Factors controlling spatiotemporal variability of soil carbon accumulation and stock estimates in a tidal salt marsh

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Abstract

Tidal salt marshes are important contributors to soil carbon (C) stocks despite their relatively small land surface area. Although it is well understood that salt marshes have soil C burial rates orders of magnitude greater than those of terrestrial ecosystems, there is a wide range in ~~storage~~accrual rates among spatially distributed marshes. In addition, wide ranges in C ~~storage~~accrual rates also exist within a single marsh ecosystem. Tidal marshes often contain multiple species of cordgrass due to variations in hydrology and soil biogeochemistry caused by microtopography and distance from tidal creeks, creating distinct subsites. Our overarching objective was to observe how soil C concentration ~~changes~~and dissolved organic carbon (DOC) vary across four plant phenophases and across three subsites categorized by unique vegetation~~,~~ and hydrology~~, and biogeochemistry, while~~. We also ~~investigating~~investigated the dominant biogeochemical controls on the spatiotemporal variability of soil C ~~concentration~~and DOC concentrations. We hypothesized that subsite biogeochemistry drives spatial heterogeneity in soil C concentration, and this causes variability in total soil C and DOC concentration at the marsh scale. In addition, we hypothesized that soil C concentration and porewater biogeochemistry vary temporally across the four plant phenophases (i.e., senescence, dormancy, green-up, maturity~~), causing further variation in marsh soil C that could lead to uncertainty in soil C estimates.~~). To test these interrelated hypotheses, we quantified soil C and DOC concentrations in 12 cm sections of soil cores (0-48 cm depth) across time (i.e., phenophase) and space (i.e., subsite), alongside several ~~porewater biogeochemical variables including dissolved organic carbon (DOC), EEMs/ UV-VIS, redox potential, pH, salinity, reduced iron (Fe^{2+}), reduced sulfur (S^{2-}), and total porewater element (Fe, Ca) concentrations in three distinct subsites.~~other porewater biogeochemical variables. Soil C concentration varied significantly ($p<0.05$) among the three subsites and was significantly greater during plant dormancy. Soil S, porewater sulfide, redox potential, and depth predicted 44% of the variability in soil C concentration. There were also significant spatial differences in the optical characterization properties of DOC across subsites. Our results show that soil C varied spatially across a marsh ecosystem up to 63% and across plant phenophase by 26%, causing variability in soil C ~~storage~~accrual rates and stocks depending on where and when

samples are taken. This shows that hydrology, biogeochemistry, and ~~ecological function~~plant phenology are major controls on ~~saltmarsh~~salt marsh C content. It is~~, therefore,~~ critical to consider ~~spatial and temporal~~spatiotemporal heterogeneity in soil C concentration and porwewater biogeochemistry to account for these sources of uncertainty in C stock estimates. We recommend that multiple locations and sampling timepoints are sampled when conducting blue C assessments to account for ~~soil carbon~~ecosystem-scale variability~~ and uncertainty in C stock estimates.~~.

1 Introduction

Coastal blue carbon (C) cycled in tidal salt marshes is critically important for global soil C sequestration despite the small relative land area (Mcowen et al. 2017). High primary productivity coupled with high sedimentation rates and slowed organic C decomposition due to flooded anoxic soils allow salt marshes to rapidly accrete and preserve soil C (Arias-Ortiz et al. 2018). Soils in such ecosystems retain approximately 15% of their yearly primary productivity in soils compared to just 1% for tropical rainforests (Duarte 2017). Restoring, protecting, and artificially creating salt marshes can facilitate removal of CO_2 from the atmosphere and storage in soils on timescales conducive to climate change mitigation goals. These ecosystems should therefore be included in climate mitigation policy (Ewers Lewis et al. 2019; Serrano et al. 2019). However, a wide range of global salt marsh soil C sequestration rates of ~ 1 to >1100 g C m^{-2} $year^{-1}$ has been reported (Wang et al. 2021). The inclusion of salt marshes in improved climate mitigation policy is, in part, contingent upon improving our understanding of the environmental variables causing wide ranges in marsh soil C concentration and thus soil sequestration rates (Saintilan et al. 2013; Macreadie et al. 2019). Understanding key controls on salt marsh soil C variability will also decrease uncertainty in Earth System Models and inform new policy aimed at protecting these valuable ecosystems.

Soil C concentrations in salt marsh ecosystems vary spatially across the globe. Part of this variation is explained by regional environmental controls such as average annual air temperature (Chmura

et al. 2003), geomorphic setting (van Ardenne et al. 2018), salinity gradients, inundation frequency (van de Broek et al. 2016; Baustian et al. 2017; Luo et al. 2019), rainfall patterns (Sanders et al. 2016; Negandhi et al. 2019), soil controls such as pH, soil moisture, and soil type, as well as plant controls such as the dominant plant species ~~and soils~~ (Bai et al. 2016; Ford et al. 2019). Soil C accumulation rates also vary based on the age of the marsh and tend to be highest in newly expanding marsh edges (Miller et al. 2022). Other logistical factors contributing to variability and heterogeneity in salt marsh blue C estimates include the type of corer used (Smeaton et al. 2020) and the depth of soil that is integrated into ~~storage~~soil C accrual rates (Bai et al. 2016; Van De Broek et al. 2016; Mueller et al. 2019). While understanding global and regional controls on soil C is important for reducing uncertainty in C estimates, understanding site-level factors is also critical because ecosystem-level variability can be just as high as regional- to global-level variability (Ewers Lewis et al. 2018). Belowground biogeochemical heterogeneity ~~is often noticeable in~~ can be attributed to the aboveground vegetation and plant controls due to striking zonation of marsh grass species across the marsh platform. This is often ~~attributable to~~associated with small spatial-scale changes in hydrologic patterns (Guimond et al. 2020b, a) based on proximity to the tidal channel that drives unique subsite biogeochemistry (Seyfferth et al. 2020) which subsequently determines the type of vegetation that can survive within a given tidal zone (Davy et al. 2011). While tidal zonation alters vegetation and belowground biogeochemistry, it remains unclear if soil C concentrations are directly or indirectly altered by these ~~dynamics.~~coupled plant and soil biogeochemical controls.

Primary production rates may partially control soil C concentration and may vary among vegetative zones. For example, the short form of *Spartina alterniflora* has a lower primary production rate than the tall form (Roman and Daiber 1984) and *Phragmites australis* has above and below ground production rates two times that of the shorter *Spartina patens* (Windham 2001). Belowground ~~productivity includes root exudates (Luo et al. 2018) in the form~~production of dissolved organic carbon (DOC~~), which could~~) can arise from root exudation (Luo et al. 2018) and influence soil C concentration because belowground productivity often exceeds above-ground productivity in these ecosystems (Frasco and Good 1982). Even though DOC exudates are considered to be labile (Yousefi Lalimi et al. 2018),

they may contribute to soil C accumulation over time due to microbial transformation (Valle et al. 2018) and association with soil minerals such as Fe oxides (Chen et al. 2014; Chen and Sparks 2015; Sowers et al. 2018a, b, 2019). The optical characterization of DOC ~~quantified by optical properties~~ of chromophoric dissolved organic carbon (CDOM) can also affect degradability (Clark et al. 2014) and may differ across the marsh platform as a result of differing plant species.

~~Subsites~~Vegetation zones or subsites can have unique biogeochemical signatures based on soil redox conditions and inundation extent and frequency. For example, high marsh areas and areas near tidal channels have soils which are at least periodically oxic to sub-oxic and are dominated by iron (III) reduction, whereas low marsh areas have continuously inundated soils and are dominated by sulfate (SO_4^{2-}) reduction (Seyfferth et al. 2020). While these biogeochemical characteristics can directly influence vegetation (Moffett and Gorelick 2016) and thus indirectly influence soil C concentrations, ~~they~~these heterogeneous biogeochemical characteristics may also directly affect soil C through the interactions of soil C cycling with soil minerals. Fe oxides have an intimate role in the C cycle and C stabilization in soils experiencing dynamic redox fluctuation (Sodano et al. 2017), as previous work has shown that 99% of the dissolved Fe in the ocean is complexed with organic ligands (Whitby et al. 2020) and ~21% of all organic C in marine sediments is bound to reactive Fe species (Lalonde et al. 2012). Fe oxides may play an important role in C stabilization in soils experiencing dynamic redox fluctuation. Fe oxides can protect DOC against microbial degradation through physiochemical protection (Blair and Aller 2012; Chen and Sparks 2015; Sodano et al. 2017; Sowers et al. 2018a; Dorau et al. 2019; Wordofa et al. 2019), but these organo-mineral assemblages can be dissociated under reducing conditions (Riedel et al. 2013; Wordofa et al. 2019; Lacroix et al. 2022; Fettrow et al. 2023a). Therefore, examining the spatial variability in soil biogeochemistry and relating those variables to soil C concentration may elucidate important mechanisms that cause the wide range in salt marsh soil C concentrations.

