



1 Reviews and syntheses: Bioturbation impacts on sediment accretion and 2 erosion in tidal marshes, with implications for carbon burial and 3 sequestration 4

5 Leigh-Ann Smit^{1,2*}, Janine B. Adams^{2,3}, Gavin M. Rishworth^{1,2}

6 ¹Department of Zoology, Nelson Mandela University, Gqeberha, South Africa

7 ²SARChI: Shallow Water Ecosystems, Institute for Coastal and Marine Research (CMR),
8 Nelson Mandela University, Gqeberha, South Africa

9 ³Department of Botany, Nelson Mandela University, Gqeberha, South Africa

10 *corresponding author: s217268250@mandela.ac.za
11

12 1.1) Abstract 13

14 Tidal marshes offer multiple ecosystem services, but are some of the most threatened coastal
15 ecosystems worldwide. One of these valued services is their ability to sequester and store
16 large amounts of carbon. Bioturbating macrofauna are ecosystem engineers that can
17 influence the geomorphology and biogeochemistry of tidal marshes. Bioturbators can
18 influence accretion and erosion processes in tidal marshes by either stabilizing or destabilizing
19 sediment. Through this reworking of sediment, they can also influence the amount of carbon
20 that can be stored. The impact of bioturbation on tidal marshes depend on a number of factors,
21 such as, species composition, burrow morphology, diet, behaviour and habitat type. This
22 review assesses the current knowledge on the role benthic bioturbators play in shaping
23 sediment processes in tidal marshes and identifies key knowledge gaps for future research.
24 For example, the impact of individual benthic species on sediment dynamics is mostly
25 unknown. Bioturbation effects cannot be generalised and predicting when and where these
26 effects will be most prominent is challenging. Future studies should investigate family and
27 species specific effects on sediment properties, such as erodibility or texture, under controlled
28 laboratory conditions and in the field. This should be compared across different habitat types
29 such as ecotones, mudflats, salt marshes and mangroves. Furthermore, the role of
30 consumers, as bioturbators, remains an understudied driver of the carbon cycle because it is
31 complex. In order to better predict how tidal marshes may persist in the face of future climate
32 change, such as sea level rise, it is important to understand the role of bioturbators on
33 sediment and carbon dynamics to enable better mitigation of global change effects through
34 conservation and restoration of tidal habitats.

35 **Keywords:** blue carbon, benthic organisms, coastal ecosystems, ecosystem engineers,
36 sediment processes



1.2) Introduction: Tidal marsh sediment and carbon processes

38

39 Tidal marshes, such as salt marshes and mangroves, are vegetated coastal ecosystems that
40 are highly important in terms of their ecological value, because they exist between terrestrial,
41 estuarine and near-shore marine environments (Barbier, 2015). These coastal habitats offer
42 natural protection against storm surges and erosion (Perkins et al., 2015), in addition to other
43 essential services such as sediment retention, flood attenuation and nutrient processing (Bos
44 et al., 2007; Hatje et al., 2021). They provide important nursery areas for estuarine and marine
45 fishes and invertebrates (Sogard and Able, 1991; Barbier et al., 2011), and are also valuable
46 for tourism and food production (Hawkins et al., 2020; Lynch et al., 2023). Another important
47 ecosystem service provided by salt marshes and mangrove forests, is their ability to sequester
48 and store carbon (Macreadie et al., 2021). The carbon sequestered by these coastal habitats
49 is referred to as blue carbon (Nellemann and Corcoran, 2009; Mcleod et al., 2011). Although
50 seagrass beds are also classified as blue carbon habitats, they are primarily a subtidal habitat
51 and therefore not strictly part of tidal marshes in the context of this review. The term 'blue
52 carbon' was coined more than a decade ago (Duarte De Paula Costa and Macreadie, 2022),
53 with blue carbon research having increased over the last decade. This growing interest allows
54 for a better understanding of the global distribution of tidal marshes and the factors that
55 determine their persistence.

56 Salt marshes cover at least 41,700-54,900 km² of the globe (McOwen et al., 2017), mangrove
57 forests 150,000 km² (Spalding, 2010), and unvegetated mudflats approximately 127,921 km²
58 of the globe (Murray et al., 2019). The Northern Hemisphere has roughly double the amount
59 of tidal marshes as the Southern Hemisphere, due to their longer coastline (He et al., 2025).
60 The long-term persistence of tidal marshes is driven by the interactions between surface
61 elevation, sea level, sediment accretion and primary production (Morris et al., 2002). Surface
62 elevation and sediment accretion is regulated by abiotic and biotic factors, which includes
63 suspended sediment supply, climate, geography and bioturbation (Ouyang et al., 2022).

64 Coastal ecosystems are some of the most threatened systems worldwide with approximately
65 35 % of mangroves and 50 % of salt marshes being lost or degraded by anthropogenic
66 activities (Van Katwijk et al., 2016; Li et al., 2018). By means of satellite observations, looking
67 at changes in water presence, land losses, and gains can be estimated. It is estimated that
68 28 000 km² of land has been eroded in tidal marshes, which is double that of land gained
69 (Mentaschi et al., 2018). Some studies have revealed that accretion rates are insufficient for
70 tidal marshes to keep pace with sea level rise (e.g. Van Wijnen and Bakker, 2001), while
71 others have found that accretion rates are high enough to keep pace with moderate rises in



72 sea level (e.g. Morris et al., 2002). A dominant driver of coastal erosion is anthropogenic
73 influence, such as the clearing of mangrove forests, as well as natural disasters, such as
74 extreme storms (Mentaschi et al., 2018). Sea level rise and a changing climate is likely to
75 enhance coastal erosion. While these ecosystems are increasingly threatened, the vegetation
76 within them is a key contributor to the ecosystem services they provide.

77 Plants capture carbon dioxide from the atmosphere which they store as organic carbon, but
78 through respiration, some of this carbon also gets released. The carbon budget of a vegetated
79 habitat is used to provide an indication as to whether it is a carbon 'sink' or a carbon 'source',
80 which is related to the accumulation and discharge of carbon (Sitch et al., 2015). Salt marshes
81 and mangroves are important carbon sinks, even though these habitats cover less than 2 %
82 of the area of the global ocean (Duarte, 2017). These blue carbon habitats store up to 70 %
83 of carbon, relative to the ocean carbon cycle (Macreadie et al., 2014). It is estimated that they
84 store up to 276 to 822 Tg of atmospheric carbon dioxide per year, worldwide (Spivak et al.,
85 2019). However, a loss or degradation of blue carbon habitats not only reduces the capacity
86 of these ecosystems to act as natural carbon sinks but if degraded and disturbed these
87 habitats directly release high amounts of carbon into the atmosphere as CO₂ emissions
88 (Pendleton et al., 2012; Hatje et al., 2021). A loss of one hectare of any blue carbon ecosystem
89 is equal to losing 10-40 hectares of native forest, in terms of carbon emissions (Macreadie et
90 al., 2017). Blue carbon includes carbon that is stored in living biomass (branches, leaves,
91 stems), non-living biomass (dead wood, leaf litter), roots and soil (McLeod et al., 2011;
92 Lovelock and Duarte, 2019a). When carbon is stored in this manner it is an important
93 ecosystem service as it is an essential component of the carbon cycle (Keller et al., 2018).
94 Blue carbon habitats, if conserved, are able to act as net carbon sinks (Spivak et al., 2019).
95 There are three factors that determines the capture and storage of carbon in these habitats:
96 the ability to maintain particulate organic carbon, high productivity and the conversion of
97 carbon dioxide into plant biomass (Alongi, 2002). The sediment biogeochemistry then leads
98 to a slow decay of organic material (Kelleway et al., 2017c).

99 The storage of carbon in tidal marshes is influenced by environmental factors such as
100 differences in moisture, nutrients, sediment supply, salinity and acidity as this is important for
101 decomposition and primary productivity (Lovelock et al., 2007). Sediment depth, type and
102 deposition is also linked to carbon storage ability (Kelleway et al., 2016b). Sediment grain size
103 has a strong influence on carbon storage because it influences the amount of organic particles
104 that can accumulate. The storage of carbon is greater in fine grained sediment because of the
105 lower oxygen exchange and porosity. Furthermore, these conditions decrease sediment redox
106 potential and the rates of remineralisation, thus enhancing carbon storage (Kelleway et al.,
107 2016b). Fine grained sediment also allows for the preservation of more organic matter



108 because of their higher surface area, which reduces the oxygen in the sediment as it is
109 consumed by detritivores which in turn decreases the decomposition of organic matter (Dahl
110 et al., 2016). Coarse grained sediment (sandy sediment) is more permeable and has more
111 aeration, increasing remineralisation of carbon (Van Ardenne et al., 2018). Carbon stored in
112 salt marsh sediment is also influenced by the community composition of vegetation due to the
113 differences in leaf and root morphology of different plant species. In general, shrubby salt
114 marsh vegetation has low carbon stock (Saintilan et al., 2013). The input of organic material
115 and the rate at which it decays is what ultimately determines the long term storage of carbon.
116 Carbon storage has been shown to be higher in mature salt marshes compared to restored or
117 new salt marshes (Alongi, 2018). Marshes that experienced rapid relative sea level rise during
118 the late Holocene have higher concentrations of soil carbon compared to those that were
119 subject to long periods of sea level stability (Rogers et al., 2019). Carbon storage is also higher
120 in salt marshes which experience limited erosion and where mangrove encroachment is
121 limited (Alongi, 2018).

122 For mangroves forests, latitude, productivity rates, the age of the forest, and elevation are
123 factors that have been linked to carbon stocks (Radabaugh et al., 2018). Mangroves are more
124 productive than salt marshes which results in salt marshes storing less carbon (Saintilan et
125 al., 2013). This has been attributed to lower redox potential, less anaerobic conditions and
126 higher tidal elevations of salt marshes which are not conducive to carbon storage (Schile et
127 al., 2017). Mangroves accumulate and store carbon over longer time periods (Lovelock and
128 Duarte, 2019). They also have a higher above and belowground biomass which enables them
129 to store more carbon (Donato et al., 2011). Mangroves are trees and therefore have a greater
130 biomass than salt marsh which are dominated by succulent herbs and grasses. Moreover,
131 water velocity is decreased by their aerial roots and more carbon rich sediment is able to be
132 deposited, as well as plant matter which further promotes the formation of carbon rich
133 sediment (Horstman et al., 2015).

134 A significant proportion of the global tidal marsh carbon is found in the temperate Northern
135 Atlantic, which has 45 % of the world's tidal marsh extent (Worthington et al., 2024). The U.S,
136 Canada and Russia are the top three countries with the highest predicted total sediment
137 organic carbon in their tidal marshes, because they have extensive marsh cover and high
138 carbon per unit area (Worthington et al., 2024). The global estimate of carbon in the top metre
139 of marsh sediment is 1.44 Pg C (Maxwell et al., 2024; Table 1). The average sediment organic
140 carbon per hectare is predicted to be about 83.1 Mg C ha⁻¹ in the 0-30 cm layer and 185.3 Mg
141 C ha⁻¹ in the 30-100 cm layer (Maxwell et al., 2024). Globally, it is estimated that mangroves
142 store around 11.7 Pg C, with most of the carbon stocks being in the sediment (Kauffman et



al., 2020). The global sediment stock of tidal flats is estimated to be 0.9 Pg C (Chen and Lee, 2022).

Table 1: Continent-level summary for tidal marsh area and sediment organic carbon (SOC).

Habitat	Region	Area (km ²)	SOC (Mg ha ⁻¹) ^a	SOC (Pg C)
Salt marsh		41,700-54,900 ^a		1.44 ^a
	Africa	2 241.37	1046.05	
	South America	4 537.76	710.53	
	North America	30 259.07	1045.54	
	Europe	11 054.68	1377.9	
	Asia	2 301.71	400.02	
	Oceania	2 378.58	172.86	
Mangrove		150,000 ^b		11.7 ^b
Tidal flats		127,921 ^c		0.9 ^c

^a (Maxwell et al., 2024)

^b (Kauffman et al., 2020)

^c (Chen and Lee, 2022)

Tidal marshes have gained interest for their recently recognised value of carbon storage, leading to extensive research on carbon stocks and factors influencing carbon sequestration and storage. Similarly, accretion and erosion dynamics of tidal marshes and the processes driving these changes is well understood. However, the influence of animal interactions on these processes is poorly understood, even though soil animals are key components of aquatic environments (Adams et al., 2025). This review provides an overview of the current knowledge on the influence of bioturbation on sediment accretion and erosion in tidal marshes, including the impact of bioturbation on carbon sequestration. Table S1 in the Supplementary material provides a summary of key bioturbation studies relating to accretion, erosion, and carbon sequestration, emphasising their methodologies and main findings that are discussed in the following pages, while Figure 2 shows where these studies were conducted.

