

# 1 Accretion, retreat and transgression of coastal wetlands 2 experiencing sea-level rise

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11 **Abstract.** The vulnerability of coastal wetlands to future sea-level rise (SLR) has been extensively studied in  
12 recent years, and models of coastal wetland evolution have been developed to assess and quantify the expected  
13 impacts. Coastal wetlands respond to SLR by vertical accretion and landward migration. Wetlands accrete due to  
14 their capacity to trap sediments and to incorporate dead leaves, branches stems and roots into the soil, and they  
15 migrate driven by the preferred inundation conditions in terms of salinity and oxygen availability. Accretion and  
16 migration strongly interact and they both depend on water flow and sediment distribution within the wetland, so  
17 wetlands under the same external flow and sediment forcing but with different configurations will respond  
18 differently to SLR. Analyses of wetland response to SLR that do not incorporate realistic consideration of flow  
19 and sediment distribution, like the bathtub approach, are likely to result in poor estimates of wetland resilience.  
20 Here, we investigate how accretion and migration processes affect wetland response to SLR using a computational  
21 framework that includes all relevant hydrodynamic and sediment transport mechanisms that affect vegetation and  
22 landscape dynamics, and it is efficient enough computationally to allow the simulation of long time periods. Our  
23 framework incorporates two vegetation species, mangrove and saltmarsh, and accounts for the effects of natural  
24 and manmade features like inner channels, embankments and flow constrictions due to culverts. We apply our  
25 model to simplified domains that represent four different settings found in coastal wetlands, including a case of a  
26 tidal flat free from obstructions or drainage features and three other cases incorporating an inner channel, an  
27 embankment with a culvert, and a combination of inner channel, embankment and culvert. We use conditions  
28 typical of SE Australia in terms of vegetation, tidal range and sediment load, but we also analyse situations with  
29 three times the sediment load to assess the potential of biophysical feedbacks to produce increased accretion rates.  
30 We find that all wetland settings are unable to cope with SLR and disappear by the end of the century, even for  
31 the case of increased sediment load. Wetlands with good drainage that improves tidal flushing are more resilient  
32 than wetlands with obstacles that result in tidal attenuation, and can delay wetland submergence by 20 years.  
33 Results from a bathtub model reveals systematic overprediction of wetland resilience to SLR: by the end of the  
34 century, half of the wetland survives with a typical sediment load, while the entire wetland survives with increased  
35 sediment load.

36  
37 *Keywords:* coastal wetlands, sea-level rise, accretion, migration, hydrodynamic model, sediment transport model,  
38 mangrove, saltmarsh.

## 39 **1 Introduction**

40 The vulnerability of coastal wetlands to future sea-level rise has been extensively studied in recent years, and  
41 models of coastal wetland evolution have been developed to assess and quantify the expected impacts (Alizad et  
42 al., 2016b; Belliard et al., 2016; Clough et al., 2016; D'Alpaos et al., 2011; Fagherazzi et al., 2012; Kirwan and  
43 Megonigal, 2013; Krauss et al., 2010; Lovelock et al., 2015b; Mogensen and Rogers, 2018; Rodriguez et al.,  
44 2017; Rogers et al., 2012; Schuerch et al., 2018). Predictions vary widely, which is not surprising given the  
45 complexity of the processes involved and the practical challenges associated with representing interactions at a  
46 variety of spatial and temporal scales. Coastal wetlands respond to SLR by vertical accretion and landward  
47 migration. Vertical accretion occurs due to the capacity of wetland vegetation to trap sediments and to incorporate  
48 dead leaves, branches stems and roots into the soil, building up their vertical elevation and counteracting  
49 submergence due to SLR. Landward migration is driven by the preferred inundation conditions of wetland  
50 vegetation, which is continuously moving up the wetland slope due to SLR. These two main processes interact,  
51 but they also integrate a number of biophysical exchanges that occur smaller scales. Accretion is a function of  
52 many other variables like the tidal regime, sediment availability and type of vegetation (Fagherazzi et al.,  
53 2012; Lovelock et al., 2015a). Vegetation preference is dictated by salinity, oxygen availability and the presence  
54 of phytotoxins in the soil (Bilskie et al., 2016; Crase et al., 2013).

55 Studies show that different modelling approaches used to address the interaction between these variables may lead  
56 to divergent results (Alizad et al., 2016a; Rogers et al., 2012). For the sake of simplicity, some previous studies  
57 have adopted an approach where water levels throughout the wetland remain the same as those observed at the  
58 inlet, i.e. the bathtub approach (D'Alpaos et al., 2011; Kirwan and Guntenspergen, 2010; Kirwan et al.,  
59 2010; Kirwan et al., 2016a; Lovelock et al., 2015b). Most of these bathtub model results show that vegetation in  
60 coastal areas can produce accretion rates similar to sea-level rise predictions, therefore maintaining their elevation  
61 in the tidal prism, except when tidal range and sediment supply are very low. However, the projections of coastal  
62 wetland resilience under high rates of SLR appears to be at odds with paleo-environmental reconstructions of  
63 wetland responses to rising seas during the early Holocene (Horton et al., 2018; Saintilan et al., 2020). One  
64 explanation of this discrepancy is that models fail to reproduce the flow attenuation caused by the friction induced  
65 by substrate cover and specific wetland features like inner channels, embankments and flow constrictions (Hunt  
66 et al., 2015) and its effects on sediment availability, which may result in overestimation of wetland accretion rates  
67 (Rodriguez et al., 2017). Bathtub models do not provide information on flow discharges or velocities, so they need  
68 an independent specification of sediment concentration.

69 On the other hand, more detailed description of hydrodynamic and sediment transport mechanisms can be  
70 incorporated into the computations of wetland dynamics using conventional two or three dimensional flow and  
71 sediment transport models (Ganju et al., 2015; Lalimi et al., 2020; Temmerman et al., 2005). A detailed description  
72 of flow and sediment transport processes can potentially result in a better estimation of wetland dynamics  
73 including accretion and migration processes, but implementation can be seriously limited by computational cost  
74 and data availability (Beudin et al., 2017).

75 Here, we investigate how accretion and migration processes affect wetland response to SLR using a computational  
76 framework that integrates detailed hydrodynamic and sediment transport mechanisms that affect vegetation and  
77 landscape dynamics and that is efficient enough to allow the simulation of long time periods. The framework  
78 consists of a fast-performance quasi-2D hydrodynamic model (Riccardi, 2000; Rodriguez et al., 2017) that we

79 have extensively tested in wetlands (Rodriguez et al., 2017;Saco et al., 2019;Sandi et al., 2018;Sandi et al.,  
80 2019;Sandi et al., 2020a;Sandi et al., 2020b) and a sediment advection transport model (Garcia et al., 2015) that  
81 we couple with vegetation formulations for preference to tidal conditions to obtain realistic predictions of wetland  
82 accretion and migration under SLR. Our framework incorporates two vegetation species, mangrove and saltmarsh,  
83 and accounts for the effects of manmade features like inner channels, embankments and flow constrictions due to  
84 culverts. We apply our model to simplified domains that represent distinct areas within a real wetland, in which  
85 we are able to characterise the effects of particular natural and manmade wetland features like vegetation types,  
86 culverts, embankments and channels.

87 Coastal wetlands are found over a broad spectrum of geomorphological settings (Woodroffe et al., 2016) and  
88 under a diverse set of anthropogenic interventions (Temmerman and Kirwan, 2015). While our results strictly  
89 apply to areas in a particular wetland in Southeast Australia, each of our selected domains focusses on specific  
90 geomorphological characteristics that may also be present in other wetlands worldwide. We study wetland  
91 evolution on domains with no drainage network or manmade structures, which is relevant for some low-tide  
92 wetland environments where no human intervention has occurred (Leong et al., 2018;Oliver et al., 2012;Tabak et  
93 al., 2016). We simulate the dynamics of internal channels, which can provide insight on wetlands studies with  
94 strong influence of natural channels (Reef et al., 2018;Silvestri et al., 2005) or manmade drainage channels  
95 (Manda et al., 2014). We carry-out simulations with embankments and culverts representing flood sheltered  
96 environments, which can resemble intentional flood attenuation works for coastal protection (Van Loon-Steensma  
97 et al., 2015) or unintentional flood attenuation as the result of roads, tracks, pipes and other infrastructure typical  
98 of heavily human-occupied coasts (Kirwan and Megonigal, 2013;Rodriguez et al., 2017;Temmerman et al., 2003).

99 Also, and in order to make our results more widely relevant, we analyse the sensitivity of our predictions  
100 to the sediment load coming into the wetland by including sediment-poor and sediment-rich simulations. The  
101 incoming sediment load has been proposed as one of the main factor influencing the resilience of coastal wetlands  
102 to SLR (Lovelock et al., 2015a;Schuerch et al., 2018) and is one of the components of predictive wetland evolution  
103 models with more uncertainty, due both to our limited understanding of sediment-flow-vegetation processes and  
104 our inability to predict sediment loads in a changing future.

## 105 **2 Experimental design and methods**

### 106 **2.1 Design of simulations**

107 The flow in tidal wetlands can be quite complex because of the interaction of the tidal flow with natural and  
108 manmade features like vegetation, topography, channels, culverts and embankments. For that reason, results for  
109 a particular wetland may have limited applicability to another wetland with different features. In this contribution,  
110 we analyse some of the most common features of wetlands in isolation in order to gain a better understanding of  
111 the contribution of each feature to the overall wetland response, and how it influences the response to sea-level  
112 rise. For that purpose, we study the response of wetlands with limited complexity using a state of the art  
113 ecogeomorphological model on four hypothetical tidal flats that characterise specific areas of a typical SE  
114 Australian coastal wetland that we have studied before (Fig.1 a,b) (Rodriguez et al., 2017). Simulation 1 uses a  
115 bathtub approach over a consistently sloping tidal flat initially vegetated by mangrove, saltmarsh and freshwater  
116 vegetation (Fig. 1c), in which water levels are considered uniform over the domain and no special features are

117 taken into account. In contrast, for Simulations 2 to 5, water levels are calculated with the hydrodynamic model,  
118 which allows for the inclusion of attenuation effects from vegetation and special features. Simulation 2 considers  
119 a vegetated sloping tidal flat with no special features, , Simulation 3 incorporates a drainage channel 0.4 m deep  
120 and 5 m wide to the vegetated tidal flat, Simulation 4 includes an embankment with a culvert (0.8 m wide and 0.5  
121 m tall) in the middle of the vegetated flat, and Simulation 5 combines both a drainage channel and an embankment  
122 with a culvert (Fig. 1d). These different setups can characterise different settings found in wetlands, but can also  
123 apply to different parts of a more complex wetland, as shown in Fig. 1b. In all simulations the tidal flat is 620 m  
124 long (main flow direction) and 310 m wide (cross-section), divided into 10m by 10m grid cells, with a gentle  
125 slope of 0.001 m/m. Boundary conditions include input tides described by a sinusoidal function with 1.3 m  
126 amplitude and 12-hour period, and a constant sediment concentration at the wetland inlet (Fig. 1c). In each  
127 simulation we tested wetland evolution under sea-level rise from 2000 to 2100 (high emissions scenario)  
128 considering two sediment input conditions, a low sediment supply representing current conditions and a high  
129 sediment supply. The high sediment supply condition simulations are justified due to the uncertainty of climatic  
130 conditions and the possibility of increases in intensity of storm patterns in the area, which may result on increased  
131 sediment loads in the Hunter River. Sediment loads may also increase due to changes in land use practices  
132 (Rodriguez et al., 2020).