While it is critical to assess spatial heterogeneity in soil C concentration, it is also important to assess temporal variability. The temporal assessment of soil C in salt marshes often considers long-term trends of historic C burial rates (Cusack et al. 2018; McTigue et al. 2019; Breithaupt et al. 2020; Cuellar-

Martinez et al. 2020), but variability of salt marsh soil C concentrations may also occur on shorter time scales such as across a single year. Several studies suggest salt marsh soil C does not significantly change across seasons throughout the year (Yu et al. 2014; Zhao et al. 2016), even though major changes in soil biogeochemical variables occur on this timescale (Koretsky et al. 2005; Negrin et al. 2011; Seyfferth et al. 2020; Trifunovic et al. 2020; Zhu et al. 2021). While soil C concentration may be stable across seasons, it is unclear if soil C concentration changes based on site-specific plant phenology. The phenophase of a marsh is associated with the greenness index of vegetation (Trifunovic et al. 2020) and is strongly associated with C dynamics in wetland systems (Desai 2010; Kang et al. 2016). Soil C concentration should be measured across plant phenophase to determine if temporal changes in phenology alter soil C concentration ~~and cause another source of~~, adding to variability in ~~ecosystem scale~~blue C estimates.

To address these knowledge gaps, we conducted a year-long study of a temperate tidal salt marsh to assess how soil C concentration and porewater biogeochemistry change in space (subsite) and time (phenophase). Our overarching research objectives were to understand how soil C and porewater DOC concentration and ~~soil biogeochemistry~~properties change across spatial and temporal scales~~,~~ and to investigate key biogeochemical ~~mechanisms influencing soil~~drivers of these C ~~concentration~~concentrations at the ecosystem level. We hypothesized that subsites would contain significantly different concentrations of soil C due to differences in soil biogeochemistry across the marsh platform. We further hypothesized that soil C concentration and associated porewater DOC and biogeochemistry would significantly differ across plant phenophase. Our results improve understanding of mechanistic controls on salt marsh soil C with implications for characterizing and reducing uncertainty in C sequestration estimates, while also adding to the body of literature that shows tidal salt marshes are critical reservoirs of sequestered C.

2.0 Methods and Materials

2.1 Field Site

This study was conducted at the St. Jones National Estuarine Research Reserve located in Dover, Delaware (Figure 1). The ecosystem is classified as a temperate mesohaline tidal salt marsh with a tidal

creek salinity ranging from 5 to 18 ppt (Capooci et al. 2019). Three separate subsites were previously identified at this site, each with a different vegetation type and hydrology (Guimond et al. 2020a; Seyfferth et al. 2020). The subsite nearest the channel is primarily colonized by the tall form of *Spartina alterniflora* and has semidiurnal tidal oscillation. This subsite is hereafter referred to as Tall Spartina (TS). Farther from the tidal channel, the elevation is slightly higher due to a natural levee and flooding of the upper 25 cm of soil occurs only during spring tides; this location has the larger cordgrass *S. cynosuroides* and is hereafter referred to as Tall Cordgrass (TC). The third subsite is farthest from the tidal channel, lowest in elevation, and is primarily colonized by the short form of *S. alterniflora* due to near continuous inundation; this subsite is hereafter referred to as Short Spartina (SS). These subsites have distinct hydro-biogeochemistry and vegetation that varies across small spatial scales and thus provides an ideal setting to understand site-level variability in soil C concentration, porewater biogeochemistry and their relationships.

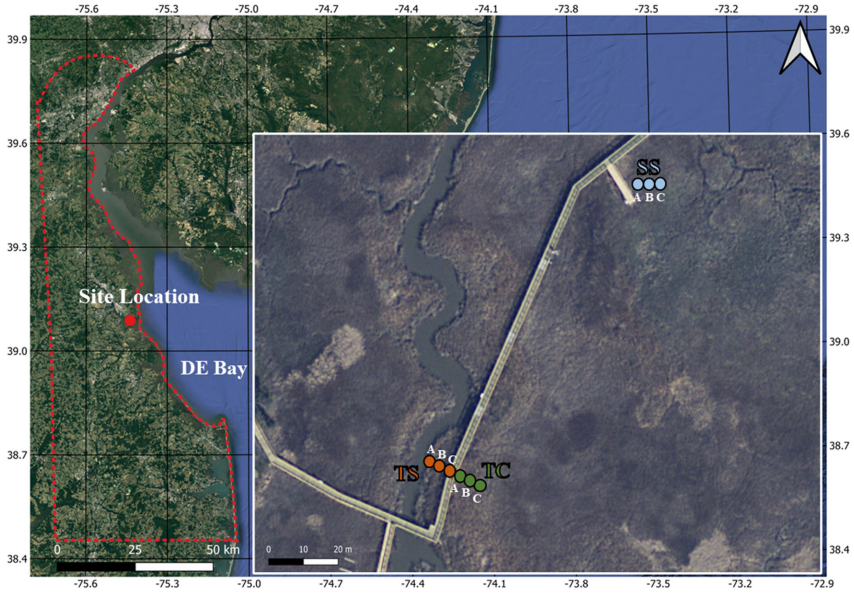

Figure 1. Map of the field site located at the St Jones Reserve near Dover, DE. Three unique subsites (TS, TC and SS) have been characterized based on previous studies at this field site showing subsite specific hydrology, vegetation, and biogeochemistry based on distance from the tidal creek (Guimond et al., 2020a; Seyfferth et al., 2020). The coring locations were sampled in triplicate (Core A, B and C), with core A

starting closest to the creek and each subsequent core in each subsite being ~~30cm~~30 cm from one another. The base layer for the map was obtained from public base layers in QGIS (© Google Maps).

2.2 Soil Sampling and Analysis

Soil cores were obtained from each of the three subsites (TS, TC, SS) in triplicate during each sampling event. Replicates were taken approximately ~~30cm~~30 cm from one another and are labeled cores A, B, and C based on distance to the tidal channel with A being closest to the channel and C the farthest (Figure 1). Sampling events occurred at four separate times of the year to coincide with each of the phenophases (i.e., senescence on 10/3/2019, dormancy on 12/3/2019, green-up on 4/29/2020, maturity on 8/13/2020), which were previously determined using the Greenness Index (Trifunovic et al. 2020). Cores were obtained at the same tidal inundation cycle each season to ensure consistent saturation during each campaign. Each sampling campaign resulted in 36 total cores (or 144 core sections, see below) that we used to understand spatiotemporal variability; unfortunately, we could not obtain more cores due to conditions of the strict soil coring permit at the estuarine preserve. Soil cores (6 cm x 48 cm) were extracted using a gouge auger that has been shown to be an effective coring technique for reducing compaction in soft marsh soils (Smeaton et al. 2020). Soil cores were quickly sectioned in the field into 12 cm increments (0-~~12cm,~~ 12-~~24cm,~~ cm, 12-24 ~~36cm~~ cm, 24-36 cm and 36-~~48cm~~48 cm relative to the soil surface) and preserved under anoxic conditions following previous methods (Seyfferth et al. 2020). For reference, the rooting zone of *Spartina* grasses is between 8-~~20cm~~20 cm (Muench and Elsey-Quirk 2019), so the upper two sections likely include C from fresh root exudates. The ~~12cm~~12 cm increments were chosen because many soil C stock papers use increments between 10-15 cm and there tends to be little variation across the ~10 cm increment in a variety of wetland soils (Baustian et al. 2017). Briefly, the soil sections were placed into 250 ml HDPE bottles which were left uncapped in gas-impermeable bags that contained oxygen scrubbers (AneroPack-Anero, Mitsubishi), and the bags were vacuum-sealed in the field. The soil samples were placed on ice during transport back to the lab. Once back in the lab, the soil sections in the gas-impermeable bags were immediately placed inside an anoxic glove bag containing ~5% hydrogen and

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~95% nitrogen. A subsample of soil was dried, ground, sieved (~~2mm~~2 mm), and powdered for analysis of total C and S (Vario EL Cube, Elementar). We clarify that we did not separate inorganic versus organic soil C and report only total soil C. Soil C and S are reported as % C (= 100% * g C/g soil dry wt.) and % S (= 100% * g S/g soil dry wt.). ~~The remaining field-moist soil was left inside the HDPE vial, capped inside the glove bag, and centrifuged for extraction of porewater using methods in the following section.~~We used soil C % to calculate soil C stocks using previously obtained bulk density measurements at our field site (Wilson and Smith 2015), and we calculated soil C accrual rates using previously obtained sedimentation rate values (Tucker 2016). The remaining field-moist soil was left inside the HDPE vial, capped inside the glove bag and centrifuged for extraction of residual porewater. The amount of porewater we obtained was a function of soil saturation that was consistent during each campaign because we sampled at the same tidal cycle each season. After centrifugation, the remaining soil sample was further dried inside of the glove bag. While this drying procedure could have introduced artificial H_2-fueled metabolism, this should be negligible because the soils were rapidly dried within the glove bag with freshly replaced desiccant and because the saturated sample was only minimally in contact with the H_2 atmosphere.

