± 0.06	0.0389 ± 0.0048
Brackish water high	0.99 ± 0.06	7.49 ± 0.09	0.0365 ± 0.0023
Saltwater low	0.63 ± 0.07 ^b	7.93 ± 0.02	0.0959 ± 0.0021
Saltwater high	0.96 ± 0.11	7.87 ± 0.03	0.0113 ± 0.0010

^a Down to 0.7 m depth. ^b Down to 0.2 m depth.

used (i.e. every 0.09 m). Continuous depth profiles for bulk density were obtained in an identical way. These continuous depth profiles were then used to calculate total SOC stocks for a volume of soil with a surface area of 1 m² and a depth equal to the total sampling depth of the marsh sediments. Root biomass was measured at discrete depths as explained above. For every layer the total root biomass for a surface area of 1 m² was calculated by rescaling the average root biomass for the three replicates to the total volume of that soil layer. Linear interpolation between measurements at different depth intervals was used to calculate the total root density per surface area of 1 m². To test whether annual above-ground biomass production was significantly different between the sites a one-way analysis of variance was used in Matlab[®], after checking for normality (Anderson–Darling test) and homogeneity of variances (Levene’s test) with a level of significance of $p < 0.05$. For the other variables only three replicates were available so no reliable significance test could be performed.

3 Results

3.1 Soil characteristics

The studied tidal marsh soils are classified as tidalic Fluvisols with a silt loam grain size (detailed grain size data are provided in the Supplement). The maximum depth of marsh sediments at the different study sites varies between 0.2 and > 1.4 m (Table 1). The average bulk density ranges from 0.40 to 0.99 g cm⁻³ (detailed bulk density data are provided in the Supplement), and both the topsoil pH and electrical conductivity increase in the downstream direction, from freshwater to salt marshes (Table 2).

3.2 Vegetation biomass production

Based on the measured total annual biomass (Fig. S1 in the Supplement, Table S2) and reported values of both above- and below-ground annual turnover rates (Table S3), annual

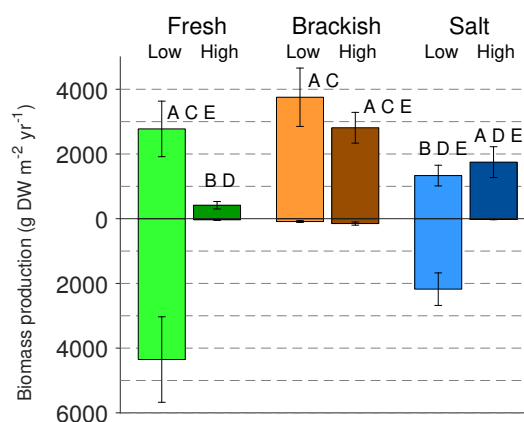


Figure 4. Annual biomass production (g dry weight m⁻² yr⁻¹), with upward pointing bars representing above-ground biomass production and downward pointing bars representing below-ground production (data are provided in Table S2). Standard deviations for above-ground biomass are calculated based on five replicates, for below-ground biomass on three replicas. Sample locations that do not share a letter have significantly ($p < 0.05$) different annual above-ground biomass production rates.

biomass production for the different sites was calculated, as shown in Fig. 4. In this figure, sample locations that do not share a letter have significantly different annual biomass production rates. The average annual above-ground biomass production is the highest for the brackish marshes, followed by the low freshwater marsh and both salt marshes. The high freshwater marsh has an above-ground biomass production that deviates from this pattern as a consequence of the fact that only fallen leaves of the willow trees were taken into account at this site, while standing willow vegetation could not be collected, so we underestimate total biomass production in this case. Upper limits for biomass production on this marsh may be deduced from other studies, which typically result in production rates of 500–1000 g dry weight m⁻² yr⁻¹ (Kopp et al., 2001). Although no clear pattern in the annual production of below-ground biomass along the estuary is observed, large differences between the sites are present (Fig. 4). Below-ground biomass production on the low freshwater marsh and the low salt marsh are 2 orders of magnitude larger compared to the other tidal marsh sites. At the former locations, most biomass is located below ground, while at the latter locations the majority of the vegetation biomass is located above ground.

3.3 Soil organic carbon depth profiles

The depth profiles of SOC show that the depth-averaged concentration decreases from freshwater to salt marshes, although the highest topsoil OC concentration is observed on the brackish marshes (Fig. 5, data on OC and C : N ratio are provided in the Supplement). In contrast to the freshwater soils, which show a gradual but limited decrease in OC con-

Table 3. Total organic carbon (OC) stock (kg OC m^{-2}) and standard deviations calculated for the full vertical sampling profiles (depths used for calculations are given in brackets) and the upper 0.6 m.

	OC stock (kg OC m^{-2})	
	Low marsh	High marsh
For the entire marsh profile		
Freshwater	32.35 ± 0.65 (1.2 m)	46.44 ± 0.80 (1.4 m)
Brackish water	20.50 ± 0.72 (0.75 m)	32.23 ± 0.31 (1.4 m)
Saltwater	2.84 ± 0.10 (0.2 m)	9.93 ± 0.34 (0.6 m)
Down to 0.6 m depth		
Freshwater	16.38 ± 0.54	21.66 ± 0.71
Brackish water	18.63 ± 0.71	19.63 ± 0.27
Saltwater	–	9.93 ± 0.34

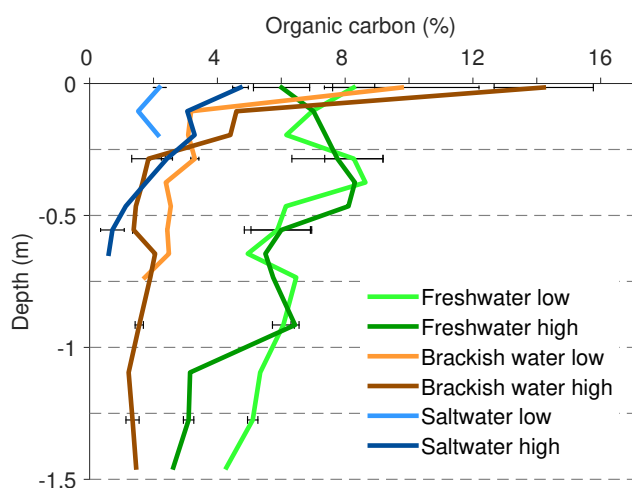


Figure 5. Depth profiles of OC concentration for all study sites. Data points show the average of three replicate soil samples. Error bars for specific depths represent the standard deviation.

centration with depth, the brackish and salt marshes show a sharp decrease in OC concentration in the upper 0.25 m.