133 The sinusoidal tide represents conditions typical of SE Australian estuaries (Rodriguez et al., 2017) and is repeated  
134 during the simulation period (100 years). However, the mean water level is gradually increased following the  
135 IPCC RCP 8.5 scenario of sea-level rise (Church et al., 2013) with an expected 0.74 m increase by year 2100 with  
136 respect to the levels in the year 2000.

137 We use as a basis for our simulations the ecogeomorphological model (EGM) framework developed by Rodriguez  
138 et al. (2017), but with the addition of a physically-based sediment transport formulation. This EGM framework  
139 has been extensively calibrated and tested in the Hunter River Estuary in Australia and, as such, vegetation  
140 functions and parameters correspond to local conditions. The framework couples multiple models to simulate  
141 interactions between overland flow hydrodynamics, vegetation establishment and growth, sediment concentration  
142 and morphodynamics of the wetland.

## 143 **2.2 Hydrodynamic model**

144 Water depth time series over the tidal flat are estimated using a finite-differences quasi-2D hydrodynamic model  
145 (Riccardi, 2000) that has been successfully applied to coastal wetlands (Rodriguez et al., 2017; Sandi et al., 2018)  
146 and floodplains (Sandi et al., 2019; Sandi et al., 2020a; Sandi et al., 2020b; Saco et al., 2019). The model solves the  
147 shallow water equations using a cells scheme, in which cells are classified into tidal flat or channel categories to  
148 speed up computations. As previously explained, the domains of all simulations are 630 m long by 310 m wide,  
149 discretised into 10mx10m cells. For cells representing channels in simulations 3 and 5, the width of the cell is  
150 reduced to 5 m and the elevation is lowered by 0.4 m. Boundary conditions include water elevations at the tidal  
151 creek and no-flow at the lateral and landward boundaries. Because the domains are wide, the effects of lateral  
152 model boundaries are minimal.

153 In each timestep, the model solves for water elevations at every cell using mass conservation in a 2D formulation,  
154 and then it solves for discharges between cells in each direction using momentum conservation in a 1D  
155 formulation. Mass conservation is solved first to compute water surface elevations:

156  $As_i \frac{dz_i}{dt} = \sum_{k=1}^j Q_{k,i},$   
 157 (1)

158 where  $As_i$  and  $z_i$  are surface wetted area and water surface elevation at cell  $i$ , respectively and  $Q_{k,i}$  are the  
 159 discharges between cell  $i$  and its  $j$  neighbouring cells. Using the water surface elevations, the model then computes  
 160 discharges between cells using the momentum or energy equation, depending on the particular characteristics of  
 161 the connection between cells. For instance, the discharge between two cells on the vegetated tidal flat is computed  
 162 as:

163  $Q_{k,i} = \frac{A_{k,i} R_{k,i}^{\frac{2}{3}}}{n_{k,i}} \left( \frac{z_k - z_i}{x_k - x_i} \right)^{\frac{1}{2}},$   
 164 (2)

165 where  $A_{k,i}$ ,  $R_{k,i}$  and  $n_{k,i}$  are respectively the cross-sectional values of area, wetted perimeter and Manning  
 166 roughness computed as an average of the values at cells  $k$  and  $i$ , and  $x_k - x_i$  is the distance between cells. Based  
 167 on Rodriguez et al. (2017) we adopt roughness coefficients for mangrove and saltmarsh cells of  $0.50 \text{ s/m}^{1/3}$  and  
 168  $0.15 \text{ s/m}^{1/3}$ , respectively. For freshwater and no-vegetated cells, the Manning's- $n$  is  $0.12 \text{ s/m}^{1/3}$ , while for channel  
 169 cell it is  $0.035 \text{ s/m}^{1/3}$ . For cells in the channel, the full momentum equation is used to account for dynamic and  
 170 backwater effects (Riccardi, 2000). If the domain includes a culvert at cell  $i$ , then the discharge between cells  $k$   
 171 and  $i$  is computed as:

172  $Q_{k,i} = \frac{(2g)^{\frac{1}{2}} (z_k - z_i)^{\frac{1}{2}}}{\left( \frac{1}{C_d^2 A_i^2} - \frac{1}{A_k^2} \right)^{\frac{1}{2}}},$   
 173 (3)

174 in which  $A_i$  and  $A_k$  are respectively the cross-sectional areas at the  $i$  and  $k$  cells and  $C_d$  is a standard discharge  
 175 coefficient for the culvert at cell  $i$  adopted as 0.8. Equation (3) considered the case of the culvert flowing under  
 176 the influence of gravity. For pressurised conditions, a different equation is used (Riccardi, 2000)

177 The model equations are solved using an implicit method and a Newton-Raphson algorithm. The time step used  
 178 in the model solution is 1s to ensure numerical stability. Further explanation about the application of this model  
 179 in a similar EGM framework can be found in Sandi et al. (2018).

### 180 2.3 Vegetation model

181 Vegetation in coastal wetlands is driven by the tidal regime, so we use water depth time-series to compute the  
 182 mean depth below high tide,  $D$ , and the hydroperiod,  $H$ , on every cell as a descriptor of the tidal regime. These  
 183 variables are the input for all the other models of the EGM framework. The first variable represents the average  
 184 maximum water depth on spring tides. In this case we use a sinusoidal wave, so  $D$  is the maximum depth. The  
 185 hydroperiod accounts for the duration of the inundation period and is computed as the proportion of time during  
 186 which a minimum water depth is present during the simulation time.

187 The values of  $H$  and  $D$  define the suitable conditions for vegetation establishment and survival at each point in  
 188 the wetland based on thresholds that have been tested for SE Australian estuaries (Rodriguez et al., 2017). Thus,  
 189 the observed threshold applies to *Avicennia marina* (grey mangrove) and to a composition of saltmarsh species

190 *Sarcocornia quinqueflora* and *Sporobolus virginicus*. Mangrove depends primarily on hydroperiod, requires  
 191 frequent inundations and establishes in areas where  $10\% < H < 50\%$  and  $D > 0.2$  m, where  $H$  is calculated as the  
 192 fraction of time where the water depth is higher than or equal to 14 cm, the typical height of the pneumatophores.  
 193 Saltmarsh tolerates prolonged inundations and can survive in areas where  $H < 80\%$ , but cannot endure inundation  
 194 depths above its height (25 cm) so we limit  $D < 0.25$  m. We consider that, if conditions suit both mangrove and  
 195 saltmarsh, mangrove will expand over saltmarsh areas (Saintilan et al., 2014). In areas not exposed to saltwater  
 196 ( $H = 0\%$ ,  $D \sim 0$  m), we assume the presence of freshwater vegetation, and if none of the above conditions applies,  
 197 areas are considered to be non-vegetated.

## 198 2.4 Sediment model

199 The original version of the framework used in the Hunter estuary applies a linear empirical relationship between  
 200 average sediment concentration in the water column and the water depth. Here, we use a more physically based  
 201 equation for fine sediment transport and deposition processes coupled to the hydrodynamic simulations. The  
 202 sediment model solves the quasi-2D continuity equation of suspended sediment neglecting horizontal diffusion  
 203 (Garcia et al., 2015). The continuity equation for the  $i$ -th cell reads as follows:

$$204 \quad A S_i \frac{d(hC)_i}{dt} = A S_i \varphi_i + \sum_{k=1}^j (QC)_{k,i},$$

205 (4)

206 where  $h_i$  is the water depth of cell  $i$  (m);  $C_i$  is the sediment concentration ( $\text{g m}^{-3}$ ),  $\varphi_i$  is the downward vertical  
 207 flux of fine sediment ( $\text{g m}^{-2} \text{ s}^{-1}$ ), and  $C_{k,i}$  are the sediment concentrations in the  $j$  neighbouring cells. For fine  
 208 grained sediment typical of estuarine environments, the downward flux can be expressed as (Krone, 1962; Mehta  
 209 and McAnally, 2008):

$$210 \quad \varphi_i = -w_s \left(1 - \frac{\tau_{bi}}{\tau_d}\right) C_i; \quad \tau_b < \tau_d,$$

211 (5)

212 where  $w_s$  is the fall/settling velocity of suspended sediment particles ( $\text{m s}^{-1}$ ),  $\tau_{bi}$  is the magnitude of bed shear  
 213 stress in cell  $i$  (Pa), and  $\tau_d$  is the critical bed shear stress for deposition (Pa). Velocities were converted to bed  
 214 shear stresses using

$$215 \quad \tau_{bi} = \rho C_f U_i^2,$$

216 (6)

217 In equation (6)  $\rho$  is the water density and  $C_f$  is a friction coefficient set at 0.05. The parameters  $w_s$  and  $\tau_d$  were  
 218 varied to reproduce similar levels of accretion observed in the wetlands where the original modelling framework  
 219 was applied (Rodriguez et al., 2017). The values obtained were  $\tau_d = 0.02$  Pa and  $w_s = 2 \times 10^{-4}$  m/s, which are  
 220 consistent with values reported by Larsen et al. (2009) and Temmerman et al. (2005). This model does not have  
 221 an erosion term, which is not a bad simplification over vegetated surfaces that receive flows that are typically very  
 222 slow.