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2.3 Porewater Extraction and Analysis

Porewater was extracted from each 12-cm soil section by centrifugation for 2 minutes under an anoxic atmosphere at 2,500 rpm. A portion of the porewater was filtered with 0.45µm PTFE syringe filters while the rest was vacuum filtered using glass fiber filters (0.7µm). The 0.45µm PTFE filtered porewater was immediately analyzed for Fe^{2+} using the ferrozine colorimetric method (Stookey 1970), S^{2-} using the methylene blue method (Cline 1969), redox potential with a ~~220mV~~220 mV offset, pH, and conductivity using calibrated probes (Orion Ross Ultra pH/ATC Triode, Orion 9179E Triode, Orion DuraProbe Conductivity Cell), and the remaining sample was acidified to 2% HNO_3 for elemental analysis using an ICP-OES. The porewater filtered with glass fiber (0.~~7µm~~7 µm) was acidified with HCl and analyzed for DOC (Vario TOC Analayzer, Elementar). To characterize the DOC, unacidified DOC samples from the plant maturity sampling event were analyzed via ultraviolet-visible (UV-VIS)/

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excitation-emission matrix spectroscopy (EEMs) (Aqualog Spectrophotometer, Horiba). The Aqualog was zeroed with double deionized water blanks, checked using the manufacturer's excitation check, corrected for inner filter effects, applied first and second order Rayleigh masking and data were normalized using the average Raman area (Gao et al. 2011; Clark et al. 2014). Measurements were taken over the wavelengths of 200-730nm with 2nm steps. Fluorescence and absorbance peaks and indices were calculated using previously established equations (Table S1).

2.4 Statistical Analysis

~~Statistical~~A three-way analysis of variance (ANOVA) was performed to understand significant interactions between factors of subsite, depth, and phenophase on soil and porewater variables. Subsequently, statistical differences between subsites and phenophase were analyzed using repeated measures ~~analysis of variance (~~ANOVA~~) (α=0.05),~~ with a post-hoc Tukey-HSD analysis to determine differences between individual subsites and ~~phenophase.~~phenophases. Assumptions of ANOVA were met by assessing for normality with QQ plots prior to analysis and transforming when necessary. Equal variance was tested to ensure homogeneity of variance between subgroups using Levene's test. Correlations with depth were analyzed using linear regression and only the significant ($p<0.05$) relationships are reported. Relationships among all measured variables were assessed using principal components analysis. In addition, a stepwise regression model was built to determine variables that significantly predict soil C concentration. This was done by maximizing the R^2-value of the model while using the least amount of variables to explain the variance. All statistical analyses were conducted in JMP (Version 16.2).

3.0 Results

3.1 Soil Carbon and Sulfur

To explore the spatiotemporal heterogeneity of soil carbon (C) and sulfur (S) at each subsite, subsamples of each collected soil increment were combusted for soil C and S concentration. Concentrations of soil C were highly variable among subsites, phenophase, depth, and replicate cores (Figure 2), indicating several ~~possibles~~patiotemporal sources of variability in marsh soil C ~~stock~~ estimates. SS ~~showed the highest~~appeared to have higher soil C concentrations, as illustrated by darker

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colors in the heat map, compared to both TS and TC. Soil C ~~was~~ also appeared higher at TS than TC, illustrated by relatively darker colors in the heat map. For all subsites, soil C concentrations changed throughout the year ~~with the highest values~~, appearing higher during plant dormancy and ~~the lowest~~lower during green-up. However, variability across individual replicates A, B, and C and with depth complicated generalities across time and space. For example, at subsite SS from 24-36 cm during senescence, core A is ~5% soil C while core C is ~10% soil C, a factor of 2 difference ~~within~~between replicates. Large ranges among replicates were also observed during green-up at TS from 12-24 cm and during maturity at TC from 36-48 cm. This exemplifies the high spatial and temporal heterogeneity inherent in marsh soils, and ~~a source~~indicates several sources of ~~variation~~uncertainty in marsh soil C estimates.

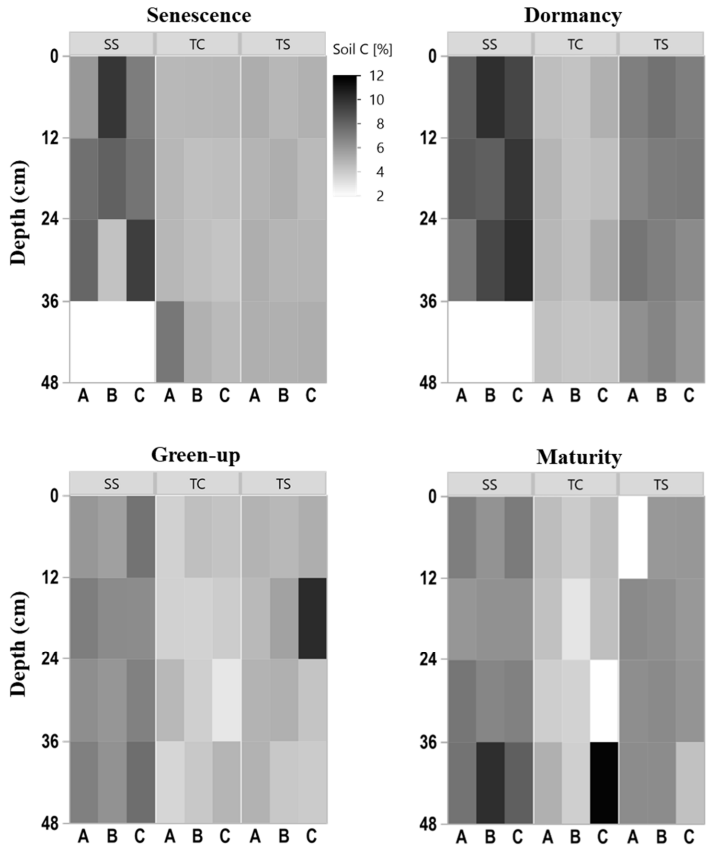
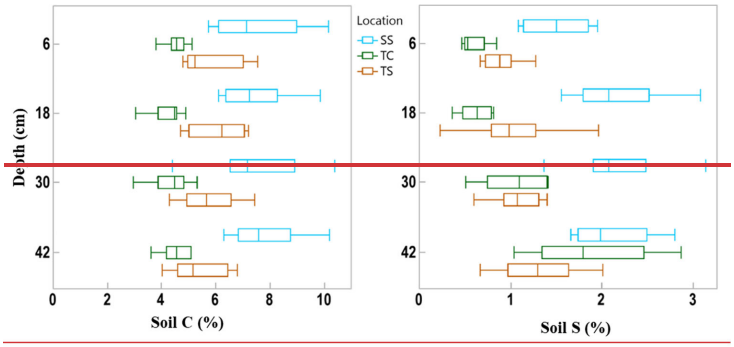


Figure 2. Heat maps of soil C concentration with depth at the three subsites (SS, TC, and TS), four phenophases, and for each replicate core (A (closest to channel), B, and C (farthest from channel)). No measurement was able to be obtained for some 12- cm sections as shown by white rectangles.

There was also variability in soil C concentration with depth (Figure 3). Subsite SS had the highest mean soil C concentration at all four depths, as well as the largest range in values. TS had the second highest mean soil C values at all four depths as well as the second largest range in values. TC had the lowest mean soil C at all four depths as well as the smallest range in values at each depth. It is clear from this graph that SS contains higher overall concentrations of soil C, followed by TS and then TC.

~~Soil~~When observing linear trends with depth, soil C at TS during dormancy significantly decreased with

depth (R^2=0.44, p=0.02) and soil C at SS during maturity significantly increased with depth (R^2=0.41, p=0.02). No other linear correlations in soil C existed with depth.



We also assessed differences in soil C and S with depth by averaging by phenology and subsite replicates (Figure 3). These results showed that there were no significant differences in soil C with depth For soil S, only the first and second depths were significantly different from one another at site SS and at TC, the deepest cores had significantly more soil S than all other depths.

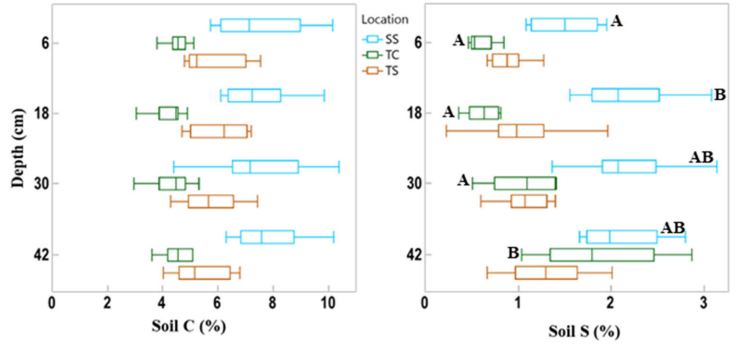


Figure 3. Box and whisker plot of soil C and S concentrations across the three subsites and separated by the four sampling depths. This indicates the difference in soil C and S variability among subsites and with depth. Whiskers indicate the minimum and maximum values, and the box indicates the upper and lower quartiles. The line in the box indicates the median. Letters with significant differences (p<0.05) with depth for each subsite are shown by different letters; subsites and depths with no letters are statistically similar.