To quantify the extent of research conducted on sediment processes and carbon in tidal marshes, a systematic literature search was performed in the web of science database using key words related to tidal marshes, carbon storage/sequestration and sediment dynamics. This search yielded 544 publications between the years 1993 and 2025. While a fair amount of research has been conducted on carbon stocks and sediment dynamics in tidal marshes, there remains a gap in our understanding of the role of bioturbators and their interaction processes on sediment dynamics. When key words relating to bioturbation were included, only 64 publications were yielded. Thus, the influence of these interactions on carbon sequestration and storage, and how this might be impacted in the face of climate change, which is a pressing future concern, is poorly understood compared to the overall science of tidal marsh carbon



173 and sediment processes. This review therefore aims to improve our understanding of how
174 bioturbators shape sediment dynamics and carbon cycling.

175 1.3) Bioturbation in coastal tidal marshes

176

177 Bioturbation in tidal marshes is associated with a number of organisms, found above and
178 below the surface sediment (Macreadie et al., 2017). Benthic invertebrates under the classes
179 Oligochaeta (worms), Gastropoda (snails), Polychaeta (polychaetes), Crustacea (crabs,
180 shrimp and malacostracans) and Bivalvia (cockles and mussels) are common bioturbators
181 found in tidal marshes (Van Der Wal and Herman, 2012). Some of the best studied groups
182 include crustaceans and molluscs (Booth et al., 2023). Bioturbators are significant
183 components of both terrestrial and aquatic ecosystems as they modify habitats, decompose
184 litter, and are also consumers (Wang et al., 2010). Bioturbation involves any transport process
185 performed by animals that affects sediment matrices, either directly or indirectly (Kristensen
186 et al., 2012), which includes burrow ventilation and particle reworking. Darwin (1881) was the
187 first to recognize the significance of animal bioturbation and its role in influencing soil
188 ecosystem processes. A dominant form of bioturbation in coastal wetlands includes that of
189 burrowing, with burrow architecture being species specific (Min et al., 2023; Fig. 1). One of
190 the most diverse groups, with special adaptations for burrow construction is Decapoda
191 (Giraldes et al., 2017; Hajializadeh et al., 2022). Burrow construction and maintenance, in
192 addition to ingestion and defecation results in particle reworking and biomixing. As a result,
193 microorganisms and organic matter are displaced within the sediment matrix, both laterally
194 and vertically (Kristensen et al., 2012). Benthic organisms can significantly affect the
195 composition of sediment, with destabilizing organisms generally decreasing mud content,
196 while stabilizing organisms can increase mud content (Arlinghaus et al., 2021). Animals that
197 rework sediment particles can be categorized as upward conveyors, downward conveyors,
198 biodiffusors and regenerators depending on their feeding type, behaviour and life style
199 (François et al., 2002). Collapsed burrows that are abandoned and become filled in, can be
200 considered as indirect bioturbation (Kristensen et al., 2012). Ventilation happens when
201 animals flush their burrows with water for feeding and respiration, and can be open with two
202 or more openings, or blind ended with one opening. This results in the rapid transport of
203 solutes from in the burrow to the overlying water (Kristensen, 2001). The activities associated
204 with bioturbation can therefore influence the physical, chemical and biological characteristics
205 of tidal marshes (Min et al., 2023).

206 Burrowing activities decreases sediment hardness, breaks up and transports sediment (Botto
207 and Iribarne, 2000), and increases the coarse particle density on the surface layers of the



208 sediment (Warren and Underwood, 1986). In addition, burrowing influences the chemistry of
209 the sediment, increases the oxygenation of the sediment and changes the pore water salinity
210 (Fanjul et al., 2007; Booth et al., 2023). Fine grained sediment, as well as sediment containing
211 high concentrations of organic matter can be trapped by crab burrows, which assists with
212 organic matter decomposition and increases the availability of nutrients (Fanjul et al., 2007).
213 The rate of nutrient and sediment turnover is further accelerated by means of excavation by
214 crabs, which transports nutrients and sediment from deep layers to the surface layers of the
215 salt marsh (Fanjul et al., 2007). Belowground processes are therefore impacted by burrowing
216 crabs which in turn influences marsh plants and trees by promoting growth (Botto et al., 2006;
217 Ngo-Massou et al., 2018). The interaction between the environment, the biology and the
218 density of a bioturbator determines the extent of the bioturbation effect (Wang et al., 2010; Xie
219 et al., 2020; Pan et al., 2023), which varies over space and time. For example, the presence
220 or absence of vegetation plays a key role in shaping this impact. When vegetation was
221 present, the quantity and quality of excavated and deposited soils (in burrow mimics) was
222 influenced, and thus, so was the burrowing effect (Wang et al., 2010). Vegetation can improve
223 nutrient concentrations, but its roots can obstruct the vertical movement of sediment.



230 **Figure 1: Examples of burrow openings of different crab species: *Scylla serrata* (A),**
231 ***Neosarmaticum africanum* (B), *Cyclograpsus punctatus*/*Parasesarma catenatum* (C**
232 **and D). The scale bar represents 10 cm in the foreground.**

233

234 1.4) Impacts of bioturbation on sediment processes

235

236 Bioturbation influences a number of sediment processes such as accretion, erosion, sediment
237 transport and deposition, which are outlined below and summarised in Table 2 and Table S1.
238 These processes are visually represented in Figure 3 and further explained in Table 3.

239 1.4.1) Accretion

240 Sedimentation is a key processes shaping tidal marshes, improving water clarity and quality
241 which helps submerged plants access sunlight (Nahlik and Mitsch, 2008). The sequestration



242 of carbon is also enhanced by sedimentation (Bernal and Mitsch, 2013) because the active
243 burial of carbon limits its exposure to oxygen thus, limiting oxidation (McCarty et al., 2009).
244 Salt marshes and mangroves persist when sediment carried by tides is deposited in vegetation
245 (Saintilan et al., 2022). This builds elevation and promotes the growth of plants which
246 increases belowground organic matter, resulting in elevation gain, slower water movement
247 and allows for more suspended sediment to settle (Kirwan and Guntenspergen, 2012). Plant
248 shoots promote the deposition of sediment while plant roots bind and stabilize the sediment
249 and can help prevent erosion (Buffington et al., 2020). Accretion therefore involves
250 sedimentation, root growth, and development of peat (Krauss et al., 2014; MacKenzie et al.,
251 2024)

252 Benthic organisms are able to facilitate sediment transport and sedimentation patterns over
253 extended periods and across surrounding areas (Arlinghaus et al., 2021). Their biological
254 activity impacts sediment structure in terrestrial, marine, and intertidal zones, either stabilizing
255 or destabilizing these environments. Some organisms enhance sediment cohesion by
256 producing an organic coating in the burrow walls from extracellular polymeric substances
257 (EPS), mainly mucus (Watling, 1991). *Sesarma reticulatum* (a crab occurring in northern
258 hemisphere temperate salt marshes) for example does this (Kristensen, 2008). These
259 biostabilization processes can therefore influence the strength of sediment in intertidal zones.
260 In a similar fashion microphytobenthic organisms form biofilms which can also improve the
261 stabilization of sediment (Decho, 2000).

262 Burrowing animals affect important ecosystem functions, while influencing the structure and
263 function of plant communities, with these effects varying in direction and magnitude regionally
264 (Vanni, 2002). Changes in the burrowing activities could have important consequences for the
265 functioning of salt marshes and mangroves. Low to moderate levels of bioturbation can be
266 beneficial to primary productivity (Kristensen et al., 2008). For example, burrowing by fiddler
267 crabs has been seen to benefit the growth of *Spartina alterniflora* by increasing soil drainage,
268 enhancing decomposition of plant debris and improving soil redox potential (Bertness, 1985).
269 Burrows can increase the surface area of the marsh allowing for the exchange of oxygen from
270 tidal water and the atmosphere which can increase the uptake of nitrogen increasing plant
271 productivity (Bradley and Morris, 1990; Sharbaugh et al., 2025)

272 Recent studies have highlighted the importance of bioturbation in determining changes in
273 surface elevation (Bennion et al., 2024). The accretion or erosion of sediment is partially
274 related to the burrowing and feeding activities of the species (Morelle et al., 2024). For
275 example, it was found that crab superfamily, whether it was an Ocypodoidea or Grapsodoidea,
276 had the biggest influence on sediment, as opposed to crab density (Rinehart et al., 2024),



277 which is related to their burrow morphology and diet (Table S1, Fig. 3). The composition of
278 crabs has the potential to influence ecosystems differently (Agusto et al., 2021). In mangroves,
279 changes in surface elevation is strongly influenced by species composition of the vegetation
280 and was positively influenced by the frequency of bioturbation. In salt marshes, however,
281 bioturbation had no significant effect on changes in surface elevation because they had lower
282 levels of bioturbation compared to the mangroves (Bennion et al., 2024) (Table S1, Fig. 3).

283 Excavated sediment through bioturbation activities, along with sediment from eroding areas
284 of the marsh, can contribute material for accretion on the surrounding marsh platform, helping
285 to increase marsh elevation (Wilson and Allison, 2008). Mussels, for example *Geukensia*
286 *demissa*, can also contribute to vertical accretion in salt marshes, as they harvest sediment
287 through their filtration activities, thus contributing to the sediment budget (Crotty et al. 2023)
288 (Table S1, Fig. 3). They also deposit faeces which is nutrient rich, indirectly increasing
289 vegetation biomass, improving soil shear strength (resistance to erosion) and stability. These
290 interactions therefore play an important role in promoting elevation gain and improving marsh
291 resilience.

292 1.4.2) Erosion

293
294 Due to coastal wetlands being situated at low elevation at the land sea interface, they are
295 susceptible to submergence and lateral erosion driven by wave activity, storm surges and
296 increased sea levels (Leonardi et al., 2018). The morphology and long term persistence of
297 tidal marshes is influenced by erosion. Erosion rates are determined by vegetation, which
298 affects sediment deposition rates and biological activity (Mudd et al., 2010; Cahoon, 2024).
299 Benthic organisms, specifically bioturbators, play a crucial role in influencing erosion
300 processes through their activities. Bioturbators can affect sediment roughness and alter its
301 characteristics, thereby influencing the erodibility of sediment (Dairain et al., 2020).
302 Bioturbators can have both direct and indirect effects on the erosion of tidal marshes. These
303 positive and negative impacts are expected to vary over time, as macrofaunal bioturbation is
304 temperature-dependent and tends to be more pronounced during warmer months (Cozzoli et
305 al., 2018). By reworking the sediment, bioturbators repack the sediment that was once
306 compact, which changes the texture and granulometry, causing larger aggregates of grains to
307 form (Grabowski et al., 2011). For example, *Scrobicularia plana* (a clam commonly found in
308 temperate European salt marshes) caused the sediment to become coarser and changed the
309 bed topography, which showed a loss by erosion (Morelle et al., 2024) (Table S1, Fig. 3). Fine
310 grained sediment such as clay and silt are more susceptible to the effects of benthos
311 (Arlinghaus et al., 2021). There are however still uncertainties with regards to the role that
312 benthic organisms play in sediment dynamics (Dairain et al. 2020; Farron et al. 2020). For



313 example, the influence of *S. reticulatum* bioturbation on the erodibility of sediment has not yet
314 been quantified as it is difficult to measure the processes in the field (Farron et al., 2020).
315 Thus, few studies have explored the connection between sediment stability and burrow
316 building bioturbators (Needham et al., 2013).

317 Burrowing activities weaken mud and clay banks in tidal marshes, making them more
318 susceptible to erosion through wave action. Dairain et al. (2020) observed that *Cerastoderma*
319 *edule* (common cockle, native to salt marshes in Europe and northwestern Africa) promotes
320 erosion of the surface sediment by increasing the roughness of the sediment, and this is due
321 to their sediment reworking activities (Table S1, Fig. 3). The same was true for the lugworm,
322 *Arenicola marina* (common in mudflats and salt marshes in Europe), which increased the
323 permeability and roughness of the sediment (Montserrat et al., 2011) (Table S1, Fig. 3).
324 *Sesarma reticulatum* contributes to changing erosion patterns by facilitating greater erosion
325 (Farron et al., 2020), which is likely driving the headward expansion of straight, low-order tidal
326 creeks in salt marshes within the Georgia Bight (Vu et al., 2017).

327 In addition to sediment disturbance, bioturbators can impact sediment cohesion and
328 erodibility. When the density of infauna were experimentally reduced in the Humber estuary
329 (UK), there was a 300 % increase in sediment stability on the intertidal mudflats (De Deckere
330 et al., 2001) (Table S1, Fig. 3). Invertebrates, such as crabs, can influence sediment stability
331 by consuming microphytobenthic organisms (Booth et al., 2023) which can indirectly promote
332 the destabilization of sediment (Daborn et al., 1993). Crabs can also contribute to sediment
333 destabilization by causing vegetation loss (Smit et al., 2024). Burrowing by *Sesarma*
334 *reticulatum* caused the upper 10-15 cm of the marsh to become oxidized which caused
335 enhanced degradation of belowground biomass of *S. alterniflora* (Wilson et al., 2012) (Table
336 S1, Fig. 3). This process reduces the shear strength of the sediment, increasing the erosion
337 potential which facilitates creek extension. Compared to the surrounding marsh platform, the
338 heads of newly formed creeks have lower topography, lack vegetation, and are densely
339 populated with both burrowing and herbivorous crabs. Over time these creek heads extend
340 further into the marsh platform as the creek migrates, which causes dieback of vegetated
341 areas and a loss of elevation of up to 50 cm (Day et al., 2011; Wilson et al., 2012). Similarly,
342 *Chasmagnathus granulatus* (a crab inhabiting the salt marshes of South America), through
343 their burrowing activities, have also been shown to increase the growth rate of tidal creeks,
344 causing larger creeks to form, which can promote salt marsh erosion (Escapa et al., 2008)
345 (Table S1, Fig. 3). In addition to their large scale effects on creek formation and vegetation
346 loss, crabs can also affect sediment structure at finer scales, through the formation of burrows.