223 Equation (4) is solved using the same numerical scheme than the water mass conservation (equation 1) providing  
 224 a time series of sediment concentrations in each cell of the domain. However, as the soil elevation model (next  
 225 section) works at a larger time scale and requires the annual concentration,  $\bar{C}$ , a weighted average is computed for  
 226 each cell:

$$\bar{C} = \frac{\sum_{t=0}^M (C_t \times h_t)}{\sum_{t=0}^M h_t}, \quad (7)$$

where  $t$  is the time in the hydrodynamic simulation with  $M$  the final step,  $C_t$  and  $h_t$  are the sediment concentration and the water depth, respectively, at time  $t$ .

The sediment transport equation based on mass conservation (eq. 4) cannot be used in the case of the bathtub simulations because the bathtub model does not provide information on water discharge and velocity. For the bathtub simulations, we used the linear relation between water depth and concentration empirically developed by Rodriguez et al. (2017). Based on the measured data, the fitted equation is:

$$\bar{C} = C_{max}(0.55D + 0.32), \quad (8)$$

where  $\bar{C}$  is the average sediment concentration ( $\text{g}/\text{m}^3$ ), and  $C_{max}$  is the concentration at the wetland inlet.

This equation is much simpler and has different parameters than the sediment transport equation; however, for very simple flow conditions it should produce comparable results. We confirmed the suitability of the simple model by comparing EGM results using the bathtub approach (with the linear sediment relation) and a full hydrodynamic and sediment transport EGM over a smooth topography. Both the hydrodynamics and the resulting elevation changes of both models were very similar (See Fig. S1 in Supplementary Materials).

## 2.5 Soil elevation change model

Our EGM framework adopts the model originally proposed by Morris et al. (2002) and later modified by Kirwan and Guntenspergen (2010) to estimate the increase in soil elevation due to accretion as function of hydrodynamic and ecological conditions. We first compute the biomass production,  $B$  ( $\text{g}/\text{m}^2$ ), by using the parabolic equation:

$$B = aD^2 + bD + c, \quad (9)$$

where  $a$ ,  $b$ , and  $c$  are parameters fitted to field data, for each vegetation type. Then, the surface elevation change rate,  $dE/dt$  ( $\text{m}/\text{year}$ ), is calculated using:

$$\frac{dE}{dt} = \bar{C}(q + kB)D, \quad (10)$$

where  $q$  is a depositional parameter and  $k$  is a vegetation sediment trapping coefficient. For all five parameters of equations (9) and (10) we used the values adopted in Rodriguez et al. (2017) and Sandi et al. (2018) (see Table 1) for an Australian wetland. Although the term  $As_i\phi_i$  in equation (4) provides an amount of settled sediment that contributes to accretion, it only considers the gravitational settling of sediment and does not include many other important accretion processes associated to the presence of vegetation. The full effects of sediment and vegetation are considered in equation (10), which produces much larger accretion values (see Fig. S2 in Supplementary Materials).

The EGM simulations use a yearly time-step, i.e. the computed biomass and accretion represent an average condition within this period. We choose a yearly time-step as vegetation dynamics does not respond instantaneously to flow and depositional processes (Alizad et al., 2016b; Saco and Rodríguez, 2013; Schuerch et al., 2018). Our model does not account for erosion and diffusion processes and also does not take into account the redistribution of deposited sediment by waves. Because of that, the resulting accretion from equation (10) is noisy

265 and vary considerably over very short distances. In order to work with a more realistic distribution of deposition  
266 over the tidal flat we smooth the topography by applying a very simple diffusion model. The diffusion model does  
267 not change the general trends of deposition and avoids localised peaks of excessive deposition.

## 268 **3 Results**

### 269 **3.1 Spatial patterns of accretion and vegetation**

270 In order to show the characteristic spatial patterns of each of the typical cases analysed we first show in Fig. 2  
271 accumulated accretion ( $\Delta E$ ) and vegetation distribution in 2050 under the expected SLR scenario for each of the  
272 five numerical simulations, including the bathtub and the other four simulations that use a hydrodynamic and  
273 sediment transport (HST) models. Details on the temporal evolution of topography and vegetation for each of the  
274 simulations are provided later in the manuscript.

275 Fig. 2 shows that accumulated accretion is homogeneous in the transverse direction for the simulations without  
276 the channel (Fig. 2a,b,d), as there is no lateral flow and the changes in sedimentation occur in the longitudinal  
277 direction only. For the simulations with the central drainage channel (Fig. 2 c,e) there is a marked concentration  
278 of flow and sediment accumulation close to the channel. Some of the accumulated accretion patterns of the  
279 simulations with the channel presented in Fig. 2 are remarkably similar to the results from Chen et al. (2010) on  
280 a similar geometry.

281 It can be seen from the figure that all simulations show a general decrease of accretion with distance to the tidal  
282 input (which can represent a tidal creek or the river), which is expected because the source of sediment is at the  
283 tidal input. However, each simulation has a characteristic elevation profile and vegetation distribution, and they  
284 are all quite different from the predictions of the bathtub model. Fig. 2a shows that the bathtub simulation displays  
285 a smoother and longer transition of accumulated accretion. A slight concentration of accretion is observed at 500  
286 m from the creek, due to the initial position of high biomass saltmarsh. The bathtub case has flood and ebb flows  
287 of the same duration, since there is no flow attenuation. This keeps the hydroperiod within a range that promotes  
288 mangrove establishment over most of the wetland. Saltmarsh is limited to the upper parts of the tidal flat.

289 The other simulations (2 to 5) use the hydrodynamic and sediment transport (HST) models instead of the bathtub  
290 approximation. In these cases, accretion presents an exponential shape with a sharper decrease than the bathtub  
291 model, and vegetation establishment is strongly controlled by the effects of vegetation roughness, channel and  
292 culverts. In contrast to the bathtub model results, all HST simulations show mangrove dieback in lower areas,  
293 which is caused by a higher hydroperiod due to attenuated ebb flows.

294 Simulation 2, with the undisturbed tidal flat (Fig. 2b), shows the effect of hydraulic resistance due to the vegetation  
295 roughness only, which generates an elevation mound closer to the tidal input than the bathtub simulation. In  
296 Simulation 3 (Fig. 2c), the inner channel increases the drainage of the surrounding areas, thus reducing the  
297 hydroperiod in the vicinity of the channel and allowing mangroves to persist close to the tidal creek. The channel  
298 also enhances sediment delivery farther from the tidal input, which causes an increase in accretion around the  
299 mid-point of the flat (300 m from the tidal creek). However, this effect is concentrated near the channel and fades  
300 away as flow is directed into the tidal flat. In Simulation 4 (Fig. 2d), the flow is restricted by an embankment and  
301 a culvert, so the hydroperiods in the upper wetland are higher. This effect reduces mangrove migration and its  
302 encroachment on saltmarsh areas. In Simulation 5 with embankment and channel (Fig. 2e), the channel promotes



303 mangrove landwards of the embankment, and also the stabilisation of saltmarsh areas in the upper sections of the  
304 tidal flat as they receive more sediment (Fig. 2e).

### 305 **3.2 Evolution of accumulated accretion profiles**

306 Fig. 3 shows the results of surface elevation change ( $\Delta E$ ) in each simulation for the years 2020, 2040, 2060 and  
307 2100 for low sediment input conditions (corresponding to contemporary rates in the Hunter estuary), in terms of  
308 accumulated accretion profiles along the main flow direction. For the simulations with the central drainage  
309 channel (Simulations 3 and 5), we have included two profiles at different transverse locations, one close to the  
310 channel and one 150 m away in the middle of the tidal flat.

311 During the first two decades, the vegetation type plays an important role in the longitudinal distribution of the  
312 accumulated accretion profiles. By 2020 (first column of Fig. 3) the profiles show a continuous decrease from  
313 the tidal input up to 300 to 350 m approximately, which coincides with the transition from mangrove to saltmarsh  
314 in the initial vegetation distributions (see Fig. 5 later in the manuscript). This occurs due to the dynamics of  
315 sediment transport (more deposition close to the tidal input) and also due to the reduction of the mangrove biomass  
316 away from the tidal creek (reductions in  $D$ , see eqn. 10). The increase in  $\Delta E$  at the transition is due to the saltmarsh  
317 having a higher biomass and trapping efficiency than mangrove at that particular value of  $D$ . Landward of the  
318 transition,  $\Delta E$  decreases with decreases in saltmarsh biomass. This general dynamics is disrupted by the presence  
319 of the culvert because it limits the amount of sediment reaching the upper areas of the tidal flat.

320 Changes in  $\Delta E$  slow down after 2060 in all simulations except for the bathtub case. This is due to reductions in  
321 vegetation as most of the lower areas of the tidal flat have experienced submergence and vegetation loss. Small  
322 increases in  $\Delta E$  occur in the upper areas in the cases in which the central channel promotes tidal flushing  
323 (Simulations 3 and 5), but this effect is concentrated in areas close to the channel.

324 None of the simulations using the HST model produces  $\Delta E$  results similar to the bathtub simulations. The  
325 simulation with the central channel (Simulation 3), presents values of  $\Delta E$  near the channel that are close to the  
326 results of the bathtub simulation during the first years, but over time, the results diverge. The increased  $\Delta E$  values  
327 are limited to areas next to the channel, and they quickly decline as the flow is directed into the tidal flat. In  
328 general, the outcomes from the HST model shows a reduction in the water levels and total accretion compared to  
329 the bathtub results. Furthermore, when the culvert is introduced in the simulation (Experiments 4 and 5), the main  
330 effect is a drastic reduction of  $\Delta E$  in the upper areas of the domain.

331 Fig. 3 results correspond to a situation with a low sediment input of  $37 \text{ g/m}^3$ , typical of current SE Australia  
332 conditions (Rodriguez et al., 2017). Similar patterns but with larger values of accumulated accretion were obtained  
333 for a higher sediment input of  $111 \text{ g/m}^3$  (Fig. S3 in Supplementary Materials).