~~Soil S also varied across 12 cm sampling increment depths (Figure 3).~~ SS had the highest mean soil S concentration at each depth, and the range of values initially increased with depth. TS has a higher mean concentration than TC at all depths except at the bottom core section. The range of soil S values increased with depth at TC while the range was more consistent with depth at TS, except for the wide range of values measured at the ~~18cm~~18 cm depth interval. Soil S at SS during maturity significantly increased with depth (R^2=0.50, p=0.01), as did TC during dormancy (R^2=0.88, p<0.0001), green-up (R^2=0.51, p=0.01), and senescence (R^2=0.42, p=0.02). ~~No other correlations between soil S existed with depth.~~

3.2 Porewater Data

3.2.1 Porewater DOC and Characterization

Porewater DOC was highly variable across subsites, phenophase, depth, and replicate cores (Figure 4). Note that the data in Figure 4 have been log transformed (natural log) due to large ranges in values across the one-year sampling campaign. Unlike soil C, which was relatively consistent with depth, DOC concentrations were highly variable with depth and even more so among replicate cores. Some of the highest individual concentrations of DOC were detected nearest the surface and rooting zone, which can extend to 20 cm below the surface (Muench and Elsey-Quirk 2019), but also at depth at SS during senescence. DOC concentrations decreased with depth at SS during green-up (R^2=0.44, p=0.02) and maturity (R^2=0.37, p=0.03) and increased with depth at TC during dormancy (R^2=0.76, p=0.0002). These results indicate the highly variable nature of porewater DOC concentrations, possibly leading to additional and complexity in marsh soil C estimates. In addition, we summarized DOC concentrations across depths and subsite (Figure 5) to better understand variability with depth. The top depth increment at 6 cm appeared to contain the greatest variability, particularly at subsite TC. Variability at TC decreased with depth, as did variability at SS. This is apparent because the range tends to decrease with depth at both TC and SS. Overall, TC seems to contain the most variability followed by TS and SS appears to contain the least amount of variability at each depth increment.

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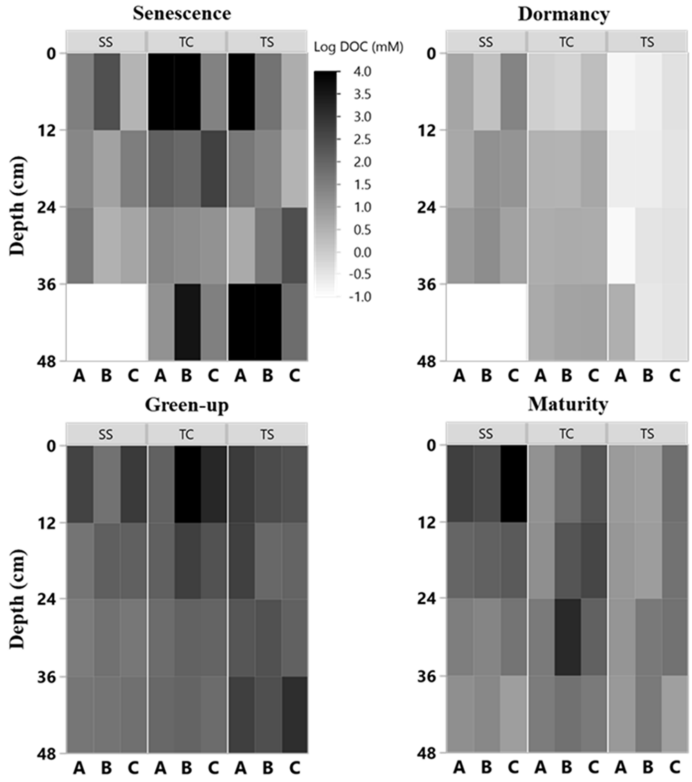

Figure 4. Heat maps of porewater DOC (natural log) concentration with depth at the three subsites (SS, TC, and TS), four phenophases, and for each replicate core (A (closest to channel), B, and C (farthest from channel)). No measurement was able to be obtained for some 12-cm sections as shown by white rectangles.

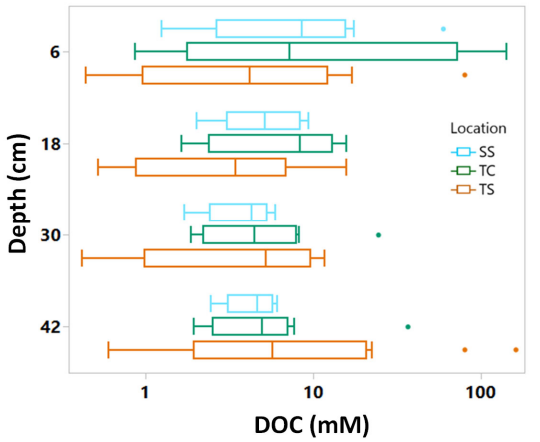

Figure 5. Box and whisker plot of porewater DOC concentrations plotted on log scale across the three subsites at all phenophases and separated by the four sampling depths. Whiskers indicate the minimum and maximum values, and the box indicates the upper and lower quartiles. Points outside the boxes indicate outliers. Due to large variability present across depths, there are no statistically significant differences between depths.

Porewater ultraviolet-visible (UV-VIS) and excitation emission matrices (EEMs) data were collected only from the maturity sampling event to further characterize DOC molecular properties (Figure 5). Optical properties (i.e., peaks, indices) from spectroscopic data were calculated and interpreted following previous studies cited in the supplemental table (Table S1). These data show significant trends with depth at SS. At SS, coble peak intensities T (R^2=0.55, p=0.01), B (R^2=0.49, p=0.01), A (R^2=0.57, p=0.004), M (R^2=0.55, p=0.01) and C (R^2=0.49, p=0.01) all significantly decreased with depth, as did the fluorescence index (FI) (R^2=0.79, p=0.0001), the biological index (BIX) (R^2=0.50, p<0.01) and absorbance at 254nm (Abs_{254}) (R^2=0.36, p=0.04), indicating decreases in CDOM with depth. To ensure the coble peaks represented changes in CDOM properties and not DOC concentration, they were normalized to DOC concentration and the relationships remained significant (p<0.05), except for the Coble B peak (R^2=0.11, p=0.20). The E_2:E_3 (R^2=0.50, p=0.01) and $SUVA_{254}$ (R^2=0.53, p=0.007) significantly increased with depth at SS, indicating a decrease in molecular weight and an increase in

aromaticity with depth. No significant trends with depth were present at TC or TS. Differences in DOC molecular properties among subsites are apparent for many of the calculated indices and peaks.

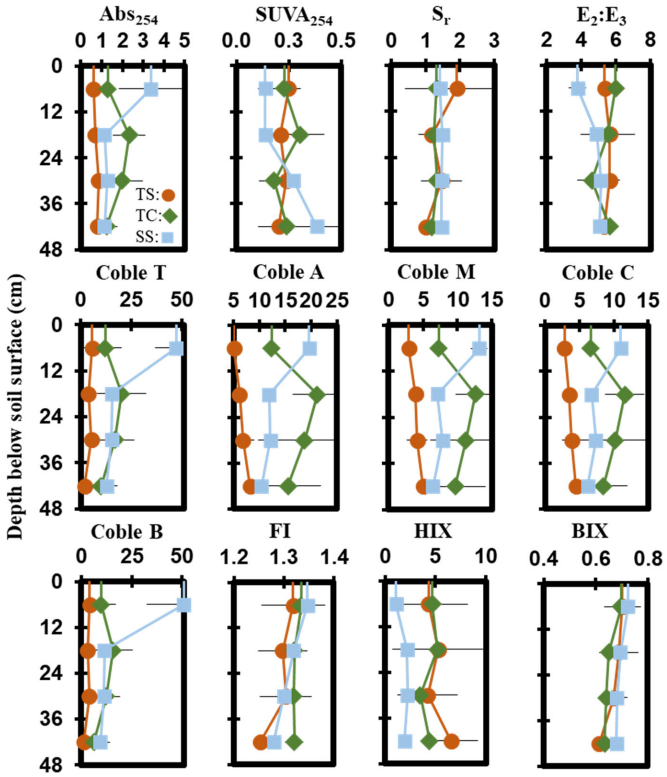

Figure ~~5~~6. Depth profiles of porewater EEMs/ UV-VIS peaks and indices down to 48cm taken during the maturity sampling event. Each point represents the mean between replicates (n=3) with error lines indicating the standard deviation (± 1 SD).