347 Crab burrows, particularly those of species that do not plug their burrows during inundation
348 function as passive sediment traps (Grabowski et al., 2011; Escapa et al., 2008). However,
349 water filled burrows often lead to a reduced bulk shear strength and density, and reduced
350 erosion thresholds, which in areas that are heavily burrowed would increase the mass of
351 sediment eroded (Grabowski et al., 2011). Sediment trapping rate is dependent on burrow
352 architecture, density and possibly bed roughness (Escapa et al., 2008), therefore, different
353 species of burrowing crabs have different effects on the erosion and transport of sediment
354 (Min et al., 2023, Fig. 1).

355 1.4.3) Sediment transport and deposition

356
357 Sediment transport is often considered to be only a physical process, as a result of sediment
358 beds responding to hydrodynamic forces in coastal habitats (Le Hir et al., 2007). However,
359 biological components are also able to influence sediment transport processes. The
360 interaction between organisms and the sediment is complex and generally context specific,
361 due to factors such as hydrodynamics, sediment composition or species specific behaviours
362 (Needham et al., 2013). The influence of individual species on sediment dynamics are
363 therefore poorly understood. This makes it difficult to predict the overall impact of organisms
364 on sediment transport. While erosion and deposition are primarily driven by hydrodynamics,
365 benthic organisms influence the extent of these processes on a spatial and seasonal scale.
366 Studies have shown that benthos can cause change of the same order of magnitude as
367 hydrodynamic processes (Arlinghaus et al., 2021).

368 Crab burrow morphology is related to biological (e.g. sex or size; Sen and Homechaudhuri,
369 2016) and environmental (e.g. vegetation or sediment composition; Penha-Lopes et al., 2009)
370 factors, with morphology influencing their effectiveness in trapping sediment and organic
371 matter. Intertidal decapods construct funnel shaped burrows which aids in the trapping of
372 organic matter and sediment (Botto et al., 2006). Funnel shaped burrows with low aspect ratios
373 trapped a greater percentage of organic matter while tubular shaped burrows with a higher
374 aspect ratio trapped a greater amount of sediment (Botto et al., 2006) (Table S1, Fig. 3).
375 Gutiérrez et al. (2006) and Wang et al. (2010) deployed burrow mimics and found that less
376 material by weight was collected in the mimics than was excavated by crabs, indicating a net
377 export of sediment material (Table S1, Fig. 3). Excavation allows for buried material to be
378 brought to the surface, increasing the amount of sediment available for export by tidal flushing.
379 The quantity of sediment and organic matter available for transport is therefore a balance
380 between material deposited into crab burrows and material excavated from them.



381 Crabs create sediment mounds when they move sediment from their burrow to the surface.
382 During flooding and ebbing tide, this fresh mound sediment is transported. It remains a
383 challenge to predict when burrowing engineers will have a significant effect on their
384 environment (Coggan et al., 2018). However, the engineering effect is anticipated to intensify
385 as crab population densities increase (Rinehart et al., 2024). For example, burrowing crabs
386 are often found to have site specific effects on ecosystems (Beheshti et al., 2021), such as
387 promoting sediment trapping in one area of the marsh, but enhancing sediment removal in
388 other areas (Escapa et al., 2008). Crabs were found to promote the trapping of sediment in
389 open mudflats and intertidal salt marsh where current speeds are low, whereas in the salt
390 marsh edge, they were increasing sediment removal (Escapa et al., 2008) (Table S1, Fig. 3).
391 This was due to funnel shaped burrows being more frequent in the low intertidal zones as well
392 as the assistance of plants in trapping sediment. In habitats with weak flow, burrowing animals
393 are expected to promote sediment trapping, whereas in high flow energy habitats, burrowing
394 activity is anticipated to increase sediment removal rates, determined by the strength of the
395 current. In addition to crabs, *Thalassinidea* which are shrimp-like organisms, commonly
396 referred to as mud or sandprawns in South Africa, also influence sediment transport and
397 deposition. These burrowing species similarly create mounds by expelling sediment from their
398 burrows (Pillay and Branch, 2011). The transport of sediment by thalassinideans is greater
399 than that achieved by diffusion processes or abiotic burial (Grigg, 2003). The sediment
400 expelled from callianassid burrows is easily eroded at low current speeds because it is
401 unconsolidated, making it more prone to resuspension and redeposition in adjacent areas
402 (Pillay et al., 2007). *Kraussillichirus kraussi* (sandprawn characteristic of temporarily closed
403 estuaries in South Africa) consumes organic matter around its burrow, thus is an effective
404 mover of sediment (Pillay and Branch, 2011). Burrowing organisms are therefore key drivers
405 of sediment transport and redistribution in tidal marshes.

406 1.5) Impact of bioturbation on carbon burial and sequestration

407

408 Consumers can influence the carbon cycle directly and indirectly. For instance, small
409 bioturbating grazers change sediment properties and remove plant biomass. While they are
410 known to have an effect, they remain an understudied driver of carbon cycling (Guimond et
411 al., 2020; Ren et al., 2022). It was estimated by Montague (1982) that *Uca pugnax* (a species
412 of fiddler crab native to salt marshes along the coast of North America) excavated an amount
413 of carbon that is equal to 20 % of what *S. alterniflora* produces belowground annually, in
414 Sapelo Island (Table S1, Fig. 3). The amount of carbon collected in burrows was lower than
415 that made available for tidal flushing by excavation (Montague, 1982). The concentration of



labile and total carbon at the marsh surface is expected to decrease with crab activities because of the lower carbon content in the sediment that is excavated in relation to that deposited into the burrow (Gutiérrez et al., 2006). Burrowing organisms, such as crabs, can influence the carbon balance of tidal marshes by releasing carbon that would otherwise remain stored deeply in the sediment. Wittingham et al. (2024) showed that small grazers cause a decrease in carbon stocks with *S. reticulatum* accounting for a loss in carbon stocks of 40-70 % (Table S1, Fig. 3). In Cape Cod where marsh die off and erosion occurred due to overgrazing by *S. reticulatum*, an estimate of 248.6 ± 4.8 gigagrams of belowground carbon was released (Coverdale et al., 2014) (Table S1, Fig. 3). A correlation exists between crab burrows and carbon content, with higher densities of crab burrows associated with decreased carbon in the topsoil (Carpenter et al., 2023). The highest carbon content was found in salt marsh with minimal burrowing by crabs.

Complex burrow networks can have an effect on soil carbon stocks. A study conducted in Kenya found that mangrove forests that had a greater abundance of sesarmid crabs, had higher soil carbon stocks (Andreotta et al., 2014) (Table S1, Fig. 3). Crabs can also directly transfer carbon to sediments through the transportation of faeces, algae, leaf litter, and exuviae into their burrows (Alongi, 2002). This vertical transport of carbon was demonstrated through radiocarbon dating of sediment cores. Modern carbon was found to depths of 115 cm (Andreotta et al., 2014), which means that crabs are supplying new organic matter to deeper sediments. It is possible that the diversity of macrofauna in these ecosystems could be an important driver of carbon dynamics (MacKenzie et al., 2021). Macrofaunal diversity means a variety of sediment reworking activities, through bioturbation and bio-irrigation, which in turn can exert control on sedimentary biogeochemical cycling, such as carbon cycling (Meysman et al., 2006). On the other hand, crabs can also decrease carbon stocks because their burrows increase sediment surface area, aiding organic matter decay as more sediment becomes oxic, which leads to carbon loss via tidal flushing (Klaassen et al., 2025).

The effects of bioturbation on carbon cycling is context specific. For instance, *Macrophthalmus japonicas*, a salt marsh crab species from East Asia, increased the mineralization of sediment organic matter (SOM), stimulating the release of organic carbon, thus slowing the accumulation of organic carbon within sediment surface layers (Nie et al., 2021) (Table S1, Fig. 3). Similarly, bioturbation by *S. reticulatum* led to the remineralization of belowground organic matter by increasing the permeability and aeration of the sediment, leading to the degradation of organic material (Wilson et al., 2012). Crabs decreased SOM and carbon content in vegetated habitats and increased SOM and carbon in unvegetated habitats (Rinehart et al., 2024). Crab bioturbation has been shown to improve benthic metabolism and exchange of dissolved organic matter from the sediment to the water column (Fanjul et al.,



2015) (Table S1, Fig. 3). It was also found that the distribution, quality and bioavailability of sedimentary organic matter is influenced by bioturbation. Furthermore, efficient remineralisation of detritus occurs at bioturbated sediment and is exported as CO₂ and DOC to the water column. Bioturbation, by crabs, therefore improves the amount of labile organic carbon of bioturbated sediments and alters the pathway of carbon export to coastal waters (Fanjul et al., 2015).

While bioturbation can contribute to carbon loss, some bioturbating organisms can promote carbon storage. Burrows of *Upogebia major* (mudshrimp found in salt marshes in East Asia) and other thalassinideans have been found to trap organic matter (Kinoshita et al., 2008), which can increase the storage of carbon. Moreover, it was found that grazing by livestock had a neutral to positive effect on carbon sequestration (Graversen et al., 2022) (Table S1, Fig. 3). Crab burrowing was found to increase the turnover of nitrogen and carbon, with excavated soil having higher inorganic carbon concentration compared to soil deposited into burrows (Wang et al., 2010). This indicates that excavation activities accelerates the mineralization of organic matter from organic to inorganic carbon (Wang et al., 2010). Such changes to organic matter availability and benthic metabolism by bioturbation have the potential to decrease the storage capacity of carbon (Gutiérrez et al., 2006). Under accelerated sea level rise, consumers' impact on the carbon cycle, through carbon consumption and marsh stability, is expected to intensify as a result of the accelerated migration rates of consumer fronts, which are clusters of consumers bordering a specific resource (Wittingham et al., 2024).

1.6) Global change impacts on tidal marsh bioturbation

Blue carbon ecosystems are threatened by climate change, particularly sea level rise (Borchert et al., 2018; MacKenzie et al., 2024), as well as increasing temperatures and alterations in precipitation regimes (Arias-Ortiz et al., 2018; Adams et al., 2025). Coastal geomorphology, sedimentation patterns, geographic locality and regional oceanographic properties cause tidal marshes to become susceptible to these threats (McLeod et al., 2010). The resilience of salt marshes and mangroves to sea level rise is determined by physical drivers, such as unrestricted landward migration or increase in surface elevation (Schuerch et al., 2018; Lovelock and Reef, 2020) as well as biological drivers such as diversity (Branoff, 2020; He et al., 2025). The extent of development along the coast and the local topography controls the area available for these ecosystems to migrate landward, however, the rate of sedimentation controls the ability of salt marshes and mangroves to resist the rise in sea levels via the gain in relative surface elevation. The ability for sediment to be retained in the intertidal



487 region is dependent on local coastal dynamics and drainage basin geology (Adams et al.
488 2019). Furthermore, the structure of a wetland ecosystem affects its resistance to a
489 disturbance as well as recovery from a disturbance, therefore, local geomorphology
490 contributes substantially towards the resilience of these systems (Phillips, 2018). Mangrove
491 and salt marsh responses to sea level rise is thus not uniform across different regions and
492 even between sites within the same mangrove or salt marsh habitat (Passeri et al., 2015;
493 Adams et al., 2025).

494 Mangroves are specifically vulnerable to changes in temperature and precipitation regimes,
495 because the distribution range globally is linked to sea surface temperature. Mangrove
496 occurrence is limited to regions that are tropical or subtropical, and this by the winter 20 °C
497 isotherm (Tomlinson 1999; Hamilton and Casey, 2016). With rising temperatures comes an
498 expansion of mangroves polewards, to higher latitudes. Expansion of mangroves leads to a
499 loss of salt marsh habitats which results in ecological shifts as well as changes in the
500 provisioning of ecosystem services, for example carbon storage (Kelleway et al., 2017a).
501 Furthermore, mangroves that are found at range limits are also commonly smaller and shrub-
502 like (Morrisey et al., 2010), which influences their capacity to store and sequester carbon (Raw
503 et al., 2021). With rising sea levels, salt marshes are expected to migrate landwards (Enwright
504 et al., 2016). If the rate of sea level rise surpasses that of surface elevation gain it will cause
505 a shift in habitat with lower intertidal regions becoming subtidal and upper intertidal species
506 will encroach the terrestrial boundary (Fagherazzi et al., 2019). In salt marshes, as sea level
507 and consequently tidal prism begins to increase, it is expected that tidal creeks will develop,
508 which has been observed in Bahamas (Kirwan and Guntenspergen, 2012).