334 The reduction in accretion in the simulations that consider the actual features of the wetland can be better  
335 appreciated in Fig. 4, in which we compare domain-average  $\Delta E$  of all simulations over time. Fig. 4 includes results  
336 for a low sediment input of  $37 \text{ g/m}^3$  (Fig. 4a) and for a high sediment input of  $111 \text{ g/m}^3$  (Fig. 4b). The figure also  
337 includes the values of mean sea-level for each year to give an idea of the submergence conditions in the wetlands.  
338 There is a clear difference between the accretion generated in the bathtub simulation, and the rest of the  
339 simulations. In our simulations, accretion is a function of sediment concentration and depth below mean high tide  
340 ( $D$ ). The bathtub assumption overpredicts both inputs over the entire domain, thus generating higher accretion  
341 values. In all HST simulations, the combination of a reduction in  $D$  because of flow attenuation and the

342 exponential decay of sediment concentration results in less accretion than in the bathtub simulation. In the case of  
343 low sediment input (Fig. 4a), by 2050 the domain-average  $\Delta E$  from the bathtub is about 2 times the values of all  
344 the other simulations, increasing to more than 3 times by 2100. In the simulations with high sediment input (Fig.  
345 4b), the accumulated accretion of bathtub simulations are 2.5 and 4 times the values of the rest of the simulations  
346 for 2050 and 2100, respectively. The simulations with the HST simulations present different levels of attenuation  
347 and accordingly different accretion levels. The lowest accretion correspond to the highly attenuated case with  
348 embankment and culvert (Sim. 4), whereas the highest accretion occur in the case of the central channel (Sim. 3)  
349 that experiences increased drainage and thus less attenuation. The cases of the tidal flat with no structures (Sim.  
350 2) and of the embankment with inner channel (Sim.5) have intermediate levels of attenuation and accretion.  
351 All simulations show a strong elevation deficit (i.e. the difference between the rate of sea level rise and wetland  
352 accretion rate  $dE/dt$ ), as none of the simulations predict that the tidal flat is capable to keep pace with SLR. For  
353 the low-sediment conditions, by 2050 the elevation deficit of the bathtub simulation is 5.5 mm/yr, while the rest  
354 of the simulations predict an elevation deficit of about 7 mm/yr. Over time, the elevation deficits increase and by  
355 2100 the bathtub prediction reach a value of 9.5 mm/yr and the HST simulations a value of 12 mm/yr.  
356 Increasing the sediment input concentration considerably changes the accretion capacity of the tidal flat,  
357 particularly according to the bathtub results. Bathtub simulations predict that the tidal flat is able to accrete in a  
358 rate that almost match the changes in sea level, so the wetland survives sea-level rise. Accretion for all other  
359 simulations are moderate, with the simulations that have the central channel (Experiments 3 and 5) responding  
360 more effectively to the increased sediment and accreting more than the other simulations (Sim. 2 and 4). Compared  
361 to the low sediment conditions, elevation deficits of the bathtub predictions reduce to 3 mm/yr and 5.5 mm/yr by  
362 2050 and 2100, respectively, while in the other simulations those values increase to about 6 mm/yr and 10 mm/yr.  
363 The structures included in the simulations have a clear effect on the average  $\Delta E$ . The inner channel promotes  
364 accretion further inland, as it conveys more water and sediment to those areas away from the tidal input. Compared  
365 to the tidal flat free of structures (Simulation 2) the inclusion of the channel (Simulation 3) is responsible for an  
366 increase in wetland accumulated accretion of about 50%. The opposite effect is observed when the embankment  
367 with culvert is introduced, as it attenuates and reduces the water and sediment flow into the upper part of the  
368 wetland. Comparing results for the tidal flat without (Simulation 2) and with (Simulation 4) embankment and  
369 culvert, we can observe a reduction on wetland accumulated accretion of 25%. The introduction of a drainage  
370 channel together with the embankment and culvert (Simulation 5) represents an intermediate situation in which  
371 the increased flushing effect of the channel and the attenuating effect of the embankment and culvert partially  
372 compensate.

373 In Fig. 4a we have also included the average accumulated accretion for the entire wetland site (Area E in Fig. 1b)  
374 using information from Rodriguez et al. (2017) and (Sandi et al., 2018). Rodriguez et al. (2017) applied a similar  
375 EGM formulation to Area E (Fig. 1c) to assess the effect of attenuation on wetland evolution under SLR  
376 considering typical ( $37 \text{ g/m}^3$ ) and increased ( $111 \text{ g/m}^3$ ) sediment conditions. (Sandi et al., 2018) further studied  
377 the effects of tidal restrictions at the wetland inlet considering typical sediment loads. The values included in the  
378 figure correspond to average accumulated accretion over the entire wetland at 2050 and 2100 for low sediment  
379 load with and without tidal restrictions (Fig. 4a) and for high sediment load without restrictions (Fig. 4b). The  
380 figures shows that the simulations without tidal restrictions result in values of accumulated accretion similar to  
381 the simulation with low attenuation (Simulations 3 and 5) for both low and high sediment loads, while predictions

382 of accumulated accretion including tidal restrictions are closer to the simulation with high attenuation (Simulations  
383 2 and 4).

### 384 **3.3 Changes in vegetation**

385 The interactions between sea-level rise, accretion and vegetation changes are complex because vegetation not  
386 only responds to vertical elevation changes but also migrates inland. In order to obtain a clear picture of the  
387 vegetation changes over time, we simplified two dimensional vegetation maps (i.e., Fig. 2) into a one-dimensional  
388 representation. The vegetation type at a given distance from the tidal input was determined by selecting the  
389 predominant (higher occurrence) vegetation in the transverse direction. Fig. 5 shows snapshots of the predominant  
390 vegetation every 20 years. As already explained, in the simulations with embankment and culvert (Simulations 4  
391 and 5), the structures are located at 310 m from the tidal input. The conditions at the beginning of the simulation  
392 (Fig. 5a) for simulations 1, 2, and 3 show mangrove occupying approximately the lower 400 m of the tidal flat  
393 and saltmarsh the next 200 m upland. For simulations 4 and 5 the presence of the embankment reduces  
394 hydroperiods in the upper areas, constraining mangrove to the lower 310 m. The embankment also limits the  
395 extent of inundation in the upper areas, reducing the extent of the saltmarsh to about 100 m from the embankment.  
396 After 20 years (Fig. 5b) the simulations 1, 2 and 3 show mangrove encroachment on saltmarsh. The upstream  
397 mangrove edge moves up to 50 m, forcing saltmarsh occurrence in areas further than 300 m from the tide input  
398 creek. In simulations 4 and 5 the embankment halts mangrove migration and increases in inundation of upper  
399 areas promote saltmarsh increase. Overall, wetland area increases due to mangrove expansion (Sim. 1, 2 and 3)  
400 or to saltmarsh expansion (Sim. 4 and 5).

401 By 2040 (Fig. 5c), mangrove has encroached further on saltmarsh in simulations 1, 2 and 3, resulting in saltmarsh  
402 squeeze at the upper end due to the landward boundary of the computational domain. Simulations 4 and 5 show  
403 very minor encroachment of mangrove on saltmarsh, which is able to migrate landward. Total wetland area  
404 remains approximately unchanged for simulations 1, 2 and 3, while it keeps increasing in simulations 4 and 5.  
405 Some areas of mudflat start appearing in the HST simulations due to extended hydroperiods.

406 Twenty years later, in 2060 (Fig. 5d), the MSL is about 30 cm higher than in 2000 and we can see considerable  
407 mudflat areas in all simulations except for the bathtub simulation (Simulation 1), which presents a uniform  
408 coverage of mangrove over the entire domain. Saltmarsh is totally absent in simulations 1, 2 and 3 due to mangrove  
409 encroachment but still remains almost unchanged in simulations 4 and 5. All simulations except the bathtub  
410 simulation show decreases in wetland extent, mostly due to saltmarsh disappearance in simulations 2 and 3 and  
411 to mangrove squeeze in simulations 4 and 5.

412 From 2080 on (Fig. 5e,f), a rapid retreat of the remaining wetland can be observed in all simulations. The retreat  
413 occurs faster for the simulations with the embankment, resulting in total wetland disappearance by 2100. The rest  
414 of the simulations still show some remnant mangrove areas by 2100, which are only significant (40%) in the case  
415 of the bathtub simulations.

416 The same trend of increase in wetland area in the first 20 years of simulation, followed by a continuous decrease  
417 starting at 40 years and ending at 100 years with almost complete wetland disappearance under the same sea level  
418 rise trajectory was observed by Rodriguez et al. (2017) and Sandi et al. (2018). Sandi et al. (2018) also reported  
419 larger wetland losses in their simulations with tidal input restrictions at the wetland inlet when compared to the  
420 case without restrictions.

421 The same analysis of vegetation evolution for the high sediment input scenario is presented in Fig. 6. With  
422 increased sediment, the patterns of vegetation change remain remarkably similar to the patterns observed in Fig.  
423 5 for the low sediment conditions, with exception of the bathtub simulations (Simulation 1). Compared to Fig. 5,  
424 the bathtub results indicate that saltmarsh is able to remain in the upper wetland areas for longer (until 2060) and  
425 that mangrove does not retreat, resulting in no wetland loss after 100 years of simulation. The other simulations  
426 without embankment (2 and 3) show a slightly slower retreat of both mangrove and saltmarsh than in Fig. 5, while  
427 the simulations with the embankment show almost the same behaviour as the in the case of low sediment. Some  
428 of simulations in Fig. 6 show localised mangrove areas that tend to establish and persist close to the tidal creek.  
429 For a more detailed analysis we can look at the vegetation evolution in terms of wetland area (mangrove and  
430 saltmarsh), wetland retreat (position of the seaward edge) and wetland transgression (position of the landward  
431 edge).

432 Fig. 7a shows that the wetland extent predicted using the bathtub approach (Simulation 1) is affected by the  
433 sediment load, with only the low sediment condition resulting in a sharp decay in extent after 2060/70. The  
434 difference in extent is due to the vegetation retreat in the low sediment case, which does not occur in the high  
435 sediment case (Fig. 7b). Wetland extent values for the HST simulations are not greatly affected by the sediment  
436 load, and they are much smaller than the values predicted by the bathtub (Fig. 7a). Wetland retreat starts first in  
437 the simulations without the channel (Simulations 2 and 4) and about 20 years later in the simulations with the  
438 channel (Simulations 3 and 5) due to increased drainage. Once the retreat starts, it occurs faster in the simulations  
439 with the embankment (Simulations 4 and 5) that delays the ebb flows and increases hydroperiods in the lower  
440 wetland areas.

441 Wetland transgression is not affected by the sediment conditions (Fig. 7c) because of the limited amount of  
442 sediment that reaches the upper wetland areas. Transgression starts later in the simulations with the embankment  
443 (Simulations 4 and 5) because of the reduced depths and sediment loads in the upper wetland areas. The presence  
444 of the channel (Simulations 3 and 5) results in earlier but more gradual transgression compared to setups with no  
445 drainage structure (Simulations 2 and 4).