3.2.2 Porewater Chemistry

Measured porewater biogeochemistry was variable across subsites, phenophase, and depth (Figure 6). Porewater redox potentials showed minimal trends with depth, except for a significant decrease with depth at SS during maturity (R^2=0.58, p=0.004), though redox showed variability between replicates (Figure S2). The pH was relatively consistent with depth, except for a significant increase with

depth at TC during dormancy (R^2=0.42, p=0.02), and a significant decrease with depth at TS during dormancy (R^2=0.56, p=0.005). Redox potential and pH formed a significant but weak negative correlation (R^2=0.12, p<0.0001) across the entire 1-year dataset.

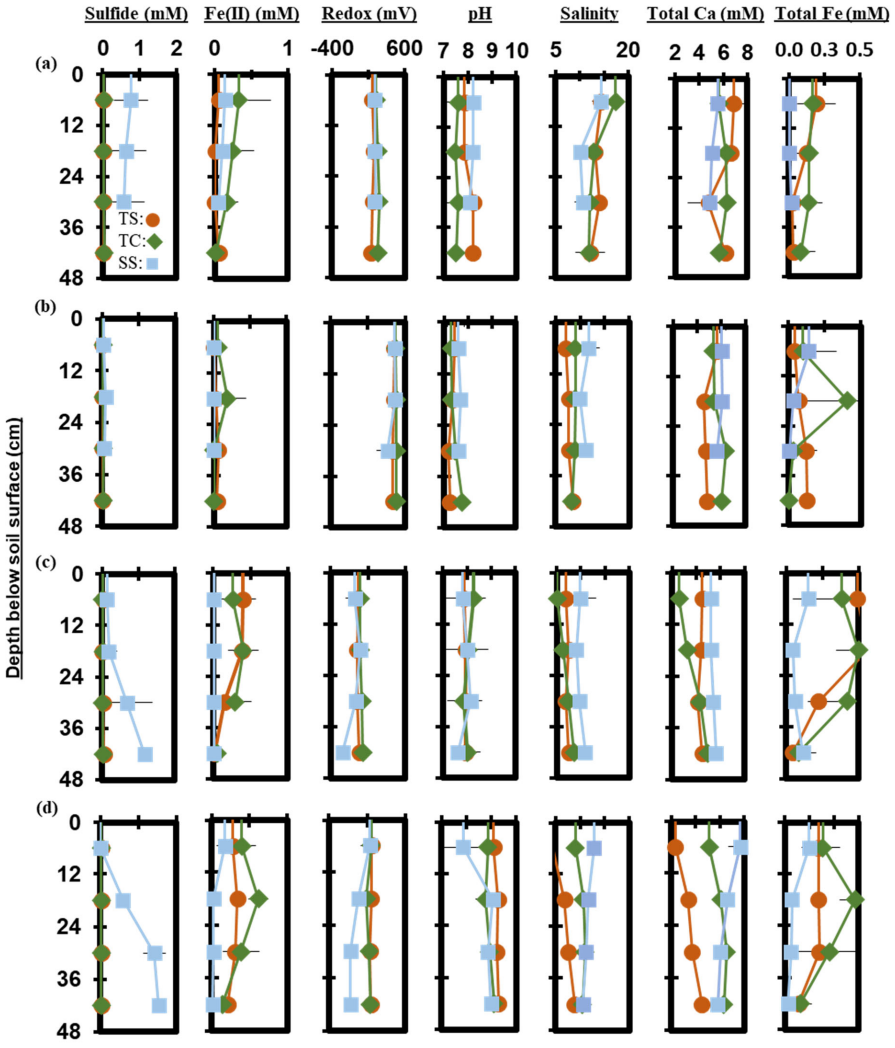

Figure ~~6~~7. Depth profiles of porewater chemistry variables down to ~~48cm~~48 cm for sampling events that occurred during plant (**a**) senescence, (**b**) dormancy, (**c**) green-up and (**d**) maturity. Each point represents the mean between replicates (n=3) with error lines indicating the standard deviation (± 1 SD).

Porewater S^{2-} varied significantly with depth. S^{2-} increased significantly with depth across the entire 1-year dataset (R^2=0.04, p=0.03). S^{2-} increased significantly with depth at SS during green-up (R^2=0.51, p=0.01) and maturity (R^2=0.86, p<0.0001). TS S^{2-} increased significantly during green-up (R^2=0.46, p=0.02) while TC S^{2-} increased significantly during maturity (R^2=0.36, p=0.04). Porewater Fe^{2+} trended negatively with S^{2-} (R^2=0.06, p=0.004) and decreased with depth (p=0.01, R^2=0.05) across the entire 1-year dataset. Significant decreases were observed at TS during green-up (R^2=0.68, p=0.001), and at SS during maturity (R^2=0.41, p=0.02). Total Fe concentration followed similar depth trends to Fe^{2+}, with a significant decrease with depth across the entire 1-year experiment (R^2=0.06, p=0.01). Total Fe decreased with depth at TS during senescence (R^2=0.41, p=0.03) and green-up (R^2=0.58, p=0.004), and at SS during maturity (R^2=0.57, p=0.01).

Porewater salinity formed varying relationships with depth. Salinity significantly decreased with depth at TC during senescence (R^2=0.52, p=0.01), and at SS during maturity (R^2=0.62, p=0.002) while salinity significantly increased with depth at TC during green-up (R^2=0.69, p=0.001) and at TS during maturity (R^2=0.87, p<0.0001). Salinity and total Ca generally increased together (p>0.0001, R^2=0.42) across the entire 1-year experiment. Total Ca increased significantly with depth at TC during green-up (R^2=0.86, p<0.0001) and at TS (R^2=0.80, p<0.0001) and TC (R^2=0.47, p=0.01) during maturity. SS total Ca significantly decreased with depth during maturity (R^2=0.60, p=0.005).

3.3 Analysis of Variance (ANOVA) Among Subsite, Depth, and Phenophase

A three-way ANOVA was run to assess the interaction between the three factors of phenology, subsite, and depth and to understand which factors are the most predictive for each variable (Table 1). Of the measured variables, only porewater DOC, sulfide, and salinity had significant interactions between all three factors; for these, one-way ANOVAs were ~~run on subsite and phenophase mean values~~ performed. These analyses showed that ~~were obtained by averaging samples from all~~ DOC was significantly higher during senescence at TS and TC in the surface than the other depths ~~across all four~~, subsites, and

phenophases ~~(for subsite comparisons) and all depths across~~, and that salinity was highest in the surface at all three subsites during senescence. In contrast, sulfide was highest during maturity at SS in the deepest core section compared to the other depths, subsites, and phenophases. For the variables without significant interaction, the three-way ANOVA showed that subsite was highly significant for soil C while phenology and depth were not significant. In contrast, phenology was only significant for porewater pH and Fe(II) while depth and subsite were not significant, and depth was only significant for soil S and sulfide (Table 1).

Table 1 Three-way ANOVA results for all variables, with interaction results of subsite, phenology and depth. Bolded p-values indicate significance ($p \leq 0.05$).

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Variable	Phenology	Subsite	Depth	Phenology*Subsite*Depth
Soil C (%)	0.06	**<0.0001**	0.95	0.96
Soil S (%)	0.99	0.89	**0.01**	0.99
DOC (mM)	.17	.91	**.02**	**.004**
Redox (mV)	.07	.31	.36	.77
pH	**<0.0001**	0.43	0.77	0.92
Fe^{2+} (mM)	**<0.0001**	0.06	0.39	0.91
Sulfide (mM)	0.80	0.91	**0.01**	**0.05**
Salinity (ppt)	**<0.0001**	**<0.0001**	0.99	**0.003**
Total Fe (mM)	0.98	0.27	0.21	0.75
Total Ca (mM)	**0.0001**	**0.003**	0.41	0.37

In addition to the three-way ANOVA, we also averaged variables by phenophase, subsite, or depth and performed one-way ANOVAs with post-hoc Tukey tests (Tables 2, 3, and Supplementary Table 2). When averaged by subsite, all ~~three subsites (for phenophase comparisons). These results show significant spatial and temporal variability in many of our measured variables. All three subsites contain significantly~~ subsites contained significantly different average concentrations of soil C, with SS having the highest average (7.5 % C), followed by TS (5.8 % C) and TC (4.6 % C) (Table 2). This indicates that on average, subsite SS contains ~29 % more soil C than TS and 63 % more soil C than TC. ~~In addition~~Site SS also had higher soil S, sulfide, and salinity and lower redox potential and Fe(II) than the other subsites. When grouped by phenophase, plant dormancy contained significantly more soil C than plant green-up~~.~~ ~~While soil S did not significantly vary across phenophase, soil S at SS was significantly higher in concentration by a factor of two than both TS and TC.~~ (Table 3). In addition, plant dormancy had significantly higher redox potential and the lowest Fe(II) and DOC than the other plant phenophases (Table 3). When averaged by depth, soil S was nearly 2x higher at the deepest depth (36-48 cm) than the surface (0-12 cm) (Supplementary Table 2).