3.4 Soil organic carbon inventories

The highest total SOC stocks are found in the freshwater marshes, followed by the brackish and salt marshes (Table 3). For every marsh, SOC stocks are greater for the high marsh compared to the low marsh, as a consequence of both deeper marsh soils and higher SOC concentrations. In order to compare the marshes directly to each other the stocks down to the largest common depth have been calculated (Table 3). Using this approach, freshwater and brackish marshes have comparable SOC stocks, while both locations on the salt marsh have significantly lower stocks. Depth profiles of cumulative SOC stocks per 0.01 m layer, after interpolation as explained in Sect. 2.4, are shown in Fig. S2.

3.5 Stable carbon isotopes

The depth profiles of stable OC isotopes ($\delta^{13}\text{C}$) are shown in Fig. 6, together with the $\delta^{13}\text{C}$ signal of above- and below-ground vegetation (data on $\delta^{13}\text{C}$ is provided in the Supplement). In general an increase in $\delta^{13}\text{C}$ values with depth is observed, although deviations from this pattern occur along the profiles. For all sites except the low salt marsh, which is characterized by *Spartina anglica* vegetation (C4 type), the $\delta^{13}\text{C}$ signal of SOC in the topsoil layer is similar to the $\delta^{13}\text{C}$ signal of standing vegetation. However, a close inspection shows differences. On the high freshwater marsh the topsoil $\delta^{13}\text{C}$ signal is higher than the signal for both above- and below-ground vegetation, while on the low freshwater marsh the topsoil $\delta^{13}\text{C}$ signal is lower than the above- and below-ground vegetation signal. On both the low and high brackish marshes, the topsoil $\delta^{13}\text{C}$ is very similar to the $\delta^{13}\text{C}$ signal of roots, while it is about 1‰ lower compared to the $\delta^{13}\text{C}$ signal of above-ground vegetation. On the high salt marsh, the topsoil $\delta^{13}\text{C}$ signal has a value in between the $\delta^{13}\text{C}$ signals of above- and below-ground vegetation, while on the low salt marsh the topsoil $\delta^{13}\text{C}$ signal is significantly lower compared to the signal of both above- and below-ground vegetation.

4 Discussion

4.1 Observed patterns in SOC storage

While our data do not allow for a full statistical or mechanistic analysis of the mechanisms controlling the long-term storage of SOC in the studied tidal marshes, some important observations can be made. A first observation is that low SOC stocks are not systematically related to low biomass production, as no statistical relationship between total annual biomass production (above and below ground) and SOC stocks is found ($R^2 = 0.01$, Fig. S3). For example, the annual biomass production on the low saltwater marsh (*Spartina an-*

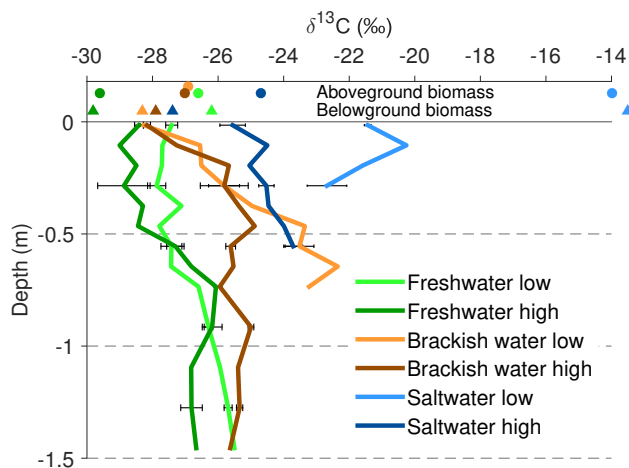


Figure 6. Depth profiles of $\delta^{13}\text{C}$, together with the $\delta^{13}\text{C}$ signal of above-ground (circles) and below-ground (triangles) biomass (values are provided in Table S2). Error bars represent the standard deviation.

glica) is relatively high (Fig. 4), while this site is characterized by the lowest SOC stocks. In addition, there is no relationship between annual root carbon production and SOC stocks ($R^2 = 0.004$, data not shown). This is rather surprising, as it has been proposed that roots contribute significantly to the subsoil OC pool in tidal marshes (Craft, 2007; Saintilan et al., 2013).

A second important observation is the very rapid decrease of SOC with depth at the brackish sites. This decrease is accompanied by a shift in $\delta^{13}\text{C}$ to less negative values with depth in the topsoil of these marshes, suggesting that on the brackish marshes a significant fraction of OC is rapidly decomposed after burial (Fig. 6). On the high brackish marsh the decline in SOC and the shift in $\delta^{13}\text{C}$ show the same tendency towards a depth of 0.3 m, while deeper along the profile, both variables remain approximately constant with depth. This indicates that a significant fraction (approx. 87 %) of deposited OC is decomposed in this top layer. In the low brackish marsh sediments the situation is different. Here the SOC concentration only decreases from the top of the profile down to a depth of 0.15 m, while the $\delta^{13}\text{C}$ signal increases throughout the profile. At this location *Spartina anglica* (a C4 plant) was possibly present during early marsh development, resulting in a more positive $\delta^{13}\text{C}$ signal (Boschker et al., 1999; Middelburg et al., 1997). Currently *Elymus athericus*, a C3 plant, is dominating the marsh vegetation. This implies that the shift in $\delta^{13}\text{C}$ with depth on the low brackish marsh could partly be the result of a shift from a C4 to C3 type vegetation, rather than resulting from the decomposition of SOC alone. This is very likely, as shifts in $\delta^{13}\text{C}$ as a consequence of kinetic fractionation during decomposition are generally of the order of 1–3 ‰ (Choi et al., 2001), while the shift we observe is much larger (ca. 5.7 ‰). However, the decrease in SOC together with the shift in $\delta^{13}\text{C}$

in the top 0.15 m suggests that, also on this marsh, significant decomposition of deposited OC (approx. 68 %) took place after burial. It should be noted that the $\delta^{13}\text{C}$ signal of the most recent sediments found on the intertidal areas can vary throughout the year (Zetsche et al., 2011). However, the $\delta^{13}\text{C}$ depth profiles we observe are an integration of these annual cycles, limiting the effect of the timing of sample collection on the observed depth profiles of $\delta^{13}\text{C}$.

On the high salt marsh a significant decrease of SOC concentration with depth also occurs. This is again accompanied by a shift in $\delta^{13}\text{C}$ towards more positive values with depth. This location is currently characterized by a mixture of C3 type vegetation. It is uncertain, however, whether the isotopic shift with depth can entirely be attributed to kinetic fractionation caused by OC decay. It is likely that at the beginning of marsh growth *Spartina anglica* was also present at this location, as it is currently present on the low part of this marsh. This would imply that, also at this location, the shift in $\delta^{13}\text{C}$ with depth is the result of a combination of decomposition of OC and a shift in vegetation from C4 to C3 type.