#### 446 **4 Discussion**

447 The interactions between all the processes related to the dynamic of coastal wetlands are quite complex  
448 (Fagherazzi et al., 2012; Reef et al., 2018; Saintilan et al., 2014), which makes the bathtub assumption limited for  
449 most applications. Places with multiple vegetation species (Cahoon et al., 2011; Rogers et al., 2006) and an  
450 intertwined channel network (D'Alpaos, 2011) present a strong heterogeneity of saltwater exposure and sediment  
451 delivery to the overbank areas that need a detailed description of flow and sediment processes (see also Coleman  
452 et al., 2020). Artificial structures constraining flow and sediment modify accretion rates (Bellafiore et al.,  
453 2014; Cahoon et al., 2011) and thus wetland evolution (Rodriguez et al., 2017; Sandi et al., 2018). Even though our  
454 simulation design focused on simplified setups, these setups comprise typical wetland features and include most  
455 of the complex processes and interactions.

456 Our results indicate that wetlands do not cope with SLR for the simulated conditions corresponding to a high  
457 emissions climate change scenario. This result was not surprising for the low sediment situation, as the inability  
458 of sediment-poor coastal wetlands to survive high levels of SLR due to low accretion rates has been reported  
459 before (Kirwan et al., 2010; Lovelock et al., 2015b; Rodriguez et al., 2017; Sandi et al., 2018; Schuerch et al., 2018).

460 However, the results for high sediment load seem to challenge some previous studies highlighting the potential of  
461 biophysical feedbacks to produce accretion rates comparable to SLR (D'Alpaos et al., 2007; Kirwan and Murray,  
462 2007; Kirwan et al., 2016b; Mudd et al., 2009; Temmerman et al., 2003) . In our case, the biophysical feedbacks  
463 with a high sediment load produced wetland accretion rates similar to SLR rates only for the bathtub simulation.  
464 Analysis of accretion rates indicate that all simulations start with similar rates in the vegetated areas, with about  
465 2.5 mm/yr and 7.5 mm/yr in the low and high sediment situation, respectively. For the low sediment case, the  
466 initial value compared very well with historic values for SE Australian conditions measured by Howe et al. (2009)  
467 and Rogers et al. (2006). For the high sediment case, an increase of the accretion value by a factor of three seems  
468 reasonable considering an increase of the sediment load by a factor of three (from 37 g/m<sup>3</sup> to 111 g/m<sup>3</sup>). Those  
469 starting values of accretion remain at approximately the same level over most of the time for the bathtub  
470 simulations, while they decrease for the HST simulations. The decrease is more marked for Simulations 2 and 4  
471 (which reach a value of about 1 to 1.5 mm/yr by 2050), than for the simulations with inner channel Simulations 3  
472 and 5 (which attain values of 2 mm/yr and 4 mm/yr by 2050 for low and high sediment conditions, respectively).  
473 The reduction of the magnitude of the biophysical feedbacks over time is due to the continuous upland migration  
474 of vegetation, which colonises upper areas with comparatively less water depth and sediment supply (see also  
475 Sandi et al. (2018)). The bathtub model predicts less migration and higher depths, so it consistently overestimates  
476 accretion rates.

477 Despite having reduced accretion rates when compared to the bathtub simulations, the HST simulations still show  
478 a noticeable difference in elevation gains depending on the sediment supply levels. Compared to the low sediment  
479 case, the high sediment supply case results in about twice the average accumulated accretion (Fig. 4). However,  
480 analysis of vegetation changes over time for low (Fig. 5) and high (Fig. 6) sediment loads reveal minimum  
481 differences between them. Analysis of Fig. 7 indicates that even though the increase in sediment load generates  
482 about twice the accretion, this extra elevation is not sufficient to prevent wetland submergence. Fig. 4 suggests  
483 that accretion rates of four times the historic values or more are needed for the wetlands to be able to cope with  
484 SLR.

485 Although the simulations carried out in this study were conducted on simplified domains they can capture the  
486 general response of more complex domains present in real wetlands, as shown by the comparison with entire  
487 wetland results from Rodriguez et al. (2017) and Sandi et al. (2018) in Fig. 4. Moreover, the features included are  
488 present in many coastal areas around the world and thus have wider implications. Our bathtub results for low  
489 sediment conditions predicting an initial increase in wetland extent early in the century and then a decrease after  
490 2060 agree with previous bathtub model predictions (Lovelock et al., 2015b; Rogers et al., 2012; Schuerch et al.,  
491 2018). However, using the HST framework our predictions indicate that the decrease may start as early as 2030  
492 for wetlands with tidal range close to 1.3 m (as represented in our study), over a wide range of sediment loads.  
493 We can expect that this accelerated wetland loss will affect many parts of the world, particularly in areas with  
494 micro to meso tidal range and heavily developed coasts, like eastern Australia (Williams and Watford, 1997),  
495 parts of eastern US (Crain et al., 2009), western US (Thorne et al., 2018) eastern China (Tian et al., 2016) and  
496 western Europe (Gibson et al., 2007). In these environments, attenuation can be important due to man-made  
497 structures, and transgression may be limited by development (Doody, 2013; Geselbracht et al., 2015; Kirwan and  
498 Megonigal, 2013), so we can expect a behaviour closer to that of simulations 4 and 5. On the other hand, wetlands  
499 with dense drainage networks like the Venice Lagoon in Italy (Silvestri et al., 2005), the Scheldt Estuary in the

500 Netherlands (Temmerman et al., 2012), the North Inlet in South Carolina, US (Morris et al., 2005), would probably  
501 behave similarly to simulation 3 and experience comparatively smaller losses of area.

502 The results presented in this study show generalized conditions of wetland dynamics under sea-level rise by using  
503 several simplified domains that focus on individual mechanisms affecting ecogeomorphic evolution. This  
504 approach can support a broader perspective on the potential fate of coastal wetlands in general, but some  
505 limitations arise as part of the model assumptions. As with most wetland evolution models, we did not consider  
506 soil processes other than accretion, disregarding swelling, compaction and deep subsidence. Measurements in  
507 wetlands of the Hunter Estuary show that long-term surface elevation changes are mostly due to accretion,  
508 supporting our assumption (Howe et al., 2009; Rogers et al., 2006). Another process that we did not consider was  
509 the effects of marsh edge retreat due to ocean or wind waves (Carniello et al., 2012; Fagherazzi et al., 2012), which  
510 can have a significant role in coastal wetland evolution. Most coastal wetlands in Australia are estuarine and not  
511 exposed to ocean waves, whereas wind effects in our wetland were not important due to the absence of large open  
512 water areas where wind waves could fully develop. We also simplified the tidal signal without including neap-  
513 spring cycles, which sped up computations but may have affected the results. However, preliminary tests including  
514 neap-spring tide variability showed only small differences in the initial landward edge of saltmarsh, which did not  
515 affect the accretion dynamics due to the small depths and low sediment availability in that area. Finally, our  
516 simulations did not include the effect of storms, which can influence sediment availability, water depths and  
517 velocities. We believe that in our case excluding storm effects is justifiable based on Rogers et al. (2013), who  
518 found that in these fine sediment environments storms affect accretion dynamics over the short term (immediate  
519 erosion or low accretion followed by increased deposition over the next months), but they do not change the long-  
520 term trend of accretion and elevation gain rates.

## 521 **5 Conclusion**

522 We conducted detailed numerical simulations on the response to SLR of four different typical coastal wetlands  
523 settings, including the case of a vegetated tidal flat free from obstructions and drainage features, and three other  
524 settings that included an inner channel, an embankment with a culvert, and a combination of inner channel,  
525 embankment and culvert. We also included a simulation using a simple bathtub approach, in which none of the  
526 features (vegetation, channels, culverts) are considered. We used conditions typical of SE Australia in terms of  
527 vegetation, tidal range and sediment load, but we also analysed simulations with an increased sediment load to  
528 assess the potential of biophysical feedbacks to enhance accretion rates.

529 We found that the distinct patterns of flow and sediment redistribution obtained from these simulations result in  
530 increased wetland vulnerability to SLR when compared to predictions using the simple bathtub approach. Changes  
531 in elevation due to accretion were between 10% and 50% of those obtained from bathtub predictions, and wetland  
532 retreat and reduction of wetland extent started 20 to 40 years earlier than for the case of the bathtub simulations,  
533 depending on wetland setting. Transgression for all settings was delayed with respect to the bathtub predictions  
534 and was limited by the presence of a hard barrier at the upland end.

535 The simulations using the full hydrodynamic and sediment transport dynamic models indicated that wetlands with  
536 good drainage (e.g. including an inner channel) were more resilient to SLR, displaying more accretion, a later  
537 retreat and reduction of wetland area and an increased transgression when compared with wetlands with strong  
538 flow impediments (e.g. including an embankment).

539 Increasing the sediment load delivered to the wetlands by a factor of three increased the accretion of all wetland  
540 settings by a factor of two. However, this extra elevation was not enough to prevent wetland submergence, as  
541 predictions of wetland evolution were very similar for low and high sediment conditions. Based on our results,  
542 we estimate that accretion rates of four times the typical historic values or more would be needed for these  
543 wetlands to cope with SLR.

544 Even though the characteristics of the wetlands studied here correspond mainly to SE Australian conditions, our  
545 results have a wider relevance because they clearly link the capacity of wetlands to accrete and migrate upland,  
546 the two mechanisms by which wetlands can gain elevation and keep up with SLR. Failure to consider the spatial  
547 coevolving nature of flow, sediment, vegetation and topographic features can result in overestimation of wetland  
548 resilience. Our results reconcile the wide discrepancy between upper thresholds of wetland resilience to sea-level  
549 rise in previous modelling studies with those emerging from paleo-stratigraphic observations.

550

551 *Data availability.* Upon acceptance of the manuscript, the hydrodynamic model and simulation results will be  
552 available from the corresponding authors on request.

553

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

555

556 *Author contribution.* A.B., P.M.S. and J.F.R. designed the study. A.B. calibrated and fitted the models and run the  
557 simulations. A.B., J.F.R., P.M.S., S.S., G.R. and N.S. analysed the results. A.B., P.M.S. and J.F.R. wrote the paper  
558 with substantial input from all co-authors.