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Table ~~1. One-way~~2. ANOVA and Post-hoc Tukey results for all assessed soil and porewater biogeochemical variables. Mean values represent average values for each subsite for subsamples from all depths and ~~phenophase~~phenophases. The mean is reported (± SD) along with a connecting letter report. Means with letters that ~~do not connect~~differ are significantly ($p<0.05$) different.

Variable	Tall Spartina (TS)	Tall Cordgrass (TC)	Short Spartina (SS)
Soil C (%)	$5.8\pm(1.2)^{B}$	$4.6\pm(1.3)^{C}$	$7.5\pm(1.4)^{A}$
Soil S (%)	$1.1\pm(0.5)^{B}$	$1.0\pm(0.6)^{B}$	$2.0\pm(0.7)^{A}$
DOC (mM)	$11.9\pm(27)^{A}$	$13.6\pm(27)^{A}$	$7\pm(9)^{A}$
Redox (mV)	$179\pm(176)^{AB}$	$211\pm(185)^{A}$	$93\pm(235)^{B}$
pH	$8.12\pm(0.8)^{A}$	$7.99\pm(0.7)^{A}$	$8.13\pm(0.6)^{A}$
Fe^{2+} (mM)	$0.15\pm(0.1)^{A}$	$0.22\pm(0.3)^{A}$	$0.04\pm(0.1)^{B}$

Sulfide (mM)	$0.02\pm(0.01)^{B}$	$0.02\pm(0.01)^{B}$	$0.6\pm(0.6)^{A}$
Salinity (ppt)	$8.8\pm(3.1)^{B}$	$9.7\pm(3)^{AB}$	$11\pm(2)^{A}$
Total Fe (mM)	$0.21\pm(0.2)^{A}$	$0.26\pm(0.3)^{A}$	$0.08\pm(0.1)^{B}$
Total Ca (mM)	$4.7\pm(1.3)^{B}$	$5.4\pm(1.2)^{A}$	$5.8\pm(0.8)^{A}$

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Table ~~2. One-way~~3. ANOVA and post-hoc Tukey results for all assessed soil and porewater biogeochemical variables. Mean values represent average values for each phenophase for subsamples from all depths and subsites. The mean is reported (± SD) along with a connecting letter report. Means with letters ~~that do not connect~~differ are significantly ($p<0.05$) different.

Variable	Senescence	Dormancy	Green-up	Maturity
Soil C (%)	$5.7\pm(1.5)^{AB}$	$6.7\pm(1.1)^{A}$	$5.3\pm(1.5)^{B}$	$6.1\pm(1.8)^{AB}$
Soil S (%)	$1.4\pm(0.7)^{A}$	$1.4\pm(0.9)^{A}$	$1.4\pm(0.7)^{A}$	$1.3\pm(0.7)^{A}$
DOC (mM)	$22.2\pm(42)^{A}$	$1.6\pm(1)^{B}$	$12.3\pm(14)^{AB}$	$7.9\pm(10)^{B}$
Redox (mV)	$193\pm(60)^{B}$	$453\pm(58)^{A}$	$-42\pm(98)^{D}$	$83\pm(111)^{C}$
pH	$7.89\pm(0.4)^{B}$	$7.45\pm(0.2)^{C}$	$7.96\pm(0.6)^{B}$	$8.94\pm(0.5)^{A}$
Fe^{2+} (mM)	$0.1\pm(0.2)^{BC}$	$0.03\pm(0.1)^{C}$	$0.2\pm(0.2)^{AB}$	$0.2\pm(0.2)^{A}$
Sulfide (mM)	$0.2\pm(0.4)^{AB}$	$0.04\pm(0.04)^{B}$	$0.2\pm(0.4)^{AB}$	$0.3\pm(0.6)^{A}$
Salinity (ppt)	$12.9\pm(2.4)^{A}$	$9.0\pm(1.8)^{BC}$	$8.0\pm(2.1)^{C}$	$9.6\pm(2.4)^{B}$
Total Fe (mM)	$0.1\pm(0.1)^{B}$	$0.1\pm(0.2)^{B}$	$0.3\pm(0.2)^{A}$	$0.3\pm(0.2)^{A}$
Total Ca (mM)	$5.8\pm(1.0)^{A}$	$5.5\pm(0.7)^{A}$	$4.5\pm(0.9)^{B}$	$5.3\pm(1.6)^{A}$

DOC concentration also varied among subsites (Table ~~1~~2) and phenology (Table ~~2~~3). The average DOC concentration at SS was approximately half of that found at TS and TC, but these results are not statistically significant due to large variability and ranges in concentration observed across the 1-year experiment. This large variability is exemplified by standard deviations that are larger than the means. In addition, DOC also varied across phenophases. Dormancy had the lowest mean DOC concentration and was significantly lower than senescence by an order of magnitude. Maturity and green-up did not have statistically different DOC concentrations. The EEMs/ UV-VIS dataset from plant maturity was analyzed based on subsites (Table ~~3~~4). There were significant differences in peaks and indices between subsites. Coble peaks T, A, M, C and Abs_{254} were significantly lower at TS than at both TC and SS by at least a factor of two which is in line with the lower DOC concentrations observed for TS at maturity ~~(Fig. 4).~~. Subsite SS had a significantly lower HIX and E_2:E_3 than both TS and TC suggesting it to have DOM with

less relative humic content and higher average molecule weight. These results indicate significantly different DOC molecular characteristics across subsites. EEMs/ UV-VIS data could not be assessed across phenology since these data were collected only during plant maturity.

Table ~~3~~4. One-way ANOVA results for UV-VIS EEMs during the plant maturity phenophase. Mean values represent average values for each subsite for subsamples from all depths. The mean is reported (± SD) along with a connecting letter report. Means with letters that ~~do not connect~~differ are significantly ($p<0.05$) different.

Parameter	Tall Spartina (TS)	Tall Cordgrass (TC)	Short Spartina (SS)
Abs_{254}	$0.7\pm(0.2)^{B}$	$1.7\pm(0.9)^{A}$	$1.7\pm(1.3)^{A}$
$SUVA_{254}$	$0.2\pm(0.1)^{A}$	$0.2\pm(0.1)^{A}$	$0.2\pm(0.1)^{A}$
s_r	$1.39\pm(0.95)^{A}$	$1.27\pm(0.33)^{A}$	$1.46\pm(0.28)^{A}$
E_2:E_3	$5.5(0.4)^{A}$	$5.4\pm(1.1)^{A}$	$4.7\pm(0.7)^{B}$
Coble T	$4.1\pm(3.8)^{B}$	$14.7\pm(10.3)^{A}$	$22.6\pm(16.2)^{A}$
Coble A	$6.6\pm(2.1)^{B}$	$16.9\pm(7.02)^{A}$	$13.5\pm(4.2)^{A}$
Coble M	$4.0\pm(1.4)^{B}$	$10.2\pm(4.4)^{A}$	$8.6\pm(3.1)^{A}$
Coble C	$3.7\pm(1.2)^{B}$	$9.2\pm(4.0)^{A}$	$7.8\pm(2.3)^{A}$
FI	$1.3\pm(0.6)^{A}$	$1.3\pm(0.02)^{A}$	$1.3\pm(0.03)^{A}$
HIX	$5.1\pm(3.0)^{A}$	$4.4\pm(3.1)^{A}$	$1.9\pm(0.6)^{B}$
BIX	$0.7\pm(0.7)^{A}$	$0.7\pm(0.03)^{A}$	$0.7\pm(0.02)^{A}$

Differences in porewater chemistry among subsites (Table ~~1~~2) and phenophase (Table ~~2~~3) were also significant. SS had the lowest average redox potential and was significantly different from TC which had the highest, while TS was not significantly different from either SS or TC. Redox potentials were even more variable between phenophase where all four phases had significantly different means. The highest mean was measured during dormancy and decreased significantly in the order senescence, maturity and green-up. The pH was not significantly different across any of the subsites but did change

437 significantly with phenology. Dormancy had the lowest pH which was significantly different from all
438 other phenophases. Senescence and green-up had a statistically similar mean pH values that were higher
439 than dormancy, and the porewater pH during maturity was statistically higher than all other phenophases.
440 S^{2-} also varied significantly among subsites. SS contained on average more than an order of
441 magnitude greater S^{2-} than both TS and TC. S^{2-} is lowest during dormancy but is only significantly
442 different than maturity which has the highest S^{2-} mean. Variability in Fe^{2+} between subsites was opposite
443 of S^{2-}. While TS and TC had low concentrations of S^{2-}, they had high concentrations of Fe^{2+}, which were
444 more than double and significantly higher than Fe^{2+} at SS. Fe^{2+} concentrations varied with phenology
445 similar to S^{2-} where dormancy had the lowest mean which was significantly different only from maturity
446 when the highest levels of Fe^{2+} were detected. Differences between subsite total Fe followed the same
447 trend as Fe^{2+}, where SS was significantly lower than both TS and TC. Total Fe was lowest during
448 dormancy and senescence, which were both statistically similar, but different from green-up and maturity.
449 SS had the highest mean salinity and was significantly different only from TS which had the
450 lowest mean salinity. Green-up had a significantly lower mean salinity than all other phenophases except
451 dormancy. Dormancy was only significantly different from senescence, which had the highest mean
452 salinity. Subsite differences in Ca were similar to salinity where SS had a significantly higher mean Ca
453 concentration than TS, but not TC. Green-up had the lowest mean Ca concentration which was
454 significantly different from all other phenophases.