509 Regions that are more flooded (e.g. seaward areas) generally have smaller, shallower burrow
510 networks compared to those in drier regions (Egawa et al., 2021). Crab activity is highest in
511 summer and lowest in winter (Egawa et al., 2021), because of this seasonal change in
512 behaviour, it could further complicate the influence of crabs on carbon budgets (Guimond et
513 al., 2020) as regional historical temperatures change lined to behavioural phenology. Changes
514 in water levels and temperature, major components of climate change, can influence the
515 distribution of crabs and the extent of bioturbation (Wilson et al., 2022). Increased flooding
516 can suppress these activities, thus leading to redox conditions becoming more anoxic in tidal
517 marshes (Pan et al., 2023). On the other hand, faunal activities can interact with climate
518 stressors. For example, cordgrass (*Spartina Alterniflora*) loss and erosion have been caused
519 by combined effects of sea level rise and *S. reticulatum* density increases in US Atlantic salt
520 marshes (Crotty et al., 2020; Morrison et al., 2024).



Crabs create burrow structures in the form of tunnels and chimneys which can potentially provide material available for erosion. Flow velocities of 60 cm/s or higher are required to erode these structures, which can be reached at tidal creek heads under typical conditions (Farron et al., 2020) (Table S1, Fig. 3). These velocities are also likely during high flow events such as storms, which are expected to increase in frequency and intensity due to climate change (Zhang and Colle, 2018; Raw et al., 2023). Rainfall events, in contrast, do not erode marsh substrate that is consolidated but rather mobilize recently deposited, unconsolidated sediment (Voulgaris and Meyers, 2004). In areas that are heavily burrowed, this would include sediment deposited in the past few tidal cycles, in addition to burrow structures and pellets. This means that storms associated with climate change will have major effects on erosion patterns, especially in regions that are heavily burrowed, which can lead to morphological changes (Farron et al., 2020). Increases in drainage density is necessary to manage the expanding tidal prism and effectively drain the marsh surface to prevent waterlogging. Crab activity at tidal creeks may help alleviate the effects of accelerating sea level rise on the marsh platform (Farron et al., 2020). In a regime of increasing sea level rise, the presence of burrowing organisms, such as crabs, may possibly increase marsh sustainability, by forming creeks or extending existing creeks, and enhancing erosion. Overpopulation of crabs, through changes in predation pressure, however can cause loss of marsh area and increase vulnerability to erosion, negatively impacting the marsh.

1.7) Synthesis and way forward

A positive sediment budget is important for the accretion and resilience of tidal marshes, as it promotes marsh elevation and enhances carbon storage by actively burying carbon. Bioturbation activities on the other hand can either stabilize or destabilise sediment, influence sediment transport and ultimately influence marsh elevation. These two processes can therefore be viewed as being interconnected rather than being independent of one another. The reworking of sediment by some organisms increases surface roughness and decreases sediment cohesion, leading to erosion and in some cases creek formation. While the stabilization of sediment is possible through burrows of other species, functioning as passive sediment traps, which in turn can promote accretion. Apart from sediment properties being affected by bioturbation activities, carbon cycling is also influenced by these activities. Activities such as burrowing and feeding can lead to a loss of carbon through increased mineralization of organic matter, or through erosion. However, bioturbators can also promote the burial of carbon by trapping sediment, and transporting organic matter such as faeces and leaf litter into their burrows.



556 This review has highlighted a number of knowledge gaps, specifically the lack of
557 understanding of the influence that bioturbators and their interactions have on sediment
558 processes and their role in carbon cycling. This is despite increasing recognition that biological
559 components have an influence on the functioning of tidal marshes. Sediment–organism
560 interactions are often context specific and complex, and our understanding of species specific
561 impacts are limited. It is challenging to predict how bioturbators might influence their
562 environment as the impact of individual species on sediment dynamics varies, therefore,
563 bioturbation effects cannot be generalized. For example, the effects of crabs from the family
564 Ocypodidae versus crabs from the family Sesamidae will have different effects on sediment
565 because of burrow morphology, diet and behaviour, all of which influence bioturbation effects.
566 Moreover, these families are often found co-occurring in the same habitat making it important
567 to understand their individual as well as combined impacts on sediment processes. Such
568 studies could be done under experimental conditions and in situ, and should be extended
569 across different habitat types as sediment characteristics and vegetation also have an
570 influence on bioturbation impacts.

571 Sediment–species interactions also have an influence on carbon cycling in tidal marshes, yet
572 consumers are an understudied driver of these processes. There is a need to quantify carbon
573 stocks, sequestration and greenhouse gas fluxes and to investigate how these processes
574 respond to bioturbation activities. Studies comparing regions with varying intensities of
575 bioturbation are important for a better understanding of the contribution of bioturbators to
576 carbon dynamics in tidal marshes. It is clear that there is no real consensus as to whether
577 bioturbation has a positive or negative influence on sediment dynamics and carbon cycling
578 (Table 2 and S1). By advancing our understanding, management and restoration efforts could
579 be improved, and better predict the resilience of tidal marshes under future climate change
580 pressures.

581 1.8) Author contributions

582

583 Conceptualization: GMR, JBA; writing original draft preparation: LS; writing review and editing:
584 GMR, JBA, LS; supervision: GMR, JBA; funding acquisition: GMR, JBA.

585 1.9) Conflict of interest

586

587 The authors declare that they have no conflict of interest.



588 **1.10) Acknowledgements**

589

590 This research was funded by the National Research Foundation (NRF) of South Africa through
591 the support of the DSI/NRF Research Chair in Shallow Water Ecosystems (UID 84375). The
592 NRF are also thanked for providing a scholarship to LS. All opinions expressed are those of
593 the authors and not necessarily also those of the funders. Thank you to the Blue Carbon
594 Working Group for thought-provoking discussions that inspired this review.

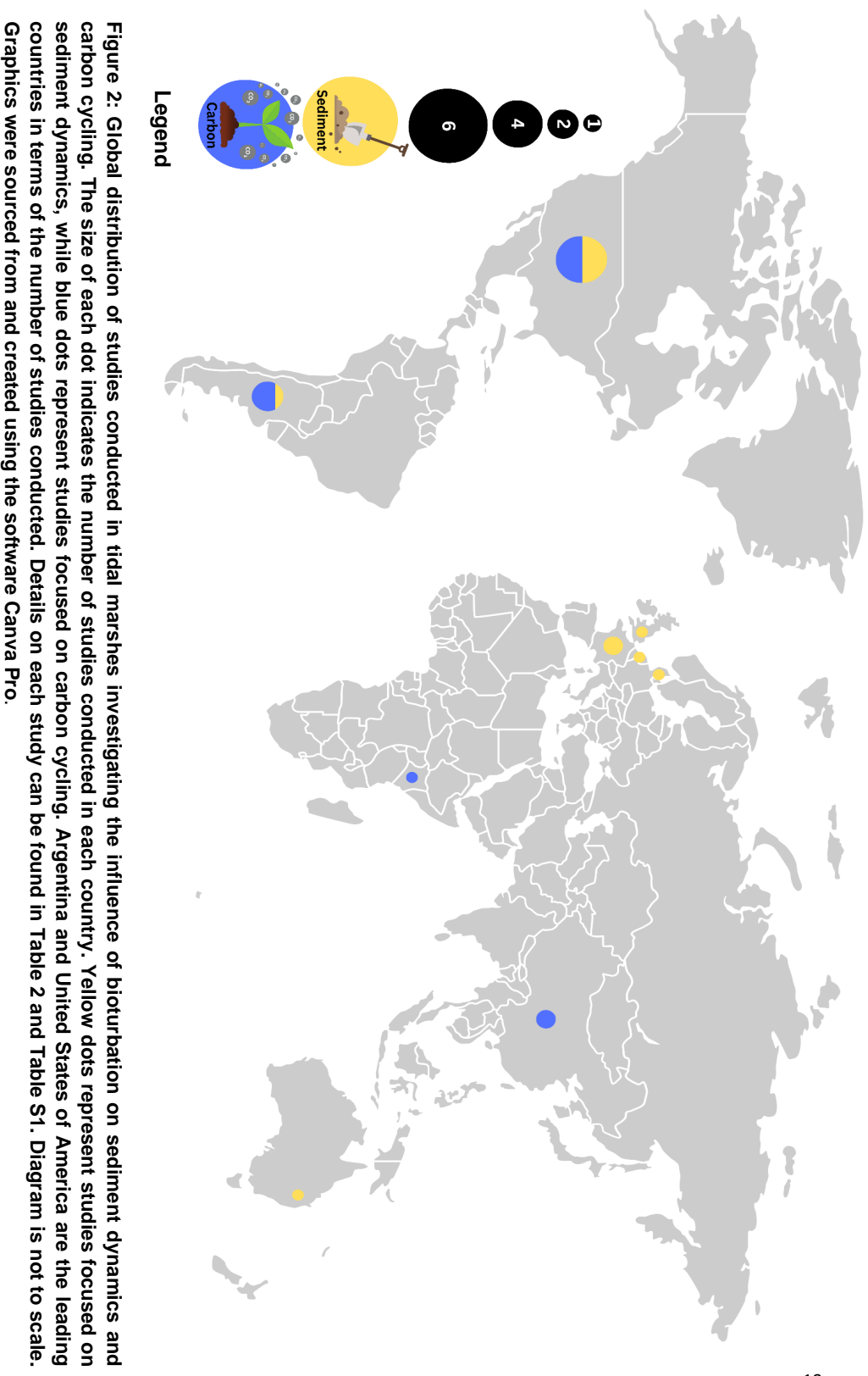


Table 2: Overview of the influence and directional effects of bioturbation on sediment and carbon dynamics across tidal habitats and continents. Adapted from Table S1.

Continent	Habitat type	Sediment/Carbon focused	Directional effect	Effect	Reference
Europe	Laboratory	Sediment	Negative	Erosion	Dairain et al., 2020
	Mudflat	Sediment	Negative	Decrease sediment stability	De Deckere et al., 2001
	Mudflat	Sediment	Negative	Decrease sediment stability	Montserat et al., 2011
	Mudflat	Sediment	Both negative & positive (species dependent)	One species caused erosion, one species caused accretion	Morelle et al., 2024
North America	Salt marsh	Carbon	Neutral to Positive	Neutral to positive effect on carbon sequestration	Graversen et al., 2022
	Salt marsh	Sediment	Negative	Change biogeochemistry of sediment	Wilson et al., 2012
	Laboratory	Sediment	Negative	Increase sediment roughness, decrease shear strength=erosion	Farron et al., 2020
	Salt marsh	Sediment	Positive	Accretion	Croft et al., 2023
South America	Salt marsh	Carbon	Negative	Hinders accretion, loss of carbon sequestration	Coverdale et al., 2014
	Salt marsh	Carbon	Negative	Decrease carbon stocks	Wittyngham et al., 2024
	Salt marsh	Carbon	Negative	Decrease carbon stocks	Montague, 1982
	Salt marsh & Mudflat	Sediment	Both negative & positive (species dependent)	One species caused sediment trapping=accretion, one species caused erosion	Escapa et al., 2008
Oceania	Mudflat	Carbon	Positive	Increase carbon sequestration	Botto et al., 2006
	Salt marsh	Carbon	Negative	Decrease carbon stocks	Guilérrez et al., 2006
	Salt marsh	Carbon	Negative	Decrease carbon stocks	Fanjul et al., 2015
	Mangrove & Salt marsh	Sediment	Neutral to positive	Increase surface elevation in mangroves, no influence in salt marsh	Bennion et al., 2024
Asia	Salt marsh	Carbon	Negative	Promote movement of carbon, can decrease long term storage of carbon	Wang et al., 2010
	Estuary	Carbon	Negative	Decrease carbon stocks	Nie et al., 2021
Africa	Mangrove	Carbon	Positive	Increase carbon storage	Andreita et al., 2014