Our observations indicate that on both the salt and brackish marshes a significant fraction of OC is lost after burial. Although in the brackish marsh sediments a larger fraction of OC is lost after burial compared to salt marshes, total SOC stocks in the brackish marsh sediments are significantly higher compared to the salt marshes.

On the freshwater marshes the situation is different. In both the low and high freshwater marsh sediments the decline in OC concentration with depth is very limited. In addition, the $\delta^{13}\text{C}$ signal does not show a significant shift in the top 0.5 m of the soil profile. Below this depth there is a limited shift in $\delta^{13}\text{C}$ toward more positive values, but the interpretation of this pattern is complicated by the effect of previous land uses on the marsh (Fig. 3). These observations indicate that at both locations on the freshwater marsh there is limited decomposition of OC after burial.

4.2 Explanations for the observed patterns in soil organic carbon stocks

An explanation is needed for the differences in depth gradients in both SOC stocks and $\delta^{13}\text{C}$ between salt and brackish marshes on the one hand and freshwater marshes on the other hand. Several factors may contribute to these differences and their possible roles are discussed below.

4.2.1 Salinity

Although the Scheldt estuary is characterized by a strong salinity gradient (Van Damme et al., 2005), it is unlikely that salinity as such is a direct factor controlling the difference in decomposition of OC that we observed. If salinity directly controlled OC decomposition, it would necessitate a positive relationship between decomposition rate and salinity. However, litterbag experiments with *Elymus athericus* on a tidal

marsh in the Scheldt estuary showed that there was an inverse relationship between soil salinity and decomposition (Hemminga et al., 1991). In addition, Hemminga et al. (1991) concluded that there is no significant variation in cellulose decomposition in tidal marsh sediments along the brackish and saltwater portion of the Scheldt estuary.

4.2.2 Vegetation type

The type of vegetation present on the different marshes is another possible controlling factor, as it has been shown that the residues of different macrophytes have a different resistance against decomposition (Buth and de Wolf, 1985; Hemminga and Buth, 1991; Valery et al., 2004). One of the factors that determines the decomposition rate of plant material is the nitrogen content, whereby plant material with a higher C : N ratio is generally more resistant to decomposition (Hemminga and Buth, 1991; Jones et al., 2016; Webster and Benfield, 1986). The C : N ratio of the vegetation present on the salt marsh (values between 27 and 30) is significantly lower compared to the vegetation present on the brackish and freshwater marshes (values between 33 and 55; Table S2). However, our OC and $\delta^{13}\text{C}$ profiles suggest that decomposition rates are highest on the brackish marshes and lowest on the freshwater marshes, while the vegetation present at these locations has comparable C : N ratios. Thus, there does not appear to be a direct relationship between the C : N ratio of the biomass and SOC decomposition.

Although our data do not allow us to isolate the effect of vegetation type on SOC stocks along the estuary, some important observations can be made. Firstly, the low and high freshwater marsh have different vegetation types (*P. australis* and *Salix* resp.). However, both soils show a similar SOC profile. In addition, the high freshwater marsh, where annual biomass production is significantly lower, has the largest SOC stock. This indicates that the effect of local biomass production on SOC stock is limited in the freshwater marshes. Secondly, both brackish marshes have the same vegetation type (*E. athericus*), while topsoil OC concentrations and total SOC stocks on the high marsh are larger compared to the low marsh. In addition, the high marsh is characterized by a somewhat lower annual biomass production (although the difference is not statistically significant). Thus, variations in local biomass production do not explain the differences in SOC stock between young and old brackish marshes either. The presence of *Spartina anglica* on the low salt marsh is indeed likely to be responsible for the low SOC stocks. While *Spartina anglica* is characterized by a high net primary productivity, the organic material produced is known to be very labile (Boschker et al., 1999; Bouillon and Boschker, 2006; Middelburg et al., 1997). Taken together, these observations indicate that local biomass production is not likely to be a dominant factor controlling overall variations in SOC stocks along the estuary as variations in OC stocks both along the salinity gradient of the estuary and

between old and young marshes at a given salinity level cannot be explained by variations in biomass production. The effect of *Spartina anglica* on the SOC stock of the low salt marsh shows, however, that in some cases the presence of a certain vegetation type may be a dominant factor.

4.2.3 Allochthonous organic carbon inputs along the estuary

The OC that is present in tidal marsh sediments is not only derived from autochthonous biomass. Estuaries are often characterized by relatively high concentrations of suspended sediment to which a significant amount of particulate organic carbon (POC) is associated (Abril et al., 2002). Due to the long residence time of water in the Scheldt estuary (2–3 months, Soetaert and Herman, 1995), organic matter is intensively processed as it moves through the estuary (Abril et al., 2002; Middelburg and Herman, 2007). In addition, mixing between fluvial and marine particles takes place (Nolting et al., 1999; Regnier and Wollast, 1993). Overall, this leads to significant variations in both the quantity and the quality of the POC that is present in the water and deposited on the marshes. Clearly, this variation may not only affect the magnitude of the OC inputs but also the decomposability of the OC that is deposited.

The freshwater marshes are located near the upstream border of the Scheldt estuary close to the maximum turbidity zone (MTZ), with average suspended sediment concentrations of ca. 0.15 g L^{-1} (Van Damme et al., 2001; Temmerman et al., 2004). The suspended sediments in this zone contain 7–10 % POC (Abril et al., 2002). The higher values are observed in summer, when phytoplankton growth is important, while the lower values are reported in winter. The POC that is present in winter may be assumed to be processed POC of terrestrial origin (Hellings et al., 1999). In addition, during the past decades a large fraction of OC that has entered the freshwater portion of the estuary originated from untreated wastewater from the city of Brussels (Abril et al., 2002; Billen et al., 2005). It has, however, been shown that this OC is mineralized on a timescale of weeks, possibly even before it enters the estuary (Muylaert et al., 2005; Servais et al., 1987).