559

560 *Acknowledgements.* P.M.S. acknowledges support from the Australian Research Council (grant FT140100610).  
561 A.B. was supported by a University of Newcastle PhD scholarship.

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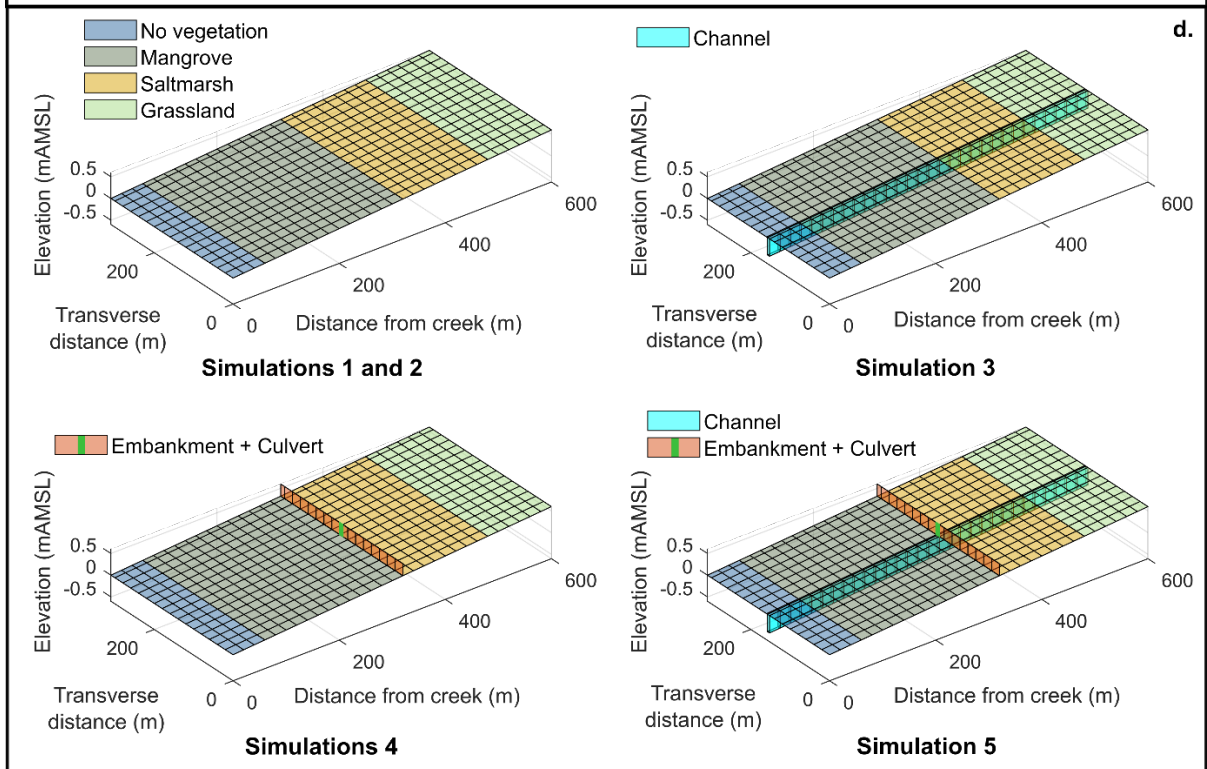
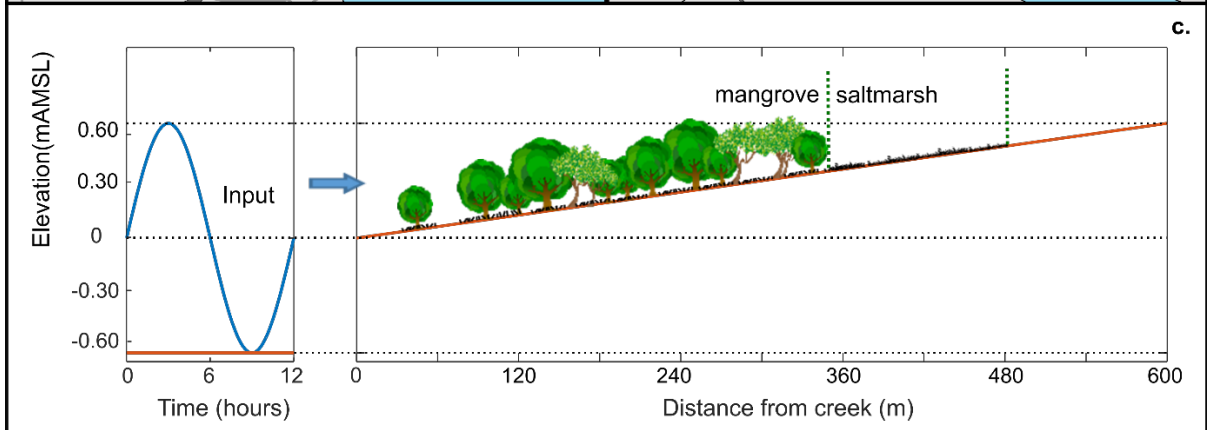
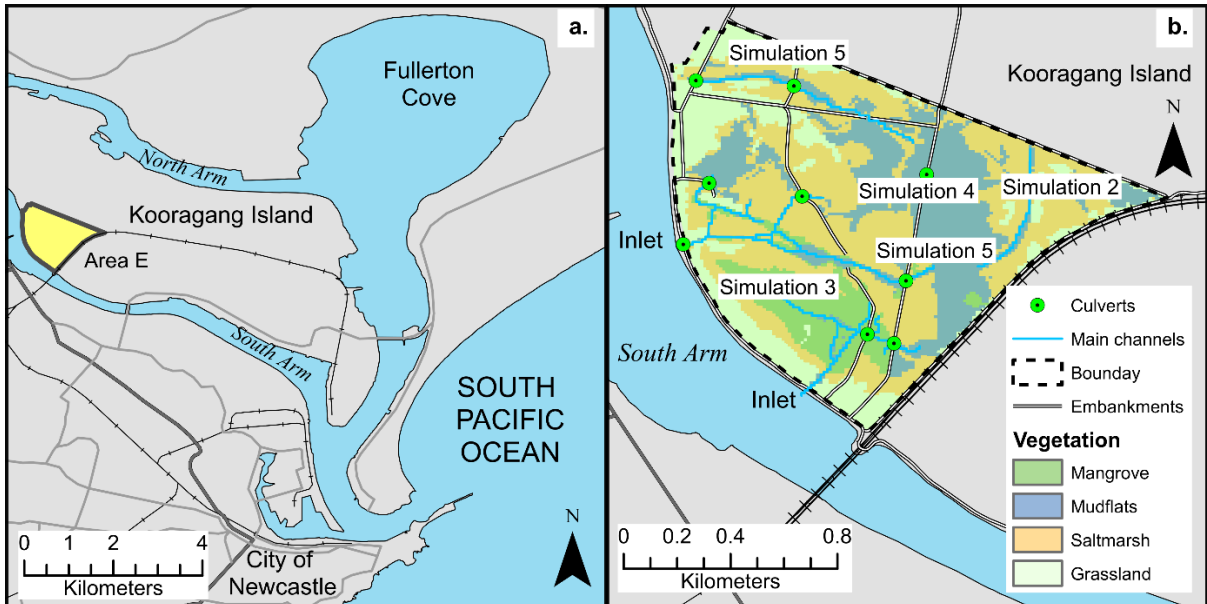
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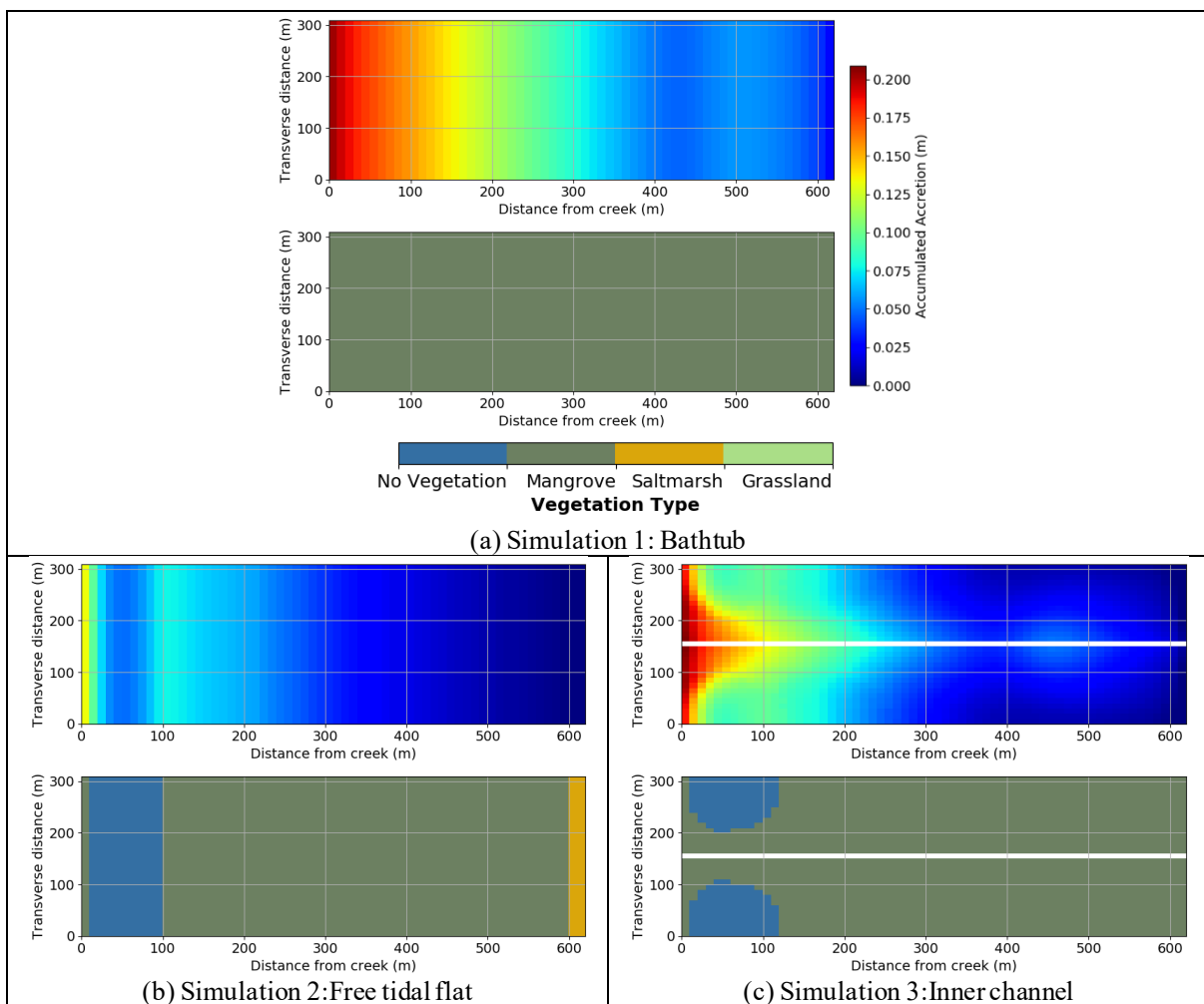
773 **Figure 1. Field site and areas within the site characterised by the numerical simulations: a) Area E of Kooragang**  
 774 **wetlands, b) areas within the wetland where the simplified simulations represent the dominant processes, c) schematic**  
 775 **longitudinal view of the domain setup and sinusoidal wave input (adapted from Rodriguez et al. (2017)), d) schematic**  
 776 **isometric view of each simulated domain and their hydraulic features. Vegetation cover is only indicative and roughly**  
 777 **corresponds to early stages of the simulations. Elevation unit, mAMSLL, stands for metres above mean sea level.**