455 **3.4 Stepwise Regression Model Results**

456 A stepwise regression model was run across the entire 1-year experiment to determine the most
457 important biogeochemical predictors of soil C concentration in our dataset (Table 45). The model results
458 indicate that depth, redox potential, soil S, and sulfide are the best predictors of soil C concentration. The
459 model R^2 value of 0.44 indicates that these variables explain 44% of the variability in our soil C
460 concentration data and the model is highly significant ($p < 0.0001$). Sulfide, redox potential, and soil S
461 each have positive estimates, meaning that these variables increase as soil C increases while depth had a

462 negative estimate, meaning that soil C tends to decrease with depth across the entire dataset. Each
463 individual predictor variable is also significant ($p<0.05$).
464
465 **Table 45.** Stepwise regression results for predicting soil carbon.

Parameter	Estimate	P-Value	Model R^2	Model P-Value
Depth	-0.03	0.003	0.44	<0.0001
Sulfide	0.96	0.04		
Redox	0.002	0.002		
Soil S%	1.3	<0.0001		

466
467 **4.0 Discussion**
468 **4.1 Subsite Differences in Soil C and Biogeochemistry**
469 We hypothesized that soil C concentration and soil biogeochemistry would differ across our
470 subsite locations. Our results support this hypothesis and suggest significant differences in both soil C
471 concentration and porewater biogeochemistry among subsites, which is consistent with prior work at this
472 field site (Seyfferth et al. 2020; Guimond et al. 2020a). This finding illustrates the importance of
473 considering multiple sampling locations when conducting blue C assessments to account for ecosystem-
474 scale variability. At SS, average soil C concentrations were 63% higher than at TC and 29% higher than
475 at TS. Even though these subsites are several to tens of meters from one another, they each had
476 statistically different mean soil C concentrations. Higher soil C at SS is not related to higher primary
477 productivity because the *Spartina alterniflora* at SS are stunted. The short form of *S. alterinflora* is
478 generally less productive than the tall form (Roman and Daiber 1984) and likely exudes less DOC from
479 the smaller root mass. This is supported by a lower average DOC concentration at SS. Also, the
480 chromophoric dissolved organic matter (CDOM) properties at SS were different than at the other subsites.
481 SS CDOM had a significantly lower E_2:E_2 than TS and TC, indicative of higher molecular weight DOC at

SS. In addition, the humification index (HIX) was significantly lower at SS indicating that the DOC at SS has been reworked by microbes less than it has been at TS and TC. Furthermore, SS consistently had lower porewater redox potentials than the other subsites; while our data represent a snapshot in time for each phenophase and subsite location, they are consistent with prior work of higher resolution porewater over time that shows SS being more strongly reducing than areas closer to the tidal channel (Guimond et al., 2020a; Seyfferth et al. 2020). Redox potentials at SS were low enough to support sulfate reduction. This is confirmed by our elevated S^{2-} porewater concentrations measured at SS. Therefore, the greatest controls on soil C concentration at SS is slower microbial oxidation of C due to strongly reducing conditions caused by nearly constant inundation and limited flushing of oxygenated surface water (Guimond et al. 2020b, a; Seyfferth et al. 2020). These conditions lead to CDOM that is less affected by microbial degradation (i.e., low HIX, low E_2:E_3) and a less energetically favorable metabolism (i.e., sulfate reduction) resulting in more soil C ~~storage~~accrual. This has important implications for soil C stock uncertainty because a greater amount of the area at St Jones is composed of subsite SS (Seyfferth et al. 2020). Sampling only near the tidal creek (TS and TC) could significantly underestimate soil C stocks, while sampling only in the marsh interior could lead to an oversimplification of soil biogeochemistry and DOC molecular properties in salt marsh ecosystems.

In contrast to SS, soil redox potentials were significantly higher at TC and soil C was significantly lower. This is likely due to TC having a slightly higher elevation on a natural levee and less reducing surface soils (Seyfferth et al. 2020). The redox potential is not low enough to support sulfate reduction but is low enough to support Fe reduction. This is supported by the abundant amount of Fe^{2+} measured in the porewater at TC. A higher redox potential and more energetically favorable electron acceptor (Fe^{3+}) likely leads to higher rates of C mineralization and explains the lower soil C concentration at TC. On the other hand, we found some of the highest concentrations of DOC at TC, particularly closer to the surface near the rooting zone. This can be explained by a greater root mass and correspondingly higher root exudation rate of the taller *S. cynosuroides* coupled with porewater flushing occurring only on a spring-neap pattern, which allows DOC to build up in porewater over time (Guimond et al. 2020a, b). A

higher concentration of freshly produced DOC and a lower concentration of soil C is also consistent with the priming effect which posits that high concentrations of freshly produced and microbially labile DOC can stimulate microbial growth leading to the degradation of older, more stable soil C (Textor et al. 2019; Zhang et al. 2021). In addition, TC CDOM fluorescence peaks (Coble, A, M, C, T), were similar to SS, indicating that SS and TC have strong sources of fluorescent CDOM.

Though TS and TC are biogeochemically more similar than SS, TS had significantly higher soil C than TC likely due to different dominant vegetation and hydrology. TS is lower in elevation and experiences diurnal tidal oscillations with slightly lower average porewater redox values than TC (Table 1), which experiences tidal oscillations on a spring-neap cycle (Guimond et al. 2020a). These differences in hydrology may cause soil C to accumulate more so under slightly stronger reducing conditions at TS compared to TC. Another unique attribute of subsite TS is the CDOM signature. The coble peaks (A, T, C, and M) and Abs_{254} were significantly lower at TS than both TC and SS, which indicates a decreased concentration of terrestrially-derived CDOM. This is likely because TS is nearest the tidal creek and therefore porewater solutes are exported to the tidal channel twice daily during ebb tide (Fettrow et al., 2023b), decreasing the marsh grass derived terrestrial CDOM signature in the near-channel porewater.

4.2 Phenophase Differences in Soil C and Biogeochemistry

We further hypothesized that soil C concentration and biogeochemistry would vary across plant phenophase, and our data support this hypothesis. Soil C was greatest during plant dormancy and was on average 26% higher than green-up, 18% higher than senescence, and 10% higher than maturity. This highlights the importance of considering the time of year soil samples are taken when conducting a blue C assessment. Likewise, many of the biogeochemical variables also changed with phenophase. The redox potential of all four phenophases were significantly different from one another, with the highest average redox potential occurring during dormancy. Higher redox potentials during dormancy are associated with significantly lower porewater Fe^{2+} and S^{2-}, indicating that microbial reduction is likely suppressed during the winter months when labile DOC produced from root exudation is less available. Dormancy also had the highest soil C concentration. We suggest this may be related to a suppressed priming affect due to low

porewater DOC concentrations and to Fe oxide formation during the high redox potential of dormancy, allowing any remaining porewater C to be pulled out of solution and into the solid phase with oxidized Fe minerals (Riedel et al. 2013; Sodano et al. 2017; ThomasArrigo et al. 2019).

We found that DOC concentrations are higher during senescence and significantly lower during plant maturity. High porewater DOC during senescence agrees with previous work showing higher belowground allocation of biomass in *Spartina* before the winter (Crosby et al. 2015). Belowground allocation of C in *S. alterniflora* has been shown to increase late into the growing season (Lytle and Hull 1980) while concentrations of soil organics have been shown to decrease during the summer months due to higher temperatures and higher rates of soil respiration (Caçador et al. 2004). Higher rates of belowground C allocation during senescence are further supported by the higher rates of soil respiration during senescence (Vázquez-Lule and Vargas 2021) due to increased labile DOC availability and associated microbial activity previously reported at this field site.

4.3 Biogeochemical Controls on Soil C

Our data reveal important biogeochemical controls on soil C concentration across space and time. The results of the stepwise regression model suggest that soil C concentrations are predicted by sulfide, soil S, redox potential, and depth. Soil C increased significantly with increasing sulfide and soil S concentration, indicated by the positive model estimate (Table 45). This is likely associated with the lower elevation, and redox potential and greater accumulation of sulphate at SS due to less tidal flushing. This may also be a result of sulfurization where inorganic sulfur, namely sulfide, may interact with organic matter via abiotic reactions (Alperin et al. 1994). Evidence suggests that this interaction can help preserve and stabilize soil C (Tegelaar et al. 1989), though spectroscopic evidence would be required to determine if this is an important process at this study site.