Sediment concentrations strongly decline downstream of the MTZ (Abril et al., 2002; Van Damme et al., 2005). At the location of the brackish and salt marshes (ca. 20 and ca. 50 km from the mouth) sediment concentrations are about 0.05 g L^{-1} (Van Damme et al., 2001; Temmerman et al., 2004). Furthermore, the POC content of these sediments decreases systematically in the downstream direction, except during the spring season when local production of OC due to phytoplankton is important in the marine part of the estuary (Abril et al., 2002). As a result, average POC concentrations vary between 4 and 6 % in the brackish water zone and between 2 and 5 % in the saltwater zone (Abril et al., 2002). The overall decline in POC content is not only explained by

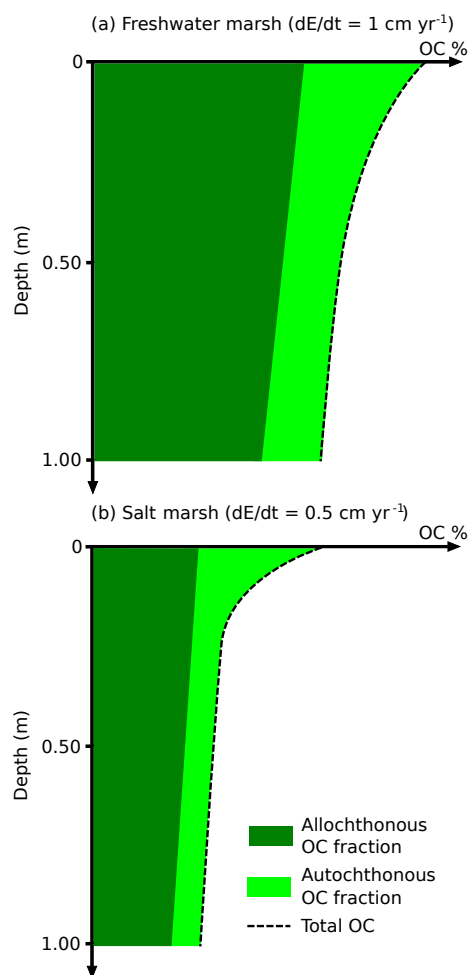


Figure 7. Conceptual diagram of the effect of both sediment deposition rate (dE/dt , E = elevation) and the relative inputs of recalcitrant allochthonous organic carbon and labile autochthonous organic carbon on the fate of buried OC in a tidal freshwater marsh (a) and salt marsh (b).

the progressive downstream mineralization of OC but also by the upstream transport of marine sediments that carry less POC.

The variations in both suspended sediment concentration and POC content have important consequences for the relative importance of allochthonous OC input on the marshes. On the freshwater marshes, both the high suspended sediment concentration and high POC loadings lead to a combination of high sedimentation rates ($10\text{--}20 \text{ mm yr}^{-1}$, with the highest sedimentation rates on the young marshes; Temmerman et al., 2004) and high inputs of allochthonous POC. On the saltwater marshes, sedimentation rates are much lower ($5\text{--}10 \text{ mm yr}^{-1}$; Temmerman et al., 2004) and the deposited sediments contain 50–70% less OC than the sediments deposited on the freshwater marsh (Abril et al., 2002). Evidently, these differences may have important effects on OC storage in tidal marsh sediments (Fig. 7). It can reasonably be

assumed that the allochthonous POC that is deposited with the sediments on the freshwater marsh consists of a large fraction of terrestrial, recalcitrant POC. This POC may be expected to have a high burial efficiency (i.e. it will decompose relatively slowly after burial) and will remain in the sediments for a considerable time. The local, autochthonous POC is fresh and will therefore be less recalcitrant. As a consequence, it may be expected to decompose much more rapidly with time and contribute much less to long-term OC storage. The latter explains why variations in biomass production and vegetation type on the marshes (both now and in the past, Fig. 3) do not explain variations in SOC storage in different marsh environments. Furthermore, the decomposition rate of both autochthonous and allochthonous POC can be expected to be inversely related to the burial rate as rapid sedimentation will protect OC from decomposition (Hartnett et al., 1998; Wang et al., 2014). Thus, OC will be better preserved when sedimentation rates are high.

Figure 7 illustrates how these factors combine. One may indeed expect to find a much less steep decline of the OC content with depth on the freshwater marsh (Fig. 7a) due to (1) the dominance of allochthonous, recalcitrant OC and (2) the rapid burial of OC. Furthermore, a relatively large fraction of labile autochthonous OC may be preserved, as it is advected rapidly to deep sediment layers. On the salt and brackish marshes a low sedimentation rate combines with low OC contents of the deposited sediments (Fig. 7b). As a consequence, autochthonous OC is a dominant input, but this OC decomposes rapidly with depth. This results in a significant decline of OC content with depth, combined with a significant increase in $\delta^{13}\text{C}$ due to kinetic isotopic fractionation. In a recent study, Hansen et al. (2016) also attributed decreasing SOC stocks with increasing salinity in the Elbe estuary (Germany) to a decreasing OC content of suspended sediments and decreasing macrophyte biomass with increasing salinity.

Thus, both sedimentation rate as well as the rate of allochthonous OC input to the marsh system appear to place important controls on OC preservation in marsh sediments. While other factors such as local biomass production and salinity gradients may also be locally important, they do not appear to be key controls in the Scheldt estuary as most autochthonous POC appears to decompose rapidly, independent of the specific environmental conditions. This finding is similar to the observations of Omengo et al. (2016), who found that the OC preserved at depth in floodplain sediments of the Tana River in Kenya consisted dominantly of processed OC that was deposited by the river, while locally produced OC contributed little to long-term OC preservation.

4.3 Effect of sampling procedure on reported estuarine OC stocks

The results of this study show that both SOC concentrations and stocks of tidal marshes vary significantly along a tem-

Table 4. Reported SOC stocks (kg OC m⁻²) of tidal marsh soils along estuarine salinity gradients.

Estuary	Sampling depth (m)	Freshwater	Oligohaline	Mesohaline	Polyhaline	Reference
Delaware, USA	0.16	7.034	2.41	3.528	–	Weston et al. (2014)
Sapelo, Doboy, Altamaha Georgia, USA	0.30	8.379	10.692	4.626	5.932	Craft (2007)
Dovey, Wales	0.10	–	2.8	1.8	2.4 (high), 1.4 (low)	Dausse et al. (2012)
Barataria Louisiana, USA	0.38	10.3	9.2	12.9	12.8	Hatton et al. (1983)
Satilla Altamaha Ogeechee Georgia, USA	0.30	8.096 ± 1.245	–	6.816 ± 0.997	6.069 ± 0.482	Wieski et al. (2010)
Barataria basin Louisiana, USA	0.50	5.37	–	4.38	2.90	Williams and Rosenheim (2015)
San Francisco Bay California, USA	0.20	–	–	7.82	5.33	Callaway et al. (2012) ^a
Louisiana, USA	1.5	65.76	–	–	56.65	Wang et al. (2011)
Elbe, Germany	1.0	–	27.05	16.04	11.31	Hansen et al. (2016) ^b
Scheldt	0.6	–	21.66 ± 0.71	19.63 ± 0.27	9.93 ± 0.34	This study ^a
Belgium, the Netherlands						