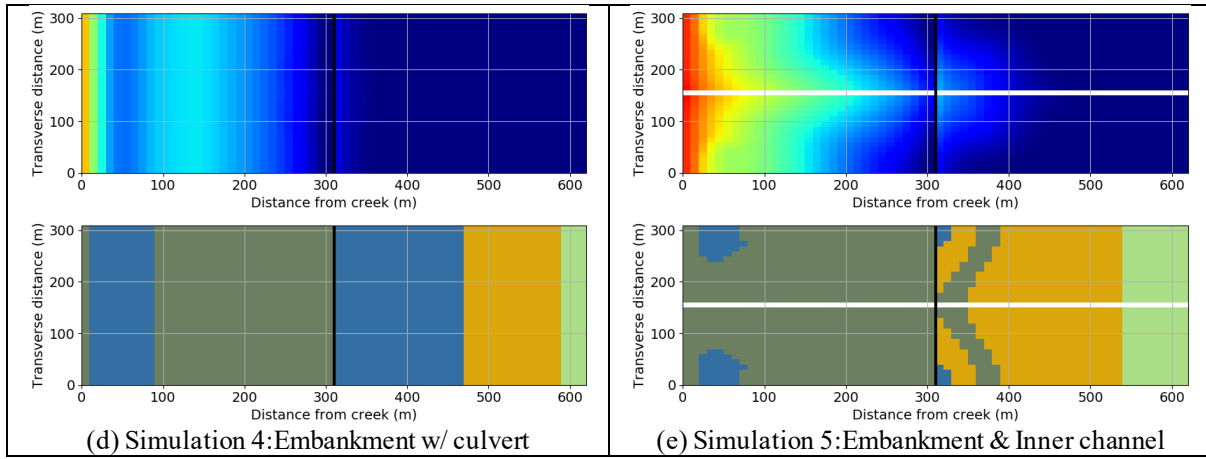
778 **Table 1: Parameters of soil surface elevation model**

Model Parameter	Mangrove	Saltmarsh
$a$ (g/m <sup>4</sup> )	-6,037.6	-16,767
$b$ (g/m <sup>3</sup> )	7,848.9	8,384
$c$ (g/m <sup>2</sup> )	-1,328.3	0
$q$ (m <sup>3</sup> /year/g)	$9 \times 10^{-5}$	$9 \times 10^{-5}$
$k$ (m <sup>5</sup> /g <sup>2</sup> )	$1.2 \times 10^{-7}$	$6.2 \times 10^{-7}$

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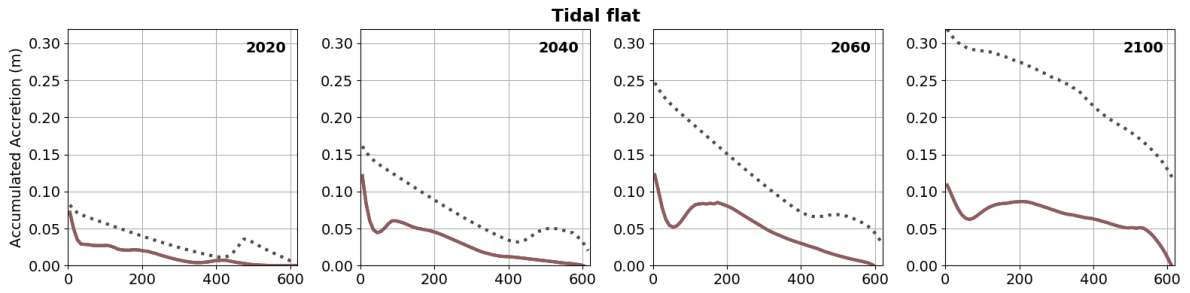
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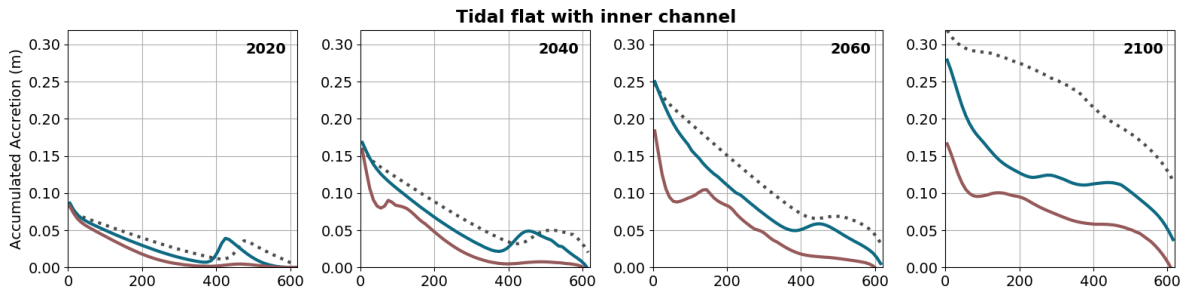


781 **Figure 2. Accumulated accretion (top) and vegetation maps (bottom) in 2050 for low sediment input corresponding to:**  
 782 **a) Simulation 1, b) Simulation 2, c) Simulation 3, d) Simulation 4, e) Simulation 5.**

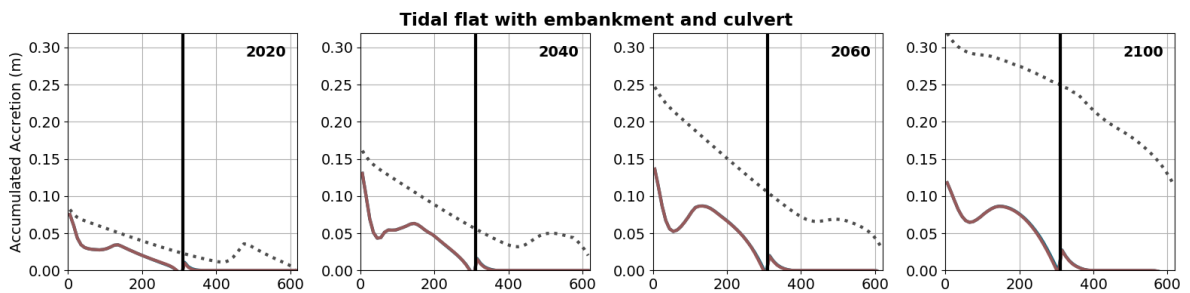
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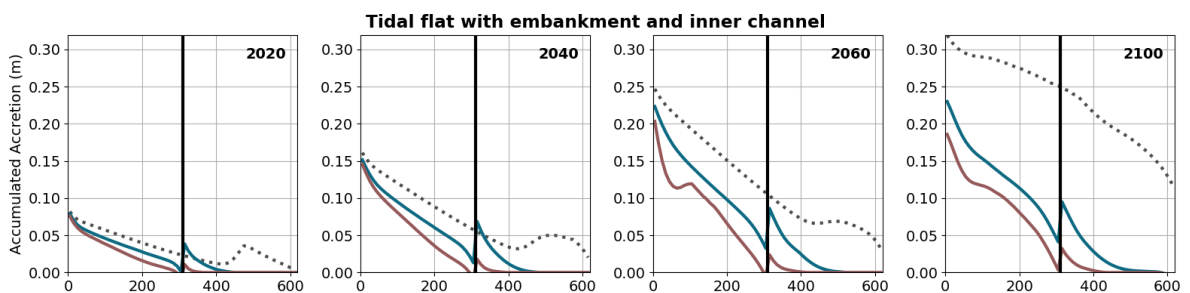
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..... Bathtub    — Channel    — Tidal Flat