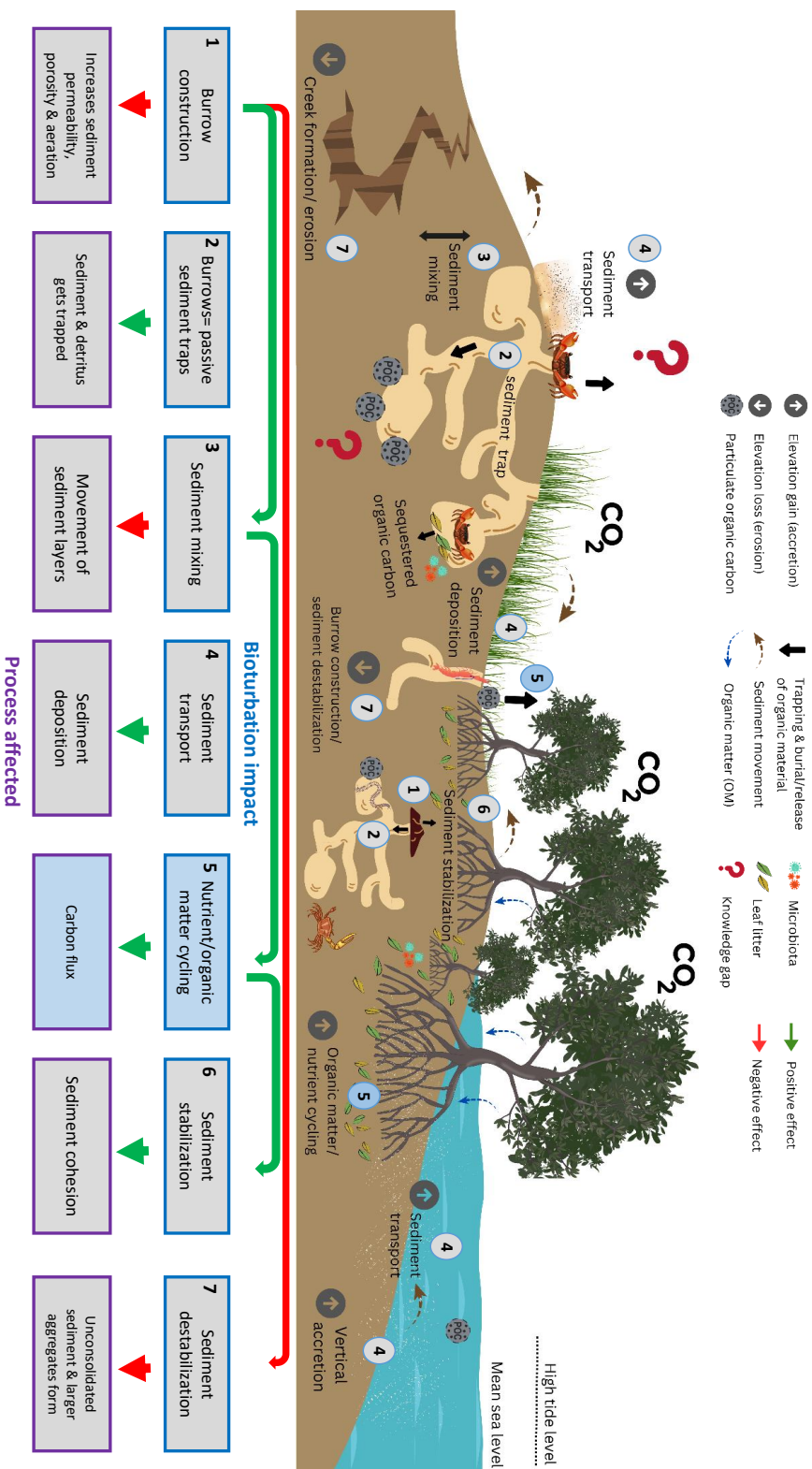


Figure 3: Conceptual diagram illustrating the processes influenced by bioturbators, specifically their impact on sediment dynamics and carbon cycling, and how they are linked. The knowledge gaps, indicated by question marks, relate to the role of benthic organisms in sediment dynamics, species specific effects, as well as the influence of bioturbators on the carbon cycle. The flow diagram indicates the graphics in terms of the bioturbation impacts and which processes are affected within the marsh, with green arrows indicating a positive effect and red indicating a negative effect. Diagram is not to scale. Graphics were sourced from and created using the software Canva Pro.



Table 3: Bioturbation impacts on sedimentation and carbon sequestration. Negative effects are italicised, while positive effects are indicated in bold. This table corresponds to the network diagram in Figure 3.

Bioturbation impact	Process	Effect on sedimentation	Effect on carbon sequestration
Sediment mixing	Movement of sediment layers	<i>Effects sediment structure- can lead to erosion^a</i>	<i>Excavate stored carbon^b</i>
Burrow construction	Increase sediment permeability, porosity & aeration	<i>Reduces sediment stability- can lead to erosion^c</i>	<i>Increases organic matter decomposition- decreases carbon sequestration^d</i>
Passive sediment trap (burrows)	Sediment and detritus gets trapped in and around burrow	Increases sediment deposition and overall concentration of sediment organic matter^e	Increases burial of organic rich sediment enhancing carbon storage^f
Sediment destabilization	Sediment becomes unconsolidated & larger aggregates are formed	<i>Increases sediment erosion^g</i>	<i>Decreases organic matter burial- decreases carbon sequestration^h</i>
Sediment transport	Sediment deposition	Causes particles to be resuspended and transportedⁱ - can contribute to accretion	If resuspended particles are trapped, it can increase carbon burial^j
Sediment stabilization	Some bioturbators promote sediment cohesion	Increases sediment strength and retention^k - can contribute to accretion	Sediment retention enhances carbon burial because active burial limits oxidation^l
Nutrient/ organic matter cycling	Organic matter transported to deeper layers	Influences plant growth^k which increases sediment trapping and stability- can contribute to accretion	Enhanced carbon storage^{e, l} – decomposition is slowed down

^a Dairain et al., 2020
^b Gutierrez et al. 2006
^c Grabowski et al. 2011
^d Nie et al., 2021
^e Botto and Iribarne 2000
^f Andreetta et al., 2014
^g Coverdale et al., 2014
^h Pillay et al., 2007
ⁱ Kristensen 2008
^j McCarthy et al., 2009
^k Botto et al., 2006



1.11) References

- Adams, J. B., Buttner, D., Hawkes, S., Human, L. R. D., Machite, A., Mfikili, A. N., Ndhlovu, A., Smit, L.-A., Rajkaran, A., Riddin, T., Rishworth, G. M., Van Deventer, H., Van Niekerk, L., Von Der Heyden, S., Whitfield, E. C., and Raw, J. L.: Blue Carbon at the southern tip of Africa: current knowledge and future perspectives for dynamic estuarine environments, *Estuarine, Coastal and Shelf Science*, 322, 109360, <https://doi.org/10.1016/j.ecss.2025.109360>, 2025.
- Agusto, L. E., Fratini, S., Jimenez, P. J., Quadros, A., and Cannicci, S.: Structural characteristics of crab burrows in Hong Kong mangrove forests and their role in ecosystem engineering, *Estuarine, Coastal and Shelf Science*, 248, 106973, <https://doi.org/10.1016/j.ecss.2020.106973>, 2021.
- Alongi, D. M.: Present state and future of the world's mangrove forests, *Environmental Conservation*, 29, 331–349, <https://doi.org/10.1017/S0376892902000231>, 2002.
- Alongi, D. M.: *Blue Carbon: Coastal Sequestration for Climate Change Mitigation*, Springer, Cham, 1 pp., 2018.
- Andreetta, A., Fusi, M., Cameldi, I., Cimò, F., Carnicelli, S., and Cannicci, S.: Mangrove carbon sink. Do burrowing crabs contribute to sediment carbon storage? Evidence from a Kenyan mangrove system, *Journal of Sea Research*, 85, 524–533, <https://doi.org/10.1016/j.seares.2013.08.010>, 2014.
- Arias-Ortiz, A., Serrano, O., Masqué, P., Lavery, P. S., Mueller, U., Kendrick, G. A., Rozaimi, M., Esteban, A., Fourqurean, J. W., Marbà, N., Mateo, M. A., Murray, K., Rule, M. J., and Duarte, C. M.: A marine heatwave drives massive losses from the world's largest seagrass carbon stocks, *Nature Climate Change*, 8, 338–344, <https://doi.org/10.1038/s41558-018-0096-y>, 2018.
- Arlinghaus, P., Zhang, W., Wrede, A., Schrum, C., and Neumann, A.: Impact of benthos on morphodynamics from a modeling perspective, *Earth-Science Reviews*, 221, 103803, <https://doi.org/10.1016/j.earscirev.2021.103803>, 2021.
- Barbier, E. B.: Valuing the storm protection service of estuarine and coastal ecosystems, *Ecosystem Services*, 11, 32–38, <https://doi.org/10.1016/j.ecoser.2014.06.010>, 2015.
- Beheshti, K. M., Wasson, K., Angelini, C., Silliman, B. R., and Hughes, B. B.: Long-term study reveals top-down effect of crabs on a California salt marsh, *Ecosphere*, 12, e03703, <https://doi.org/10.1002/ecs2.3703>, 2021.
- Bennion, V., Dwyer, J. M., Twomey, A. J., and Lovelock, C. E.: Decadal Trends in Surface Elevation and Tree Growth in Coastal Wetlands of Moreton Bay, Queensland, Australia, *Estuaries and Coasts*, 47, 1955–1971, <https://doi.org/10.1007/s12237-024-01325-y>, 2024.
- Bernal, B. and Mitsch, W. J.: Carbon Sequestration in Two Created Riverine Wetlands in the Midwestern United States, *Journal of Environmental Quality*, 42, 1236–1244, <https://doi.org/10.2134/jeq2012.0229>, 2013.
- Bertness, M. D.: Fiddler Crab Regulation of *Spartina alterniflora* Production on a New England Salt Marsh, *Ecology*, 66, 1042–1055, <https://doi.org/10.2307/1940564>, 1985.



- Booth, J. M., Fusi, M., Marasco, R., and Daffonchio, D.: The microbial landscape in bioturbated mangrove sediment: A resource for promoting nature-based solutions for mangroves, *Microbial Biotechnology*, 16, 1584–1602, <https://doi.org/10.1111/1751-7915.14273>, 2023.
- Borchert, S. M., Osland, M. J., Enwright, N. M., and Griffith, K. T.: Coastal wetland adaptation to sea level rise: Quantifying potential for landward migration and coastal squeeze, *Journal of Applied Ecology*, 55, 2876–2887, <https://doi.org/10.1111/1365-2664.13169>, 2018.
- Bos, A. R., Bouma, T. J., De Kort, G. L. J., and Van Katwijk, M. M.: Ecosystem engineering by annual intertidal seagrass beds: Sediment accretion and modification, *Estuarine, Coastal and Shelf Science*, 74, 344–348, <https://doi.org/10.1016/j.ecss.2007.04.006>, 2007.
- Botto, F. and Iribarne, O.: Contrasting Effects of Two Burrowing Crabs (*Chasmagnathus granulata* and *Uca uruguayensis*) on Sediment Composition and Transport in Estuarine Environments, *Estuarine, Coastal and Shelf Science*, 51, 141–151, <https://doi.org/10.1006/ecss.2000.0642>, 2000.
- Botto, F., Iribarne, O., Gutierrez, J., Bava, J., Gagliardini, A., and Valiela, I.: Ecological importance of passive deposition of organic matter into burrows of the SW Atlantic crab *Chasmagnathus granulatus*, *Marine Ecology Progress Series*, 312, 201–210, <https://doi.org/10.3354/meps312201>, 2006.
- Bradley, P. M. and Morris, J. T.: Influence of Oxygen and Sulfide Concentration on Nitrogen Uptake Kinetics in *Spartina Alterniflora*, *Ecology*, 71, 282–287, <https://doi.org/10.2307/1940267>, 1990.
- Branoff, B. L.: Mangrove Disturbance and Response Following the 2017 Hurricane Season in Puerto Rico, *Estuaries and Coasts*, 43, 1248–1262, <https://doi.org/10.1007/s12237-019-00585-3>, 2020.
- Buffington, K. J., Janousek, C. N., Thorne, K. M., and Dugger, B. D.: Spatiotemporal Patterns of Mineral and Organic Matter Deposition Across Two San Francisco Bay-Delta Tidal Marshes, *Wetlands*, 40, 1395–1407, <https://doi.org/10.1007/s13157-019-01259-3>, 2020.
- Cahoon, D. R.: Measuring and Interpreting the Surface and Shallow Subsurface Process Influences on Coastal Wetland Elevation: A Review, *Estuaries and Coasts*, 47, 1708–1734, <https://doi.org/10.1007/s12237-024-01332-z>, 2024.
- Carpenter, S., Evans, C., Pittman, S. J., Antonopoulou, M., Bejarano, I., Das, H. S., Möller, M., Peel, K., Samara, F., Stamoulis, K. A., and Mateos-Molina, D.: Multi-habitat carbon stock assessments to inform nature-based solutions for coastal seascapes in arid regions, *Frontiers in Marine Science*, 10, 1239904, <https://doi.org/10.3389/fmars.2023.1239904>, 2023.
- Chen, Z. L. and Lee, S. Y.: Tidal Flats as a Significant Carbon Reservoir in Global Coastal Ecosystems, *Frontiers in Marine Science*, 9, 900896, <https://doi.org/10.3389/fmars.2022.900896>, 2022.
- Coggan, N. V., Hayward, M. W., and Gibb, H.: A global database and “state of the field” review of research into ecosystem engineering by land animals, *Journal of Animal Ecology*, 87, 974–994, <https://doi.org/10.1111/1365-2656.12819>, 2018.