^a Data for high marshes only. ^b Average for all unmanaged sites.

perate estuary, with freshwater marshes having the highest stocks, followed by brackish and salt marshes (Fig. 5 and Table 3). This tendency is in agreement with observations in other studies (Table 4). However, the differences reported in previous studies are almost always much smaller than the differences we find. As the estuaries reported in Table 4 cover a large geographical range, differences in environmental conditions will have an influence on the reported SOC stocks. For example, the estuaries reported in Table 4 that are located along the southern coast of the USA experience significantly higher average temperatures compared to the Scheldt estuary. In addition, the vegetation species present on the tidal marshes, as well as differences in regional geology and estuarine morphology, will play a role. However, as the studies listed in Table 4 report SOC stocks in tidal marsh sediments along a salinity gradient, similarities in the factors controlling these stocks are also present. For example, the freshwater marshes will receive considerably more OC from terrestrial sources, while the influence of OC inputs from marine sources will be the largest in the salt marshes. Moreover, macrophyte production is generally considerably higher at freshwater marshes compared to salt marshes (e.g. Dausse et al., 2012; Hansen et al., 2016).

In addition to these factors, differences in sampling procedures can also explain some discrepancies. In most studies, marshes were sampled to a limited depth (Table 4). Generally, the differences in SOC content between different marshes reported in Table 4 are smallest for the top layers and increase with depth. As a consequence, the difference in SOC inventory will increase if a larger sampling depth is considered. Evidently, considering a larger sampling depth will also lead to higher estimates of SOC stocks. This is one of the factors explaining why our stock estimates are generally much higher than those reported in the other studies in

Table 4, especially for the freshwater marshes, and why we find larger differences in total SOC stocks between different marshes.

It is important to consider whether carbon stocks should be compared by calculating stocks down to a certain depth or whether the total stock present in the marsh sediments should be taken into account. While it is simpler and more transparent to consider a certain depth, this approach does not account for the differences in dynamics between marshes. As Fig. 2 shows, marsh accumulation rates are significantly higher for the freshwater marshes. This automatically implies that, when different marshes are sampled to a common depth, the time frame that is accounted for will be shorter for those marshes that have the highest accumulation rates (Elschot et al., 2015). We suggest that the establishment of a correct time frame, from which sedimentation rates and their variations over time can be deduced, is indispensable for a correct interpretation of differences in SOC stocks (as well as C sequestration rates) between marshes.

4.4 Implications of sea level rise for estuarine soil organic carbon stocks

As global sea level is predicted to continue to rise during the next centuries, progressive intrusion of saltwater further into estuaries may be expected (Robins et al., 2016; Ross et al., 2015). As it is shown that freshwater and brackish tidal marshes store more SOC compared to salt marshes (Table 3), one may expect that this will lead to a decrease in OC sequestration at locations where brackish marshes are replaced by salt marshes. Furthermore, the MTZ is predicted to shift more inland (Robins et al., 2016). Because the Scheldt estuary is completely embanked and the tidal wave is stopped by sluices in the city of Ghent, the total area of freshwater

marshes is likely to decline after sea level rise (Barendregt and Swarth, 2013). As we have shown that SOC sequestration rates are the largest in the freshwater portion of the estuary, the amount of OC sequestration in the estuary is therefore likely to decline with sea level rise. Moreover, as a consequence of the upstream migration of the MTZ, terrestrial organic matter will not be able to travel as far downstream in the estuary. Therefore, tidal marshes which are now located at the downstream end of the MTZ will receive less stable terrestrial OC in the future. On the other hand, overall sedimentation rates are expected to increase with a rising sea level, which will automatically lead to an increase in the rate of OC deposition as well as of OC burial rates, resulting in an increase of the OC sequestration rate per unit surface area.

Saltwater intrusion can also influence the decomposition of previously sequestered OC, with some studies concluding that saltwater intrusion will enhance decomposition of organic matter (Craft, 2007; Morrissey et al., 2014; Weston et al., 2006, 2011), while others find that decomposition rates will decrease (Hemminga et al., 1991). From these studies and from the analysis by Chambers et al. (2011), it is clear that this effect is highly dependent on local factors, such as the concentration of elements in the sea water that intrudes the estuary. Therefore, no reliable estimation of the impact of saltwater intrusion on OC mineralization in the Scheldt estuary can be made.

The above illustrates that our current understanding of the future evolution of the Scheldt estuary is still insufficient to make a quantitative assessment of how SOC stocks in the tidal marsh environment may change in the future.

5 Conclusions

As reported data on estuarine gradients of SOC are very scarce and, more importantly, often based on shallow soil sampling, additional research is needed to better constrain estimates of global estuarine OC stocks. This study shows that the quantification of SOC stocks in tidal marsh sediments critically depends on the sampling depth. Gradients in SOC concentrations with depth strongly vary between marsh types, so a full inventory can only be made if sampling is carried out over the entire depth of the marsh sediments. Even if such data are available, interpretation has to be done with care, as sedimentation rates may vary considerably within a single estuary, making it complex to convert inventories to sedimentation or preservation rates. In the Scheldt estuary, total SOC stocks are largest in a freshwater and brackish tidal marsh and significantly lower in a saltwater marsh. These variations are to some extent controlled by variations in autochthonous biomass production, but our data strongly suggest that the key control on long-term OC preservation is the relative contribution of terrestrial, allochthonous OC to the total OC input, while OC burial rate may also be important. The impact of future sea level rise on OC stocks in

tidal marsh sediments will be determined by an interplay of different factors, including the evolution of the spatial extent of marshes in different salinity zones and sediment and OC deposition rates. Our study allowed us to identify the factors that are important controls on OC storage but further research may be needed to resolve this issue.

6 Data availability

The following data are provided in the Supplement: depth profiles of (i) bulk density, (ii) OC, $\delta^{13}\text{C}$ and CN values and (iii) grain size in Excel format and (iv) data on vegetation characteristics and sample locations in pdf format.

The Supplement related to this article is available online at doi:10.5194/bg-13-6611-2016-supplement.

Competing interests. The authors declare that they have no conflict of interest.