Depth also has an important control on soil C concentration and the estimate was negative, indicating that soil C decreases with depth. This is consistent with the literature suggesting higher soil C concentration at the surface and decreasing with depth in coastal salt marshes (Bai et al. 2016). While depth was an important predictor of soil C from the stepwise regression model, our depth profiles (Figure

4) indicate only small changes with depth. This may be a result of only sampling to 48 cm and integrating across 12 cm increments, or it may be a result of our method design of extracting porewater from the soils and running porewater DOC as a separate fraction of C from the solid phase soil C. Because our porewater DOC results indicate higher concentrations near the surface, the removal of porewater DOC prior to soil C analysis may lead to lower concentrations of soil C at the surface because in most studies, porewater DOC is typically incorporated into the bulk soil C measurements upon soil drying and not extracted as a separate fraction of C (i.e., porewater DOC). We suggest future studies consider porewater DOC as a separate component of the overall soil C concentration, particularly because the variability with depth is much higher for porewater DOC than soil C and porewater DOC is presumed to be more labile and mobile than particulate OC. Therefore, when porewater is extracted from the soil, the measured soil C concentration may appear less variable with depth and time leading to more consistent estimates of the more stable solid-phase soil C.

Redox potential was the final significant predictor in the stepwise regression model and increased significantly with soil C. We expected to see a negative relationship between soil C and redox potential due to higher C preservation under reducing conditions, but an overall positive relationship between redox potential and soil C in the model indicates an additional and possibly more important mechanism related to shifting biogeochemistry throughout the year. We observed more oxic conditions at all subsites during plant dormancy in the winter, probably due to the cold winter conditions that allow for the higher dissolved oxygen concentrations in water and porewaters observed previously (Trifunovic et al. 2020). Despite more oxygenated conditions and higher redox potentials in winter, the microbial activity likely decreased during winter, allowing elevated soil C during the winter months when plants were dormant. In addition, the less reducing and more oxygenated conditions in winter likely promoted the formation of Fe oxides that incorporated solution-phase C into the solid phase via coprecipitation. While there is an abundance of evidence showing the importance of Fe oxides in soil C storage in non-wetland ecosytstems (Lalonde et al. 2012; Riedel et al. 2013; Sowers et al. 2018a, b, 2019; Adhikari et al. 2019), recent studies have shown the important role of Fe oxides in C cycling in tidal salt marshes (Seyfferth et al. 2020;

Fettrow et al. 2023a), but few studies track C cycling during the cool winter months. Variations in Fe oxide complexation with C due to phenological phase should be further investigated.

4.4 Variability in Soil ~~Carbon Storage~~C Accrual Rates and Soil C Stocks

Based on soil accretion rates obtained from a previous study near our core locations (Tucker 2016), bulk density at each of the three subsites previously obtained (Wilson and Smith 2015), and our mean soil C concentrations averaged across depth for each subsite within phenophases, we calculated the soil C accumulation rates and soil C stocks at each of the three subsites within each of the four phenophases (Figure 7). These accumulation rates are in range of previously reported values for mesohaline tidal salt marshes (Chmura et al. 2003; Lovelock et al. 2014; Ye et al. 2015; Mcleod et al. 2016; Macreadie et al. 2017, 2020), as are the soil C stock estimates (Zhao et al. 2016; Ewers Lewis et al. 2018; van Ardenne et al. 2018; Ouyang and Lee 2020; Gorham et al. 2021). These results further illustrate that soil C ~~storage~~accrual rates and soil C stocks are highly dynamic and change based on time and space within a single ecosystem. The largest difference between rates and stocks occurred between SS dormancy and TC green-up, in which the average ~~storage~~accrual rates varied by 75% and the average stocks varied by 96%. Therefore, within the same ecosystem and between phenophases, soil C ~~storage~~accrual rates and stocks can vary substantially, leading to variability and uncertainty. To account for spatial and temporal heterogeneity in soil C ~~storage~~accrual rates and stocks, we suggest taking soil cores across multiple vegetation zones (if they exist) and across both the growing and non-growing seasons. Our recommendation follows Howard et al. (2014), who suggest linear plot selection when an obvious feature (i.e., tidal creek) is present and a feature that likely has a strong control on local environmental conditions based on distance from this feature. But we also point out that selecting plot locations based on variation in vegetation is also important, since changing aboveground vegetation is often a sign of changing belowground biogeochemical conditions in tidal systems. This way, ~~more~~the source of variability can be accounted for, leading to less uncertainty in blue C estimates.

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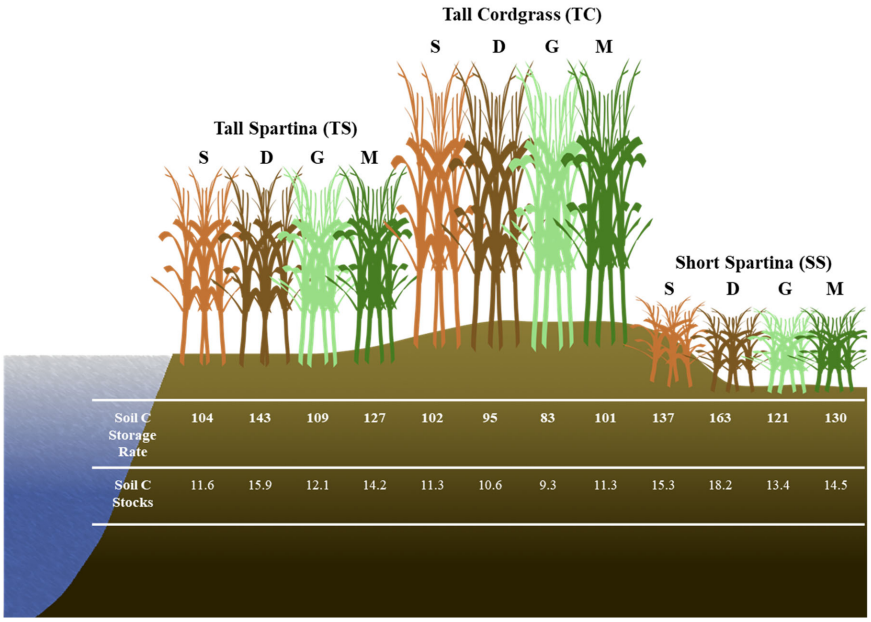

Figure ~~7~~8. Conceptual diagram illustrating the spatial and temporal variability of soil C ~~storage~~accrual rates (g C m^{-2} yr^{-1}) and soil C stocks (kg C m^{-2}) based on subsites by phenophase. Soil C stocks are 0 to ~~48cm~~48 cm depth. S= senescence, D= dormancy, G= green-up, M= maturity.

5.0 Conclusion

Our results highlight the variability in soil C in time and space at the site level. We found that some level of uncertainty in estimates of stocks and accumulation rates is likely related to spatial and temporal variability of soil C and biogeochemistry at the marsh scale. Subsites that were only a few meters from one another contained significantly different soil C concentrations, likely ~~used~~using different metabolic pathways for C mineralization, contained significantly different porewater CDOM molecular properties and led to considerable variation in soil C ~~storage~~accrual rates and soil C stock estimates. The biogeochemical controls that were best correlated with soil C concentration were redox potential, soil S, sulfide, and depth, indicating that the redox potential and sulfur content of the soils are critical in

controlling how much soil C accumulates in coastal marsh ecosystems. We also found that soil C concentration and thus soil C ~~storage~~accrual rates and soil C stock estimates, varies significantly across the phenophases of the marsh grasses. Plant dormancy contained the highest mean soil C concentration, possibly a result of high redox potential during winter months that causes remaining porewater DOC to be incorporated into the solid phase with oxidized minerals such as Fe oxides and lower microbial activity. These results demonstrate the importance of considering marsh-scale spatial and temporal heterogeneity when conducting a blue C assessment. Based on these results, we suggest taking soil cores from multiple locations within a marsh and in replicate, particularly if multiple vegetation types ~~of marsh grass~~ are present, and at different seasons to account for both spatial and temporal variability. These recommendations may help lead to less uncertainty in blue C estimates.

Statements and Declarations

Competing Interests: The authors have no relevant financial or non-financial interests to disclose.

Author Contributions: All authors contributed to the study conception and design. Material preparation, data collection, and analysis were performed by Sean Fettrow. The first draft of the manuscript was written by Sean Fettrow and Angelia Seyfferth with edits by Holly Michael and Andrew Wozniak. All authors commented on previous versions of the manuscript. All authors read and approved the final manuscript

Data Availability Statement

Data is available on Figshare (DOI: 10.6084/m9.figshare.24274417)

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