- Coverdale, T. C., Brisson, C. P., Young, E. W., Yin, S. F., Donnelly, J. P., and Bertness, M. D.: Indirect Human Impacts Reverse Centuries of Carbon Sequestration and Salt Marsh Accretion, *PLoS ONE*, 9, e93296, <https://doi.org/10.1371/journal.pone.0093296>, 2014.
- Cozzoli, F., Bouma, T. J., Ottolander, P., Lluch, M. S., Ysebaert, T., and Herman, P. M. J.: The combined influence of body size and density on cohesive sediment resuspension by bioturbators, *Scientific Reports*, 8, 3831, <https://doi.org/10.1038/s41598-018-22190-3>, 2018.
- Crotty, S. M., Ortals, C., Pettengill, T. M., Shi, L., Olabarrieta, M., Joyce, M. A., Altieri, A. H., Morrison, E., Bianchi, T. S., and Craft, C.: Sea-level rise and the emergence of a keystone grazer alter the geomorphic evolution and ecology of southeast US salt marshes, *Proceedings of the National Academy of Sciences*, 117, 17891–17902, 2020.
- Crotty, S. M., Pinton, D., Canestrelli, A., Fischman, H. S., Ortals, C., Dahl, N. R., Williams, S., Bouma, T. J., and Angelini, C.: Faunal engineering stimulates landscape-scale accretion in southeastern US salt marshes, *Nature Communications*, 14, 881, <https://doi.org/10.1038/s41467-023-36444-w>, 2023.
- Daborn, G., Amos, C., Brylinsky, M., Cristian, H., Drapeau, G., Faas, R., Grant, J., Long, B., Paterson, D., Perillo, G., and Piccolo, M.: An ecological cascade effect: migratory birds affect stability of intertidal sediments, *Limnology and Oceanography*, 38, 225–231, 1993.
- Dahl, M., Deyanova, D., Lyimo, L. D., Näslund, J., Samuelsson, G. S., Mtolera, M. S. P., Björk, M., and Gullström, M.: Effects of shading and simulated grazing on carbon sequestration in a tropical seagrass meadow, *Journal of Ecology*, 104, 654–664, <https://doi.org/10.1111/1365-2745.12564>, 2016.
- Dairain, A., Maire, O., Meynard, G., Richard, A., Rodolfo-Damiano, T., and Orvain, F.: Sediment stability: can we disentangle the effect of bioturbating species on sediment erodibility from their impact on sediment roughness? *Marine Environmental Research*, 162, 105147, <https://doi.org/10.1016/j.marenvres.2020.105147>, 2020.
- Darwin, C.: in: *The formation of vegetable mould through the action of worms with observation of their habits*, vol. 16, John Murray, London, 1881.
- Day, J. W., Kemp, G. P., Reed, D. J., Cahoon, D. R., Boumans, R. M., Suhayda, J. M., and Gambrell, R.: Vegetation death and rapid loss of surface elevation in two contrasting Mississippi delta salt marshes: The role of sedimentation, autocompaction and sea-level rise, *Ecological Engineering*, 37, 229–240, <https://doi.org/10.1016/j.ecoleng.2010.11.021>, 2011.
- De Deckere, E. M. G. T., Tolhurst, T. J., and De Brouwer, J. F. C.: Destabilization of Cohesive Intertidal Sediments by Infauna, *Estuarine, Coastal and Shelf Science*, 53, 665–669, <https://doi.org/10.1006/ecss.2001.0811>, 2001.
- Decho, A. W.: Microbial biofilms in intertidal systems: an overview, *Continental Shelf Research*, 20, 1257–1273, [https://doi.org/10.1016/S0278-4343\(00\)00022-4](https://doi.org/10.1016/S0278-4343(00)00022-4), 2000.
- 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, *Nature Geoscience*, 4, 293–297, <https://doi.org/10.1038/ngeo1123>, 2011.



- Duarte, C. M.: Reviews and syntheses: Hidden forests, the role of vegetated coastal habitats in the ocean carbon budget, *Biogeosciences*, 14, 301–310, <https://doi.org/10.5194/bg-14-301-2017>, 2017.
- Duarte De Paula Costa, M. and Macreadie, P. I.: The Evolution of Blue Carbon Science, *Wetlands*, 42, 109, <https://doi.org/10.1007/s13157-022-01628-5>, 2022.
- Egawa, R., Sharma, S., Nadaoka, K., and MacKenzie, R. A.: Burrow dynamics of crabs in subtropical estuarine mangrove forest, *Estuarine, Coastal and Shelf Science*, 252, 107244, <https://doi.org/10.1016/j.ecss.2021.107244>, 2021.
- Enwright, N. M., Griffith, K. T., and Osland, M. J.: Barriers to and opportunities for landward migration of coastal wetlands with sea-level rise, *Frontiers in Ecology & the Environment*, 14, 307–316, <https://doi.org/10.1002/fee.1282>, 2016.
- Escapa, M., Perillo, G. M. E., and Iribarne, O.: Sediment dynamics modulated by burrowing crab activities in contrasting SW Atlantic intertidal habitats, *Estuarine, Coastal and Shelf Science*, 80, 365–373, <https://doi.org/10.1016/j.ecss.2008.08.020>, 2008.
- Fagherazzi, S., Anisfeld, S. C., Blum, L. K., Long, E. V., Feagin, R. A., Fernandes, A., Kearney, W. S., and Williams, K.: Sea Level Rise and the Dynamics of the Marsh-Upland Boundary, *Front. Environ. Sci.*, 7, 25, <https://doi.org/10.3389/fenvs.2019.00025>, 2019.
- Fanjul, E., Grela, M., and Iribarne, O.: Effects of the dominant SW Atlantic intertidal burrowing crab *Chasmagnathus granulatus* on sediment chemistry and nutrient distribution, *Marine Ecology Progress Series*, 341, 177–190, <https://doi.org/10.3354/meps341177>, 2007.
- Fanjul, E., Escapa, M., Montemayor, D., Addino, M., Alvarez, M. F., Grela, M. A., and Iribarne, O.: Effect of crab bioturbation on organic matter processing in South West Atlantic intertidal sediments, *Journal of Sea Research*, 95, 206–216, <https://doi.org/10.1016/j.seares.2014.05.005>, 2015.
- Farron, S. J., Hughes, Z. J., FitzGerald, D. M., and Strom, K. B.: The impacts of bioturbation by common marsh crabs on sediment erodibility: A laboratory flume investigation, *Estuarine, Coastal and Shelf Science*, 238, 106710, <https://doi.org/10.1016/j.ecss.2020.106710>, 2020.
- Fourqurean, J. W., Duarte, C. M., Kennedy, H., Marbà, N., Holmer, M., Mateo, M. A., Apostolaki, E. T., Kendrick, G. A., Krause-Jensen, D., McGlathery, K. J., and Serrano, O.: Seagrass ecosystems as a globally significant carbon stock, *Nature Geoscience*, 5, 505–509, <https://doi.org/10.1038/ngeo1477>, 2012.
- François, F., Gerino, M., Stora, G., Durbec, J.-P., and Poggiale, J.-C.: Functional approach to sediment reworking by gallery-forming macrobenthic organisms: modeling and application with the polychaete *Nereis diversicolor*, *Marine Ecology Progress Series*, 229, 127–136, 2002.
- Giraldes, B. W., Al-Maslamani, I., and Smyth, D.: A new species of leucosiid crab (Decapoda: Brachyura: Leucosiidae) from the Arabian Gulf, *Zootaxa*, 4250, <https://doi.org/10.11646/zootaxa.4250.4.9>, 2017.
- Grabowski, R. C., Droppo, I. G., and Wharton, G.: Erodibility of cohesive sediment: The importance of sediment properties, *Earth-Science Reviews*, 105, 101–120, <https://doi.org/10.1016/j.earscirev.2011.01.008>, 2011.



- Graversen, A. E. L., Banta, G. T., Masque, P., and Krause-Jensen, D.: Carbon sequestration is not inhibited by livestock grazing in Danish salt marshes, *Limnology & Oceanography*, 67, <https://doi.org/10.1002/lno.12011>, 2022.
- Grigg, N. J.: Benthic bulldozers and pumps: laboratory and modelling studies of bioturbation and bioirrigation, 2003.
- Guimond, J. A., Seyfferth, A. L., Moffett, K. B., and Michael, H. A.: A physical-biogeochemical mechanism for negative feedback between marsh crabs and carbon storage, *Environmental Research Letters*, 15, 034024, <https://doi.org/10.1088/1748-9326/ab60e2>, 2020.
- Gutiérrez, J. L., Jones, C. G., Groffman, P. M., Findlay, S. E. G., Iribarne, O. O., Ribeiro, P. D., and Bruschetti, C. M.: The Contribution of Crab Burrow Excavation to Carbon Availability in Surficial Salt-marsh Sediments, *Ecosystems*, 9, 647–658, <https://doi.org/10.1007/s10021-006-0135-9>, 2006.
- Hajjalizadeh, P., Safaie, M., Naderloo, R., and Shojaei, M. G.: Spatial and Temporal Distribution of Brachyuran Crabs in Mangroves of the Persian Gulf, *Wetlands*, 42, 99, <https://doi.org/10.1007/s13157-022-01623-w>, 2022.
- Hamilton, S. E. and Casey, D.: Creation of a high spatio-temporal resolution global database of continuous mangrove forest cover for the 21st century (CGMFC-21), *Global Ecology and Biogeography*, 25, 729–738, <https://doi.org/10.1111/geb.12449>, 2016.
- Hatje, V., Masqué, P., Patire, V. F., Dórea, A., and Barros, F.: Blue carbon stocks, accumulation rates, and associated spatial variability in Brazilian mangroves, *Limnology & Oceanography*, 66, 321–334, <https://doi.org/10.1002/lno.11607>, 2021.
- Hawkins, S. J., Allcock, A. L., Bates, A. E., Evans, A. J., Firth, L. B., McQuaid, C. D., Russell, B. D., Smith, I. P., Swearer, S. E., and Todd, P. A. (Eds.): *Oceanography and Marine Biology: An Annual Review*, Volume 58, Taylor & Francis, Erscheinungsort nicht ermittelbar, 1 pp., 2020.
- He, Q., Li, Z., Daleo, P., Lefcheck, J. S., Thomsen, M. S., Adams, J. B., and Bouma, T. J.: Coastal wetland resilience through local, regional and global conservation, *Nature Reviews Biodiversity*, 1, 50–67, 2025.
- Holdredge, C., Bertness, M., Herrmann, N., and Gedan, K.: Fiddler crab control of cordgrass primary production in sandy sediments, *Marine Ecology Progress Series*, 399, 253–259, <https://doi.org/10.3354/meps08331>, 2010.
- Horstman, E. M., Dohmen-Janssen, C. M., Bouma, T. J., and Hulscher, S. J. M. H.: Tidal-scale flow routing and sedimentation in mangrove forests: Combining field data and numerical modelling, *Geomorphology*, 228, 244–262, <https://doi.org/10.1016/j.geomorph.2014.08.011>, 2015.
- Kauffman, J. B., Adame, M. F., Arifanti, V. B., Schile-Beers, L. M., Bernardino, A. F., Bhomia, R. K., Donato, D. C., Feller, I. C., Ferreira, T. O., and Jesus García, M. del C.: Total ecosystem carbon stocks of mangroves across broad global environmental and physical gradients, *Ecological monographs*, 90, e01405, 2020.
- Keller, D. P., Lenton, A., Littleton, E. W., Oschlies, A., Scott, V., and Vaughan, N. E.: The Effects of Carbon Dioxide Removal on the Carbon Cycle, *Current Climate Change Reports*, 4, 250–265, <https://doi.org/10.1007/s40641-018-0104-3>, 2018.



- Kelleway, J. J., Saintilan, N., Macreadie, P. I., Skilbeck, C. G., Zawadzki, A., and Ralph, P. J.: Seventy years of continuous encroachment substantially increases 'blue carbon' capacity as mangroves replace intertidal salt marshes, *Global Change Biology*, 22, 1097–1109, <https://doi.org/10.1111/gcb.13158>, 2016.
- Kelleway, J. J., Cavanaugh, K., Rogers, K., Feller, I. C., Ens, E., Doughty, C., and Saintilan, N.: Review of the ecosystem service implications of mangrove encroachment into salt marshes, *Global Change Biology*, 23, 3967–3983, <https://doi.org/10.1111/gcb.13727>, 2017a.
- Kelleway, J. J., Saintilan, N., Macreadie, P. I., Baldock, J. A., and Ralph, P. J.: Sediment and carbon deposition vary among vegetation assemblages in a coastal salt marsh, *Biogeosciences*, 14, 3763–3779, <https://doi.org/10.5194/bg-14-3763-2017>, 2017b.
- Kinoshita, K., Wada, M., Kogure, K., and Furota, T.: Microbial activity and accumulation of organic matter in the burrow of the mud shrimp, *Upogebia major* (Crustacea: Thalassinidea), *Marine Biology*, 153, 277–283, 2008.
- Kirwan, M. L. and Guntenspergen, G. R.: Feedbacks between inundation, root production, and shoot growth in a rapidly submerging brackish marsh, *Journal of Ecology*, 100, 764–770, <https://doi.org/10.1111/j.1365-2745.2012.01957.x>, 2012.
- Klaassen, M., Moosdorf, N., and Zimmer, M.: Combined effect of burrowing mangrove crabs and tides on carbon fluxes, *Bulletin of Marine Science*, 101, 1221–1236, <https://doi.org/10.5343/bms.2024.0012>, 2025.
- Krauss, K. W., McKee, K. L., Lovelock, C. E., Cahoon, D. R., Saintilan, N., Reef, R., and Chen, L.: How mangrove forests adjust to rising sea level, *New phytologist*, 202, 19–34, 2014.
- Kristensen, E.: Impact of polychaetes (*Nereis* spp. and *Arenicola marina*) on carbon biogeochemistry in coastal marine sediments, *Geochemical Transactions*, 2, 92, <https://doi.org/10.1186/1467-4866-2-92>, 2001.
- Kristensen, E.: Mangrove crabs as ecosystem engineers; with emphasis on sediment processes, *Journal of Sea Research*, 59, 30–43, <https://doi.org/10.1016/j.seares.2007.05.004>, 2008.
- Kristensen, E., Bouillon, S., Dittmar, T., and Marchand, C.: Organic carbon dynamics in mangrove ecosystems: A review, *Aquatic Botany*, 89, 201–219, <https://doi.org/10.1016/j.aquabot.2007.12.005>, 2008.
- Kristensen, E., Penha-Lopes, G., Delefosse, M., Valdemarsen, T., Quintana, C., and Banta, G.: What is bioturbation? The need for a precise definition for fauna in aquatic sciences, *Marine Ecology Progress Series*, 446, 285–302, <https://doi.org/10.3354/meps09506>, 2012.
- Le Hir, P., Monbet, Y., and Orvain, F.: Sediment erodability in sediment transport modelling: Can we account for biota effects? *Continental Shelf Research*, 27, 1116–1142, <https://doi.org/10.1016/j.csr.2005.11.016>, 2007.
- Leonardi, N., Carnacina, I., Donatelli, C., Ganju, N. K., Plater, A. J., Schuerch, M., and Temmerman, S.: Dynamic interactions between coastal storms and salt marshes: A review, *Geomorphology*, 301, 92–107, <https://doi.org/10.1016/j.geomorph.2017.11.001>, 2018.