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References

- Abril, G., Nogueira, M., Etcheber, H., Cabeçadas, G., Lemaire, E., and Brogueira, M.: Behaviour of Organic Carbon in Nine Contrasting European Estuaries, *Estuar. Coast. Shelf Sci.*, 54, 241–262, doi:10.1006/ecss.2001.0844, 2002.
- Barendregt, A. and Swarth, C. W.: Tidal Freshwater Wetlands: Variation and Changes, *Estuar. Coast.*, 36, 445–456, doi:10.1007/s12237-013-9626-z, 2013.
- Billen, G., Garnier, J., and Rousseau, V.: Nutrient fluxes and water quality in the drainage network of the Scheldt basin over the last 50 years, *Hydrobiologia*, 540, 47–67, doi:10.1007/s10750-004-7103-1, 2005.
- Boschker, H. T. S., de Brouwer, J. F. C., and Capenberg, T. E.: The contribution of macrophyte-derived organic matter to microbial biomass in salt-marsh sediments: Stable carbon isotope analysis of microbial biomarkers, *Limnol. Oceanogr.*, 44, 309–319, doi:10.4319/lo.1999.44.2.0309, 1999.

- Bouillon, S. and Boschker, H. T. S.: Bacterial carbon sources in coastal sediments: a cross-system analysis based on stable isotope data of biomarkers, *Biogeosciences*, 3, 175–185, doi:10.5194/bg-3-175-2006, 2006.
- Bridgman, S. D., Megonigal, J. P., Keller, J. K., Bliss, N. B., and Trettin, C.: The carbon balance of North American wetlands, *Wetlands*, 26, 889–916, 2006.
- Buth, G. J. C., and de Wolf, L.: Decomposition of *Spartina anglica*, *Elytrigia pungens* and *Halimione portulacoides* in a Dutch salt marsh in association with faunal and habitat influences, *Vegetatio*, 62, 337–355, doi:10.1007/BF00044761, 1985.
- Callaway, J. C., Borgnis, E. L., Turner, R. E., and Milan, C. S.: Carbon Sequestration and Sediment Accretion in San Francisco Bay Tidal Wetlands, *Estuar. Coast.*, 35, 1163–1181, doi:10.1007/s12237-012-9508-9, 2012.
- Chambers, L. G., Reddy, K. R., and Osborne, T. Z.: Short-Term Response of Carbon Cycling to Salinity Pulses in a Freshwater Wetland, *Soil Sci. Soc. Am. J.*, 75, 2000, doi:10.2136/sssaj2011.0026, 2011.
- Cherry, J. A., McKee, K. L., and Grace, J. B.: Elevated CO₂ enhances biological contributions to elevation change in coastal wetlands by offsetting stressors associated with sea-level rise, *J. Ecol.*, 97, 67–77, doi:10.1111/j.1365-2745.2008.01449.x, 2009.
- Chmura, G. L., Anisfeld, S. C., Cahoon, D. R., and Lynch, J. C.: Global carbon sequestration in tidal, saline wetland soils, *Global Biogeochem. Cy.*, 17, 22-1–22-12, doi:10.1029/2002gb001917, 2003.
- Choi, Y., Hsieh, Y., and Wang, Y.: Vegetation succession and carbon sequestration in a coastal wetland in northwest Florida: Evidence from carbon isotopes, *Global Biogeochem. Cy.*, 15, 311–319, 2001.
- Craft, C.: Freshwater input structures soil properties, vertical accretion, and nutrient accumulation of Georgia and U.S. tidal marshes, *Limnol. Oceanogr.*, 52, 1220–1230, doi:10.4319/lo.2007.52.3.1220, 2007.
- Dausse, A., Garbutt, A., Norman, L., Papadimitriou, S., Jones, L. M., Robins, P. E., and Thomas, D. N.: Biogeochemical functioning of grazed estuarine tidal marshes along a salinity gradient, *Estuar. Coast. Shelf Sci.*, 100, 83–92, doi:10.1016/j.ecss.2011.12.037, 2012.
- De Leeuw, J., Olf, H., and Bakker, J. P.: Year-to-Year variation in peak above-ground biomass of six salt-marsh angiosperm communities as related to rainfall deficit and inundation frequency, *Aquat. Bot.*, 36, 139–151, 1990.
- Donato, D. C., Kauffman, J. B., Murdiyarso, D., Kurnianto, S., Stidham, M., and Kanninen, M.: Mangroves among the most carbon-rich forests in the tropics, *Nat. Geosci.*, 4, 293–297, doi:10.1038/ngeo1123, 2011.
- Duarte, C. M., Losada, I. J., Hendriks, I. E., Mazarrasa, I., and Marbà, N.: The role of coastal plant communities for climate change mitigation and adaptation, *Nature Climate Change*, 3, 961–968, doi:10.1038/nclimate1970, 2013.
- Elschot, K., Bakker, J. P., Temmerman, S., Van De Koppel, J., and Bouma, T. J.: Ecosystem engineering by large grazers enhances carbon stocks in a tidal salt marsh, *Mar. Ecol.-Prog. Ser.*, 537, 9–21, doi:10.3354/meps11447, 2015.
- Fagherazzi, S., Kirwan, M. L., Mudd, S. M., Guntenspergen, G. R., Temmerman, S., Rybczyk, J. M., Reyes, E., Craft, C., and Clough, J.: Numerical models of salt marsh evolution: Ecological, geomorphic, and climatic factors, *Rev. Geophys.*, 50, 1–28, doi:10.1029/2011RG000359, 2012.
- Govers, G., Merckx, R., Van Oost, K., and van Wesemael, B.: Managing Soil Organic Carbon for Global Benefits: A STAP Technical Report, Global Environmental Facility, Washington, D.C., 2013.
- Groenendijk, A. M.: Primary production of 4 dominant salt-marsh angiosperms in the southwestern Netherlands, *Vegetatio*, 57, 143–152, 1984.
- Hansen, K., Butzeck, C., Eschenbach, A., Gröngroft, A., Jensen, K., and Pfeiffer, E. M.: Factors influencing the organic carbon pools in tidal marsh soils of the Elbe estuary (Germany), *J. Soils Sediments*, 1–14, doi:10.1007/s11368-016-1500-8, 2016.
- Hartnett, H. E., Keil, R. G., Hedges, J. I., and Devol, A. H.: Influence of oxygen exposure time on organic carbon preservation in continental margin sediments, *Nature*, 391, 572–574, 1998.
- Hatton, R. S., Delaune, R. D., and Patrick, W. H. J.: Sedimentation, accretion, and subsidence in marshes of Barataria Basin, Louisiana, *Limnol. Oceanogr.*, 28, 494–502, doi:10.4319/lo.1983.28.3.0494, 1983.
- Hellings, L., Dehairs, F., Tackx, M., Keppens, E., and Baeyens, W.: Origin and fate of organic carbon in the freshwater part of the Scheldt Estuary as traced by stable carbon isotope composition, *Biogeochemistry*, 47, 167–186, 1999.
- Hemminga, M. A. and Buth, G. J. C.: Decomposition in Salt Marsh Ecosystems of the S.W. Netherlands: The Effects of Biotic and Abiotic Factors, *Vegetatio*, 92, 73–83, 1991.
- Hemminga, M. A., De Leeuw, J., De Munck, W., and Koutstaal, B. P.: Decomposition in estuarine salt marshes: the effect of soil salinity and soil water content, *Vegetatio*, 94, 25–33, doi:10.1007/BF00044913, 1991.
- Jones, J. A., Cherry, J. A., and McKee, K. L.: Species and tissue type regulate long-term decomposition of brackish marsh plants grown under elevated CO₂ conditions, *Estuar. Coast. Shelf Sci.*, 169, 38–45, doi:10.1016/j.ecss.2015.11.033, 2016.
- Kirwan, M. L. and Blum, L. K.: Enhanced decomposition offsets enhanced productivity and soil carbon accumulation in coastal wetlands responding to climate change, *Biogeosciences*, 8, 987–993, doi:10.5194/bg-8-987-2011, 2011.
- Kirwan, M. L. and Mudd, S. M.: Response of salt-marsh carbon accumulation to climate change, *Nature*, 489, 550–553, doi:10.1038/nature11440, 2012.
- Kopp, R. F., Abrahamson, L. P., White, E. H., Volk, T. A., Nowak, C. A., and Fillhart, R. C.: Willow biomass production during ten successive annual harvests, *Biomass Bioenerg.*, 20, 1–7, doi:10.1016/S0961-9534(00)00063-5, 2001.
- McLeod, E., Chmura, G. L., Bouillon, S., Salm, R., Björk, M., Duarte, C. M., Lovelock, C. E., Schlesinger, W. H., and Silliman, B. R.: A blueprint for blue carbon: toward an improved understanding of the role of vegetated coastal habitats in sequestering CO₂, *Front. Ecol. Environ.*, 9, 552–560, doi:10.1890/110004, 2011.
- Meire, P., Ysebaert, T., Van Damme, S., Van Den Bergh, E., Maris, T., and Struyf, E.: The Scheldt estuary: A description of a changing ecosystem, *Hydrobiologia*, 540, 1–11, doi:10.1007/s10750-005-0896-8, 2005.
- Middelburg, J. J. and Nieuwenhuize, J.: Carbon and nitrogen stable isotopes in suspended matter and sediments from the