- Li, S., Cui, B., Xie, T., Bai, J., Wang, Q., and Shi, W.: What drives the distribution of crab burrows in different habitats of intertidal salt marshes, Yellow River Delta, China, *Ecological Indicators*, 92, 99–106, <https://doi.org/10.1016/j.ecolind.2017.11.003>, 2018.
- Lovelock, C. E. and Duarte, C. M.: Dimensions of Blue Carbon and emerging perspectives, *Biology Letters*, 15, 20180781, <https://doi.org/10.1098/rsbl.2018.0781>, 2019.
- Lovelock, C. E. and Reef, R.: Variable Impacts of Climate Change on Blue Carbon, *One Earth*, 3, 195–211, <https://doi.org/10.1016/j.oneear.2020.07.010>, 2020.
- Lovelock, C. E., Feller, I. C., Ellis, J., Schwarz, A. M., Hancock, N., Nichols, P., and Sorrell, B.: Mangrove growth in New Zealand estuaries: the role of nutrient enrichment at sites with contrasting rates of sedimentation, *Oecologia*, 153, 633–641, <https://doi.org/10.1007/s00442-007-0750-y>, 2007.
- Lynch, A. J., Cooke, S. J., Arthington, A. H., Baigun, C., Bossenbroek, L., Dickens, C., Harrison, I., Kimirei, I., Langhans, S. D., Murchie, K. J., Olden, J. D., Ormerod, S. J., Owuor, M., Raghavan, R., Samways, M. J., Schinegger, R., Sharma, S., Tachamo-Shah, R., Tickner, D., Tweddle, D., Young, N., and Jähnig, S. C.: People need freshwater biodiversity, *WIREs Water*, 10, e1633, <https://doi.org/10.1002/wat2.1633>, 2023.
- MacKenzie, R., Sharma, S., and Rovai, A. R.: Environmental drivers of blue carbon burial and soil carbon stocks in mangrove forests, in: *Dynamic Sedimentary Environments of Mangrove Coasts*, Elsevier, 275–294, <https://doi.org/10.1016/B978-0-12-816437-2.00006-9>, 2021.
- MacKenzie, R. A., Krauss, K. W., Cormier, N., Eperiam, E., Van Aardt, J., Kargar, A. R., Grow, J., and Klump, J. V.: Relative Effectiveness of a Radionuclide (^{210}Pb), Surface Elevation Table (SET), and LiDAR At Monitoring Mangrove Forest Surface Elevation Change, *Estuaries and Coasts*, 47, 2080–2092, <https://doi.org/10.1007/s12237-023-01301-y>, 2024.
- Macreadie, P. I., Baird, M. E., Trevathan-Tackett, S. M., Larkum, A. W. D., and Ralph, P. J.: Quantifying and modelling the carbon sequestration capacity of seagrass meadows – A critical assessment, *Marine Pollution Bulletin*, 83, 430–439, <https://doi.org/10.1016/j.marpolbul.2013.07.038>, 2014.
- Macreadie, P. I., Nielsen, D. A., Kelleway, J. J., Atwood, T. B., Seymour, J. R., Petrou, K., Connolly, R. M., Thomson, A. C., Trevathan-Tackett, S. M., and Ralph, P. J.: Can we manage coastal ecosystems to sequester more blue carbon? *Frontiers in Ecology & the Environment*, 15, 206–213, <https://doi.org/10.1002/fee.1484>, 2017.
- Macreadie, P. I., Costa, M. D., Atwood, T. B., Friess, D. A., Kelleway, J. J., Kennedy, H., Lovelock, C. E., Serrano, O., and Duarte, C. M.: Blue carbon as a natural climate solution, *Nature Reviews Earth & Environment*, 2, 826–839, 2021.
- Maxwell, T. L., Spalding, M. D., Friess, D. A., Murray, N. J., Rogers, K., Rovai, A. S., Smart, L. S., Weilguny, L., Adame, M. F., and Adams, J. B.: Soil carbon in the world's tidal marshes, *Nature Communications*, 15, 10265, 2024.
- McCarty, G., Pachepsky, Y., and Ritchie, J.: Impact of Sedimentation on Wetland Carbon Sequestration in an Agricultural Watershed, *Journal of Environmental Quality*, 38, 804–813, <https://doi.org/10.2134/jeq2008.0012>, 2009.



- McKenzie, L. J., Nordlund, L. M., Jones, B. L., Cullen-Unsworth, L. C., Roelfsema, C., and Unsworth, R. K.: The global distribution of seagrass meadows, *Environmental Research Letters*, 15, 074041, 2020.
- Mcleod, E., Poulter, B., Hinkel, J., Reyes, E., and Salm, R.: Sea-level rise impact models and environmental conservation: A review of models and their applications, *Ocean & Coastal Management*, 53, 507–517, <https://doi.org/10.1016/j.ocecoaman.2010.06.009>, 2010.
- 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, *Frontiers in Ecology & the Environment*, 9, 552–560, <https://doi.org/10.1890/110004>, 2011.
- Mcowen, C., Weatherdon, L., Bochove, J.-W., Sullivan, E., Blyth, S., Zockler, C., Stanwell-Smith, D., Kingston, N., Martin, C., Spalding, M., and Fletcher, S.: A global map of saltmarshes, *Biodiversity Data Journal*, 5, e11764, <https://doi.org/10.3897/BDJ.5.e11764>, 2017.
- Mentaschi, L., Voudoukas, M. I., Pekel, J.-F., Voukouvalas, E., and Feyen, L.: Global long-term observations of coastal erosion and accretion, *Scientific Reports*, 8, 12876, <https://doi.org/10.1038/s41598-018-30904-w>, 2018.
- Meysman, F. J., Middelburg, J. J., and Heip, C. H.: Bioturbation: a fresh look at Darwin's last idea, *Trends in Ecology & Evolution*, 21, 688–695, 2006.
- Min, W. W., Kandasamy, K., and Balakrishnan, B.: Crab Species-Specific Excavation and Architecture of Burrows in Restored Mangrove Habitat, *JMSE*, 11, 310, <https://doi.org/10.3390/jmse11020310>, 2023.
- Montague, C. L.: The influence of fiddler crab burrows and burrowing on metabolic processes in salt marsh sediments, in: *Estuarine Comparisons*, Elsevier, 283–301, <https://doi.org/10.1016/B978-0-12-404070-0.50023-5>, 1982.
- Montserrat, F., Suykerbuyk, W., Al-Busaidi, R., Bouma, T. J., Van Der Wal, D., and Herman, P. M. J.: Effects of mud sedimentation on lugworm ecosystem engineering, *Journal of Sea Research*, 65, 170–181, <https://doi.org/10.1016/j.seares.2010.09.003>, 2011.
- Morelle, J., Huguet, A., Richard, A., Laverman, A. M., Roose-Amsaleg, C., Parlanti, E., Sourzac, M., Mesnage, V., Lecoq, N., Deloffre, J., Viollier, E., Maire, O., and Orvain, F.: Antagonistic impacts of benthic bioturbator species: Interconnected effects on sedimentary properties, biogeochemical variables, and microbial dynamics, *Journal of Experimental Marine Biology and Ecology*, 573, 152000, <https://doi.org/10.1016/j.jembe.2024.152000>, 2024.
- Morris, J. T., Sundareshwar, P., Nietch, C. T., Kjerfve, B., and Cahoon, D. R.: Responses of coastal wetlands to rising sea level, *Ecology*, 83, 2869–2877, 2002.
- Morrisey, D. J., Swales, A., Dittmann, S., Morrison, M. A., Lovelock, C. E., and Beard, C. M.: The ecology and management of temperate mangroves, *Oceanography and marine biology: an annual review*, 48, 43–160, 2010.
- Morrison, E. S., Bianchi, T. S., Kenney, W. F., Brenner, M., Prince, K., Williams, S., Ortals, C., Cordero, O., Crotty, S. M., and Angelini, C.: Influence of the Keystone Grazer, *Sesarma reticulatum*, on the Hydrology and Organic Matter Cycling in Salt Marshes



- of the Southeastern USA, *Estuaries and Coasts*, 47, 994–1011, <https://doi.org/10.1007/s12237-024-01336-9>, 2024.
- Mudd, S. M., D'Alpaos, A., and Morris, J. T.: How does vegetation affect sedimentation on tidal marshes? Investigating particle capture and hydrodynamic controls on biologically mediated sedimentation, *Journal of Geophysical Research: Earth Surface*, 115, 2010.
- Murray, N. J., Phinn, S. R., DeWitt, M., Ferrari, R., Johnston, R., Lyons, M. B., Clinton, N., Thau, D., and Fuller, R. A.: The global distribution and trajectory of tidal flats, *Nature*, 565, 222–225, 2019.
- Nahlik, A. M. and Mitsch, W. J.: The Effect of River Pulsing on Sedimentation and Nutrients in Created Riparian Wetlands, *Journal of Environmental Quality*, 37, 1634–1643, <https://doi.org/10.2134/jeq2007.0116>, 2008.
- Needham, H. R., Pilditch, C. A., Lohrer, A. M., and Thrush, S. F.: Density and habitat dependent effects of crab burrows on sediment erodibility, *Journal of Sea Research*, 76, 94–104, <https://doi.org/10.1016/j.seares.2012.12.004>, 2013.
- Nellemann, C. and Corcoran, E.: Blue carbon: the role of healthy oceans in binding carbon: a rapid response assessment, UNEP/Earthprint, 2009.
- Ngo-Massou, V. M., Din, N., Kenne, M., and Dongmo, A. B.: Brachyuran crab diversity and abundance patterns in the mangroves of Cameroon, *Regional Studies in Marine Science*, 24, 324–335, <https://doi.org/10.1016/j.rsma.2018.09.010>, 2018.
- Nie, L., Li, Y., Hou, Y., Di, L., Xi, M., and Yu, Z.: Dynamics of organic carbon under bioturbation by mud crabs (*Macrophthalmus japonicus*) and clamworms (*Perinereis aibuhitensis*) in an estuary ecosystem, *Journal of Experimental Marine Biology and Ecology*, 534, 151474, <https://doi.org/10.1016/j.jembe.2020.151474>, 2021.
- Ouyang, X., Connolly, R. M., and Lee, S. Y.: Revised global estimates of resilience to sea level rise for tidal marshes, *Environmental Challenges*, 9, 100593, 2022.
- Pan, F., Xiao, K., Cai, Y., Li, H., Guo, Z., Wang, X., Zheng, Y., Zheng, C., Bostick, B. C., and Michael, H. A.: Integrated effects of bioturbation, warming and sea-level rise on mobility of sulfide and metalloids in sediment porewater of mangrove wetlands, *Water Research*, 233, 119788, <https://doi.org/10.1016/j.watres.2023.119788>, 2023.
- Passeri, D. L., Hagen, S. C., Medeiros, S. C., Bilske, M. V., Alizad, K., and Wang, D.: The dynamic effects of sea level rise on low-gradient coastal landscapes: A review, *Earth's Future*, 3, 159–181, <https://doi.org/10.1002/2015EF000298>, 2015.
- 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, <https://doi.org/10.1371/journal.pone.0043542>, 2012.
- Penha-Lopes, G., Bartolini, F., Limbu, S., Cannicci, S., Kristensen, E., and Paula, J.: Are fiddler crabs potentially useful ecosystem engineers in mangrove wastewater wetlands? *Marine Pollution Bulletin*, 58, 1694–1703, <https://doi.org/10.1016/j.marpolbul.2009.06.015>, 2009.



- Perkins, M. J., Ng, T. P. T., Dudgeon, D., Bonebrake, T. C., and Leung, K. M. Y.: Conserving intertidal habitats: What is the potential of ecological engineering to mitigate impacts of coastal structures? *Estuarine, Coastal and Shelf Science*, 167, 504–515, <https://doi.org/10.1016/j.ecss.2015.10.033>, 2015.
- Phillips, J. D.: Coastal wetlands, sea level, and the dimensions of geomorphic resilience, *Geomorphology*, 305, 173–184, <https://doi.org/10.1016/j.geomorph.2017.03.022>, 2018.
- Pillay, D. and Branch, G.: Bioengineering effects of burrowing thalassinidean shrimps on marine soft-bottom ecosystems, *Oceanography and Marine Biology: An Annual Review*, 49, 137–192, 2011.
- Pillay, D., Branch, G. M., and Forbes, A. T.: The influence of bioturbation by the sandprawn *Callinassa kraussi* on feeding and survival of the bivalve *Eumarcia paupercula* and the gastropod *Nassarius kraussianus*, *Journal of Experimental Marine Biology and Ecology*, 344, 1–9, 2007.
- Radabaugh, K. R., Moyer, R. P., Chappel, A. R., Powell, C. E., Bociu, I., Clark, B. C., and Smoak, J. M.: Coastal Blue Carbon Assessment of Mangroves, Salt Marshes, and Salt Barrens in Tampa Bay, Florida, USA, *Estuaries and Coasts*, 41, 1496–1510, <https://doi.org/10.1007/s12237-017-0362-7>, 2018.
- Raw, J., Tsipa, V., Banda, S., Riddin, T., van Niekerk, L., and Adams, J.: Scoping Study: A Blue Carbon Sinks Assessment for South Africa, 2021.
- Raw, J. L., Van Der Stocken, T., Carroll, D., Harris, L. R., Rajkaran, A., Van Niekerk, L., and Adams, J. B.: Dispersal and coastal geomorphology limit potential for mangrove range expansion under climate change, *Journal of Ecology*, 111, 139–155, <https://doi.org/10.1111/1365-2745.14020>, 2023.
- Ren, L., Jensen, K., Porada, P., and Mueller, P.: Biota-mediated carbon cycling—A synthesis of biotic-interaction controls on blue carbon, *Ecology Letters*, 25, 521–540, <https://doi.org/10.1111/ele.13940>, 2022.
- Rinehart, S. A., Dybiec, J. M., Walker, J. B., Simpson, L., and Cherry, J. A.: Effects of burrowing crabs on coastal sediments and their functions: A systematic meta-analysis, *Ecosphere*, 15, e4927, <https://doi.org/10.1002/ecs2.4927>, 2024.
- Rogers, K., Kelleway, J. J., Saintilan, N., Megonigal, J. P., Adams, J. B., Holmquist, J. R., Lu, M., Schile-Beers, L., Zawadzki, A., Mazumder, D., and Woodroffe, C. D.: Wetland carbon storage controlled by millennial-scale variation in relative sea-level rise, *Nature*, 567, 91–95, <https://doi.org/10.1038/s41586-019-0951-7>, 2019.
- Saintilan, N., Rogers, K., Mazumder, D., and Woodroffe, C.: Allochthonous and autochthonous contributions to carbon accumulation and carbon store in southeastern Australian coastal wetlands, *Estuarine, Coastal and Shelf Science*, 128, 84–92, <https://doi.org/10.1016/j.ecss.2013.05.010>, 2013.
- Saintilan, N., Kovalenko, K., Guntenspergen, G., Rogers, K., Lynch, J., Cahoon, D., Lovelock, C., Friess, D., Ashe, E., Krauss, K., Cormier, N., Spencer, T., Adams, J., Raw, J., Ibanez, C., Scarton, F., Temmerman, S., Meire, P., Maris, T., Thorne, K., Brazner, J., Chmura, G., Bowron, T., Gamage, V., Cressman, K., Endris, C., Marconi, C., Marcum, P., Laurent, K., Reay, W., Raposa, K., Garwood, J., and Khan,



- N.: Constraints on the adjustment of tidal marshes to accelerating sea level rise, *Science*, 377, 523–527, 2022.
- Sarker, S., Masud-Ul-Alam, M., Hossain, M. S., Rahman Chowdhury, S., and Sharifuzzaman, S.: A review of bioturbation and sediment organic geochemistry in mangroves, *Geological Journal*, 56, 2439–2450, <https://doi.org/10.1002/gj.3808>, 2021.
- Schile, L. M., Kauffman, J. B., Crooks, S., Fourqurean, J. W., Glavan, J., and Megonigal, J. P.: Limits on carbon sequestration in arid blue carbon ecosystems, *Ecological Applications*, 27, 859–874, <https://doi.org/10.1002/eap.1489>, 2017.
- Schuerch, M., Spencer, T., Temmerman, S., Kirwan, M. L., Wolff, C., Lincke, D., Mcowen, C., Pickering, M., Reef, R., Vafeidis, A., Hinkel, J., Nicholls, R., and Brown, S.: Future response of global coastal wetlands to sea-level rise, *Nature*, 561, 231–234, 2018.
- Sen, S. and Homechaudhuri, S.: Comparative Burrow Architectures of Resident Fiddler Crabs (Ocypodidae) in Indian Sundarban Mangroves to Assess Their Suitability as Bioturbating Agents, *Proceedings of the Zoological Society*, 71, 17–24, <https://doi.org/10.1007/s12595-016-0178-7>, 2016.
- Sharbaugh, M., Rinehart, S. A., Dybiec, J. M., and Cherry, J. A.: Burrowing Crab Effects on Plants and Sediments Peak in Early Successional Ecotones Along A Chronosequence of Tidal Marsh Restoration, *Ecosystems*, 28, 35, <https://doi.org/10.1007/s10021-025-00980-x>, 2025.
- Sitch, S., Friedlingstein, P., Gruber, N., Jones, S. D., Murray-Tortarolo, G., Ahlström, A., Doney, S. C., Graven, H., Heinze, C., Huntingford, C., Levis, S., Levy, P. E., Lomas, M., Poulter, B., Viovy, N., Zaehle, S., Zeng, N., Armeth, A., Bonan, G., Bopp, L., Canadell, J. G., Chevallier, F., Ciais, P., Ellis, R., Gloor, M., Peylin, P., Piao, S. L., Le Quéré, C., Smith, B., Zhu, Z., and Myneni, R.: Recent trends and drivers of regional sources and sinks of carbon dioxide, *Biogeosciences*, 12, 653–679, <https://doi.org/10.5194/bg-12-653-2015>, 2015.
- Smit, L.-A., Adams, J. B., Hawkes, S. A., Peer, N., and Rishworth, G. M.: Proportional top-down effects of grapsoid crabs on growth of *Spartina maritima* cordgrass in southern African salt marshes, *Marine Ecology Progress Series*, 739, 49–64, 2024.
- Sogard, S. M. and Able, K. W.: A comparison of eelgrass, sea lettuce macroalgae, and marsh creeks as habitats for epibenthic fishes and decapods, *Estuarine, Coastal and Shelf Science*, 33, 501–519, [https://doi.org/10.1016/0272-7714\(91\)90087-R](https://doi.org/10.1016/0272-7714(91)90087-R), 1991.
- Spalding, M., Kainuma, M., and Collins, L.: *World Atlas of Mangroves*, Routledge, <https://doi.org/10.4324/9781849776608>, 2010.
- Spivak, A. C., Sanderman, J., Bowen, J. L., Canuel, E. A., and Hopkins, C. S.: Global-change controls on soil-carbon accumulation and loss in coastal vegetated ecosystems, *Nature Geoscience*, 12, 685–692, <https://doi.org/10.1038/s41561-019-0435-2>, 2019.
- Van Ardenne, L. B., Jolicouer, S., Bérubé, D., Burdick, D., and Chmura, G. L.: The importance of geomorphic context for estimating the carbon stock of salt marshes, *Geoderma*, 330, 264–275, <https://doi.org/10.1016/j.geoderma.2018.06.003>, 2018.
- Van Der Wal, D. and Herman, P. M. J.: Ecosystem Engineering Effects of *Aster tripolium* and *Salicornia procumbens* Salt Marsh on Macrofaunal Community Structure,



- Estuaries and Coasts, 35, 714–726, <https://doi.org/10.1007/s12237-011-9465-8>, 2012.
- Van Katwijk, M. M., Thorhaug, A., Marbà, N., Orth, R. J., Duarte, C. M., Kendrick, G. A., Althuizen, I. H. J., Balestri, E., Bernard, G., Cambridge, M. L., Cunha, A., Durance, C., Giesen, W., Han, Q., Hosokawa, S., Kiswara, W., Komatsu, T., Lardicci, C., Lee, K., Meinesz, A., Nakaoka, M., O'Brien, K. R., Paling, E. I., Pickerell, C., Ransijn, A. M. A., and Verduin, J. J.: Global analysis of seagrass restoration: the importance of large-scale planting, *Journal of Applied Ecology*, 53, 567–578, <https://doi.org/10.1111/1365-2664.12562>, 2016.
- Van Wijnen, H. and Bakker, J.: Long-term surface elevation change in salt marshes: a prediction of marsh response to future sea-level rise, *Estuarine, Coastal and Shelf Science*, 52, 381–390, 2001.
- Vanni, M. J.: Nutrient Cycling by Animals in Freshwater Ecosystems, *Annual Review of Ecology, Evolution, and Systematics*, 33, 341–370, <https://doi.org/10.1146/annurev.ecolsys.33.010802.150519>, 2002.
- Voulgaris, G. and Meyers, S. T.: Net effect of rainfall activity on salt-marsh sediment distribution, *Marine Geology*, 207, 115–129, <https://doi.org/10.1016/j.margeo.2004.03.009>, 2004.
- Vu, H. D., Wie, ski, K., and Pennings, S. C.: Ecosystem engineers drive creek formation in salt marshes, *Ecology*, 98, 162–174, <https://doi.org/10.1002/ecy.1628>, 2017.
- Wang, J. Q., Zhang, X. D., Jiang, L. F., Bertness, M. D., Fang, C. M., Chen, J. K., Hara, T., and Li, B.: Bioturbation of Burrowing Crabs Promotes Sediment Turnover and Carbon and Nitrogen Movements in an Estuarine Salt Marsh, *Ecosystems*, 13, 586–599, <https://doi.org/10.1007/s10021-010-9342-5>, 2010.
- Warren, J. H. and Underwood, A. J.: Effects of burrowing crabs on the topography of mangrove swamps in New South Wales, *Journal of Experimental Marine Biology and Ecology*, 102, 223–235, [https://doi.org/10.1016/0022-0981\(86\)90178-4](https://doi.org/10.1016/0022-0981(86)90178-4), 1986.
- Watling, L.: The sedimentary milieu and its consequences for resident organisms, *American Zoologist*, 31, 789–796, 1991.
- Wilson, C. A. and Allison, M. A.: An equilibrium profile model for retreating marsh shorelines in southeast Louisiana, *Estuarine, Coastal and Shelf Science*, 80, 483–494, <https://doi.org/10.1016/j.ecss.2008.09.004>, 2008.
- Wilson, C. A., Hughes, Z. J., and FitzGerald, D. M.: The effects of crab bioturbation on Mid-Atlantic saltmarsh tidal creek extension: Geotechnical and geochemical changes, *Estuarine, Coastal and Shelf Science*, 106, 33–44, <https://doi.org/10.1016/j.ecss.2012.04.019>, 2012.
- Wilson, C. A., Hughes, Z. J., and FitzGerald, D. M.: Causal relationships among sea level rise, marsh crab activity, and salt marsh geomorphology, *Proceedings of the National Academy of the United States of America*, 119, e2111535119, <https://doi.org/10.1073/pnas.2111535119>, 2022.
- Wittingham, S. S., Johnson, D. S., Chen, Y., and Kirwan, M. L.: A grazing crab drives saltmarsh carbon storage and recovery, *Ecology*, 105, e4385, <https://doi.org/10.1002/ecy.4385>, 2024.



- Worthington, T., Spalding, M., Landis, E., Maxwell, T. L., Navarro, A., Smart, L. S., and Murray, N. J.: The global distribution of tidal marshes from earth observation data, 2024.
- Xie, T., Dou, P., Li, S., Cui, B., Bai, J., Wang, Q., and Ning, Z.: Potential Effect of Bioturbation by Burrowing Crabs on Sediment Parameters in Coastal Salt Marshes, Wetlands, 40, 2775–2784, <https://doi.org/10.1007/s13157-020-01341-1>, 2020.
- Zhang, Z. and Colle, B. A.: Impact of Dynamically Downscaling Two CMIP5 Models on the Historical and Future Changes in Winter Extratropical Cyclones along the East Coast of North America, Journal of Climate, 31, 8499–8525, <https://doi.org/10.1175/JCLI-D-18-0178.1>, 2018.