- Schelde Estuary, *Mar. Chem.*, 60, 217–225, doi:10.1016/S0304-4203(97)00104-7, 1998.
- Middelburg, J. J. and Herman, P. M. J.: Organic matter processing in tidal estuaries, *Mar. Chem.*, 106, 127–147, doi:10.1016/j.marchem.2006.02.007, 2007.
- Middelburg, J. J., Nieuwenhuize, J., Lubberts, R. K., van de Plassche, O., and Vandeplassche, O.: Organic carbon isotope systematics of coastal marshes, *Estuar. Coast. Shelf Sci.*, 45, 681–687, doi:10.1006/ecss.1997.0247, 1997.
- Morrissey, E. M., Gillespie, J. L., Morina, J. C., and Franklin, R. B.: Salinity affects microbial activity and soil organic matter content in tidal wetlands, *Glob. Change Biol.*, 20, 1351–1362, doi:10.1111/gcb.12431, 2014.
- Muylaert, K., Dasseville, R., De Brabandere, L., Dehairs, F., and Vyverman, W.: Dissolved organic carbon in the freshwater tidal reaches of the Schelde estuary, *Estuar. Coast. Shelf Sci.*, 64, 591–600, doi:10.1016/j.ecss.2005.04.010, 2005.
- Nelleman, C., Corcoran, E., Duarte, E., Valdés, L., De Young, C., Fonseca, L., and Grimsditch, G. (Eds.): *Blue carbon*, United Nations Environment Programme, GRID-Arendal, 2009.
- Neubauer, S. C.: Contributions of mineral and organic components to tidal freshwater marsh accretion, *Estuar. Coast. Shelf Sci.*, 78, 78–88, doi:10.1016/j.ecss.2007.11.011, 2008.
- Nolting, R. F., Helder, W., De Baar, H. J. W., and Gerringa, L. J. A.: Contrasting behaviour of trace metals in the Scheldt estuary in 1978 compared to recent years, *J. Sea Res.*, 42, 275–290, doi:10.1016/S1385-1101(99)00036-2, 1999.
- Omengo, F. O., Geeraert, N., Bouillon, S., and Govers, G.: Deposition and fate of organic carbon in floodplains along a tropical semiarid lowland river (Tana River, Kenya), *J. Geophys. Res.-Biogeo.*, 121, 1131–1143, doi:10.1002/2015JG003288, 2016.
- Pendleton, L., Donato, D. C., Murray, B. C., Crooks, S., Jenkins, W. A., Sifleet, S., Craft, C., Fourqurean, J. W., Kauffman, J. B., Marbà, N., Megonigal, P., Pidgeon, E., Herr, D., Gordon, D., and Baldera, A.: Estimating Global “Blue Carbon” Emissions from Conversion and Degradation of Vegetated Coastal Ecosystems, *PLoS One*, 7, e43542, doi:10.1371/journal.pone.0043542, 2012.
- Poffenbarger, H. J., Needelman, B. A., and Megonigal, J. P.: Salinity influence on methane emissions from tidal marshes, *Wetlands*, 31, 831–842, doi:10.1007/s13157-011-0197-0, 2011.
- Regnier, P. and Wollast, R.: Distribution of Trace-Metals in Suspended Matter of the Scheldt Estuary, *Mar. Chem.*, 43, 3–19, doi:10.1016/0304-4203(93)90212-7, 1993.
- Robins, P. E., Skov, M. W., Lewis, M. J., Giménez, L., Davies, A. G., Malham, S. K., Neill, S. P., McDonald, J. E., Whitton, T. A., Jackson, S. E., and Jago, C. F.: Impact of climate change on UK estuaries: A review of past trends and potential projections, *Estuar. Coast. Shelf Sci.*, 169, 119–135, doi:10.1016/j.ecss.2015.12.016, 2016.
- Rocha, A. V. and Goulden, M. L.: Why is marsh productivity so high? New insights from eddy covariance and biomass measurements in a Typha marsh, *Agr. Forest Meteorol.*, 149, 159–168, doi:10.1016/j.agrformet.2008.07.010, 2009.
- Ross, A. C., Najjar, R. G., Li, M., Mann, M. E., Ford, S. E., and Katz, B.: Sea-level rise and other influences on decadal-scale salinity variability in a coastal plain estuary, *Estuar. Coast. Shelf Sci.*, 157, 79–92, doi:10.1016/j.ecss.2015.01.022, 2015.
- Saintilan, N., Rogers, K., Mazumder, D., and Woodroffe, C.: Allochthonous and autochthonous contributions to carbon accumulation and carbon store in southeastern Australian coastal wetlands, *Estuar. Coast. Shelf Sci.*, 128, 84–92, doi:10.1016/j.ecss.2013.05.010, 2013.
- Scharlemann, J. P., Tanner, E. V., Hiederer, R., and Kapos, V.: Global soil carbon: understanding and managing the largest terrestrial carbon pool, *Carbon Manag.*, 5, 81–91, doi:10.4155/cmt.13.77, 2014.
- Servais, P., Billen, G., and M.-C., H.: Determination of the biodegradable fraction of dissolved organic matter in waters, *Water Res.*, 21, 445–450, 1987.
- Soetaert, K. and Herman, P. M. J.: Estimating estuarine residence times in the Westerschelde (The Netherlands) using a box model with fixed dispersion coefficients, *Hydrobiologia*, 311, 215–224, doi:10.1007/BF00008582, 1995.
- Temmerman, S., Govers, G., Meire, P., and Wartel, S.: Modelling long-term tidal marsh growth under changing tidal conditions and suspended sediment concentrations, Scheldt estuary, Belgium, *Mar. Geol.*, 193, 151–169, doi:10.1016/S0025-3227(02)00642-4, 2003.
- Temmerman, S., Govers, G., Wartel, S., and Meire, P.: Modelling estuarine variations in tidal marsh sedimentation: response to changing sea level and suspended sediment concentrations, *Mar. Geol.*, 212, 1–19, doi:10.1016/j.margeo.2004.10.021, 2004.
- Tolman, M. E. and Pranger, D. P.: *Toelichting bij de Vegetatiekartering Westerschelde 2010*, Rijkswaterstaat, Ministerie van Verkeer en Waterstaat, Delft, 2012.
- Valery, L., Bouchard, V., and Lefevre, J. C.: Impact of the invasive native species *Elymus athericus* on carbon pools in a salt marsh, *Wetlands*, 24, 268–276, doi:10.1672/0277-5212(2004)024[0268:IOTINS]2.0.CO;2, 2004.
- Van Braeckel, A., Coen, L., Peeters, P., Plancke, Y., Mikkelsen, J., and Van den Bergh, E.: *Historische evolutie van Zeescheldebhabitats*, Kwantitatieve en kwalitatieve analyse van invloedsfactoren. Rapporten van het Instituut voor Natuur- en Bosonderzoek (59), Instituut voor Natuur- en Bosonderzoek, Brussel i.s.m. het Waterbouwkundig Laboratorium, Antwerpen, 2012.
- Van Damme, S., De Winder, B., Ysebaert, T., and Meire, P.: Het “bijzondere” van de Schelde: de abiotiek van het Schelde-estuarium, *Levende Nat.*, 102, 37–39, 2001.
- Van Damme, S., Struyf, E., Maris, T., Ysebaert, T., Dehairs, F., Tackx, M., Heip, C., and Meire, P.: Spatial and temporal patterns of water quality along the estuarine salinity gradient of the Scheldt estuary (Belgium and The Netherlands): results of an integrated monitoring approach, *Hydrobiologia*, 540, 29–45, doi:10.1007/s10750-004-7102-2, 2005.
- Wang, J. J., Dodla, S. K., Delaune, R. D., Hudnall, W. H., and Cook, R. L.: Soil carbon characteristics in two Mississippi river deltaic Marshland profiles, *Wetlands*, 31, 157–166, doi:10.1007/s13157-010-0130-y, 2011.
- Wang, Z., Van Oost, K., and Govers, G.: Predicting the long-term fate of buried organic carbon in colluvial soils, *Global Biogeochem. Cy.*, 29, 65–79, doi:10.1002/2014GB004912, 2014.
- Webster, J. R. and Benfield, E. F.: Vascular Plant Breakdown in Freshwater Ecosystems, *Annu. Rev. Ecol. Syst.*, 17, 567–594, 1986.
- Weston, N. B., Dixon, R. E., and Joye, S. B.: Ramifications of increased salinity in tidal freshwater sediments: Geochemistry and microbial pathways of organic matter mineralization, *J. Geophys. Res.-Biogeo.*, 111, 1–14, doi:10.1029/2005JG000071, 2006.

- Weston, N. B., Vile, M. A., Neubauer, S. C., and Velinsky, D. J.: Accelerated microbial organic matter mineralization following salt-water intrusion into tidal freshwater marsh soils, *Biogeochemistry*, 102, 135–151, doi:10.1007/s10533-010-9427-4, 2011.
- Weston, N. B., Neubauer, S. C., Velinsky, D. J., and Vile, M. A.: Net ecosystem carbon exchange and the greenhouse gas balance of tidal marshes along an estuarine salinity gradient, *Biogeochemistry*, 120, 163–189, doi:10.1007/s10533-014-9989-7, 2014.
- Whigham, D. F.: Primary Production in Tidal Freshwater Wetlands, in *Tidal freshwater wetlands*, edited by: Barendregt, A., Whigham, D., and Baldwin, A., 320 pp., Margraf Publishers GmbH, 2009.
- Wieski, K., Guo, H. Y., Craft, C. B., and Pennings, S. C.: Ecosystem Functions of Tidal Fresh, Brackish, and Salt Marshes on the Georgia Coast, *Estuar. Coast.*, 33, 161–169, doi:10.1007/s12237-009-9230-4, 2010.
- Williams, E. K. and Rosenheim, B. E.: What happens to soil organic carbon as coastal marsh ecosystems change in response to increasing salinity? An exploration using ramped pyrolysis, *Geochem. Geophys. Geosy.*, 16, 2322–2335, doi:10.1002/2015GC005839, 2015.
- Woodwell, G. M., Rich, P. H., and Mall, C. S. A.: Carbon in estuaries, in *Carbon and the Biosphere*, edited by: Woodwell, G. M. and Pecari, E. V., US AEC, 1973.
- Zetsche, E., Thornton, B., Midwood, A. J., and Witte, U.: Utilization of different carbon sources in a shallow estuary identified through stable isotope techniques, *Cont. Shelf Res.*, 31, 832–840, doi:10.1016/j.csr.2011.02.006, 2011.