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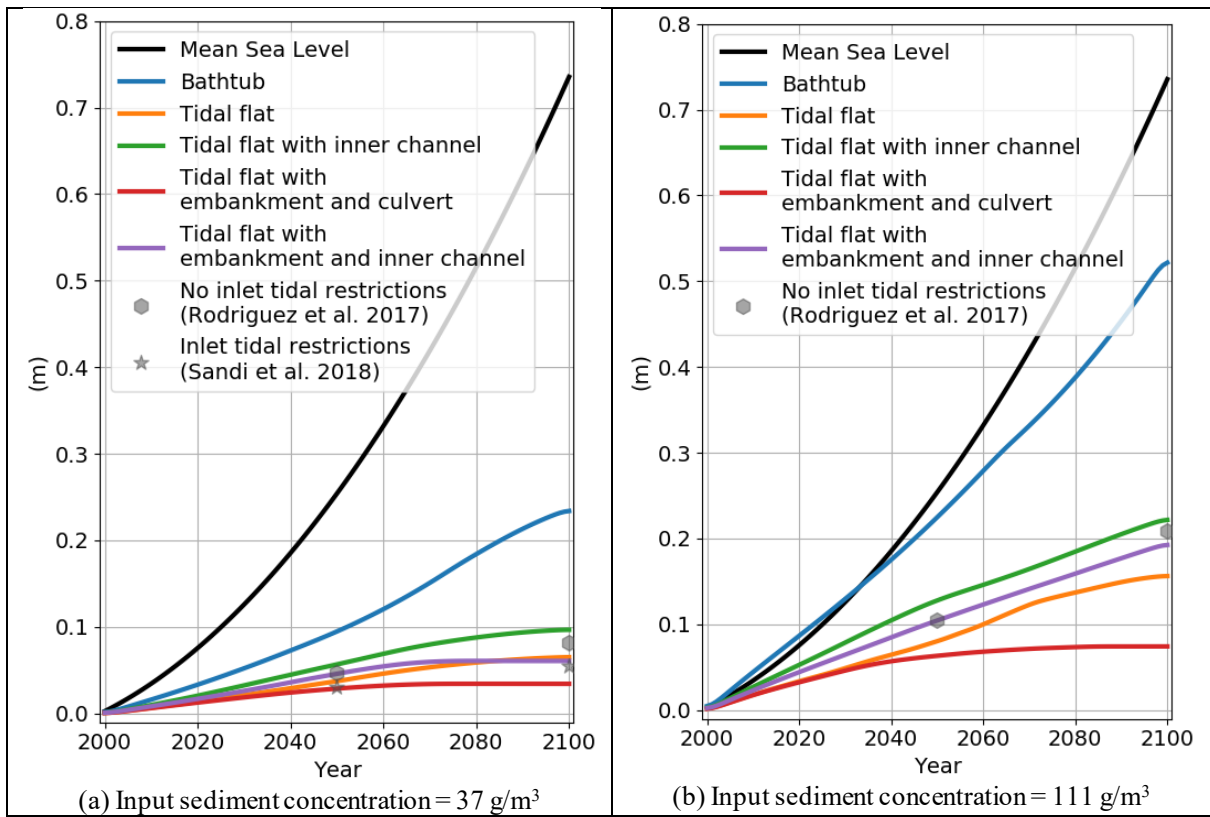
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**Figure 3. Longitudinal profiles of accumulated accretion ( $\Delta E$ , m) for a sediment supply of  $37 \text{ g/m}^3$ . The vertical black line represents the embankment with culvert. The “channel” profile represents the elevation gain near the central channel, while the “tidal flat” profile is situated in the middle of the tidal flat. Note: simulation starts in the year 2000.**

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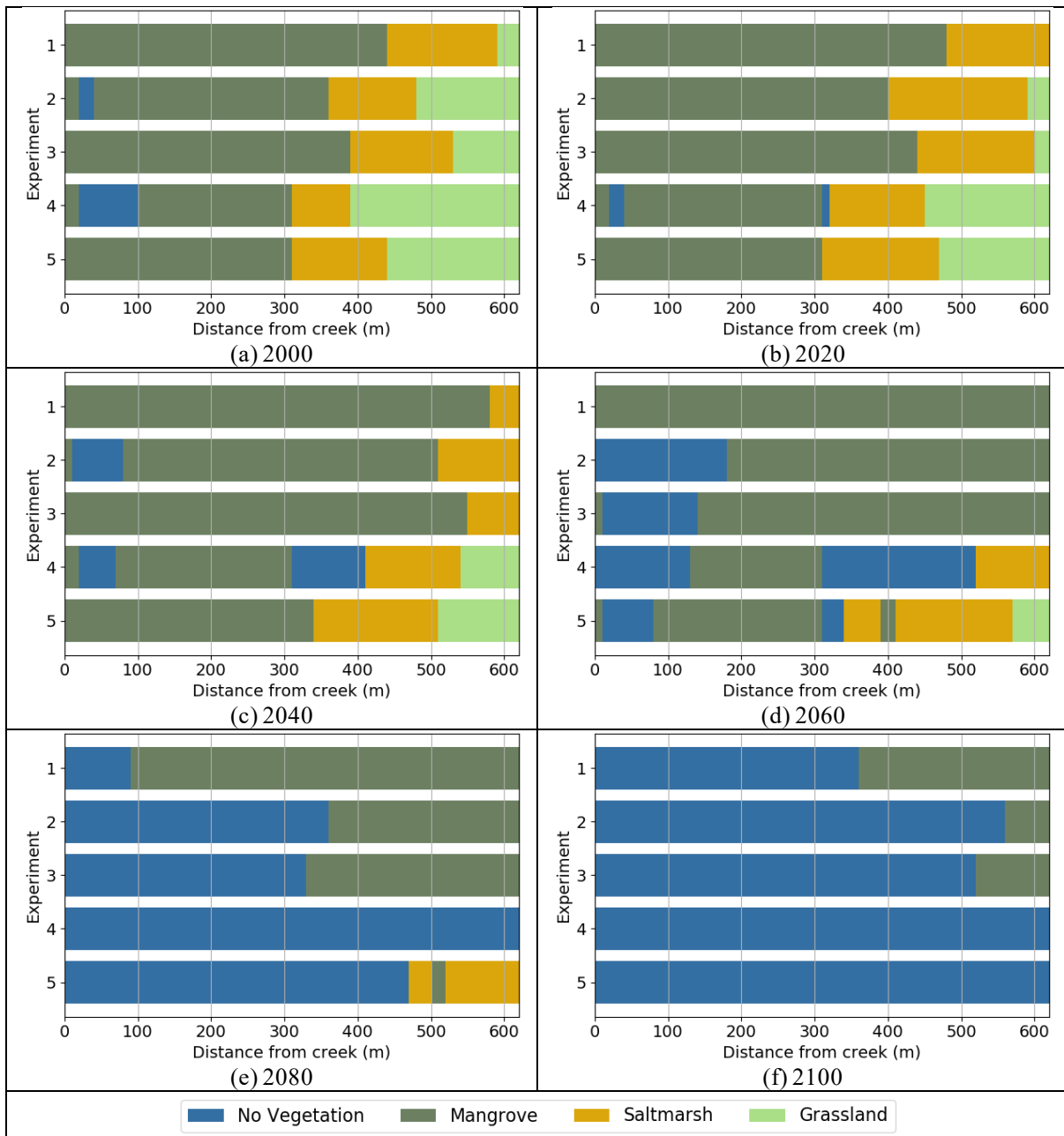
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793 **Figure 4. Sea-Level Rise and domain-average accumulated accretion over time for all simulations for a) low sediment**  
 794 **input and b) high sediment input. Results from Rodriguez et al. (2017) and Sandi et al. (2018) corresponding to the**  
 795 **entire Area E wetland are included for comparison.**

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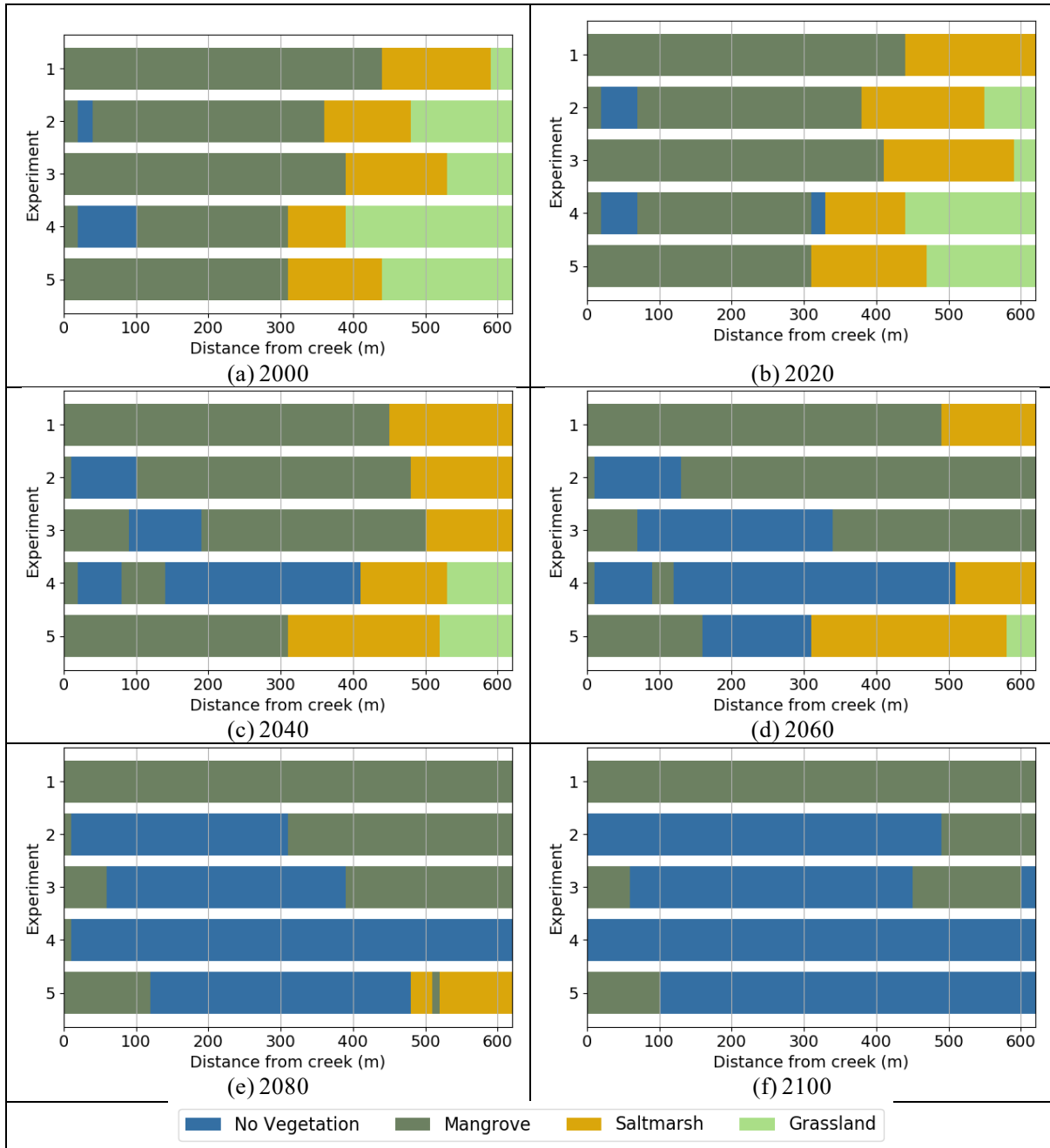


809 **Figure 5. Predominant position occupied by each vegetation type in the tidal flat from 2000 to 2100. Simulations for low**  
 810 **sediment input,  $SSC = 37 \text{ g/m}^3$ . Experiments: 1 Bathtub, 2 Free tidal flat, 3 Inner channel, 4 Embankment with culvert**  
 811 **and 5 Embankment and inner channel.**

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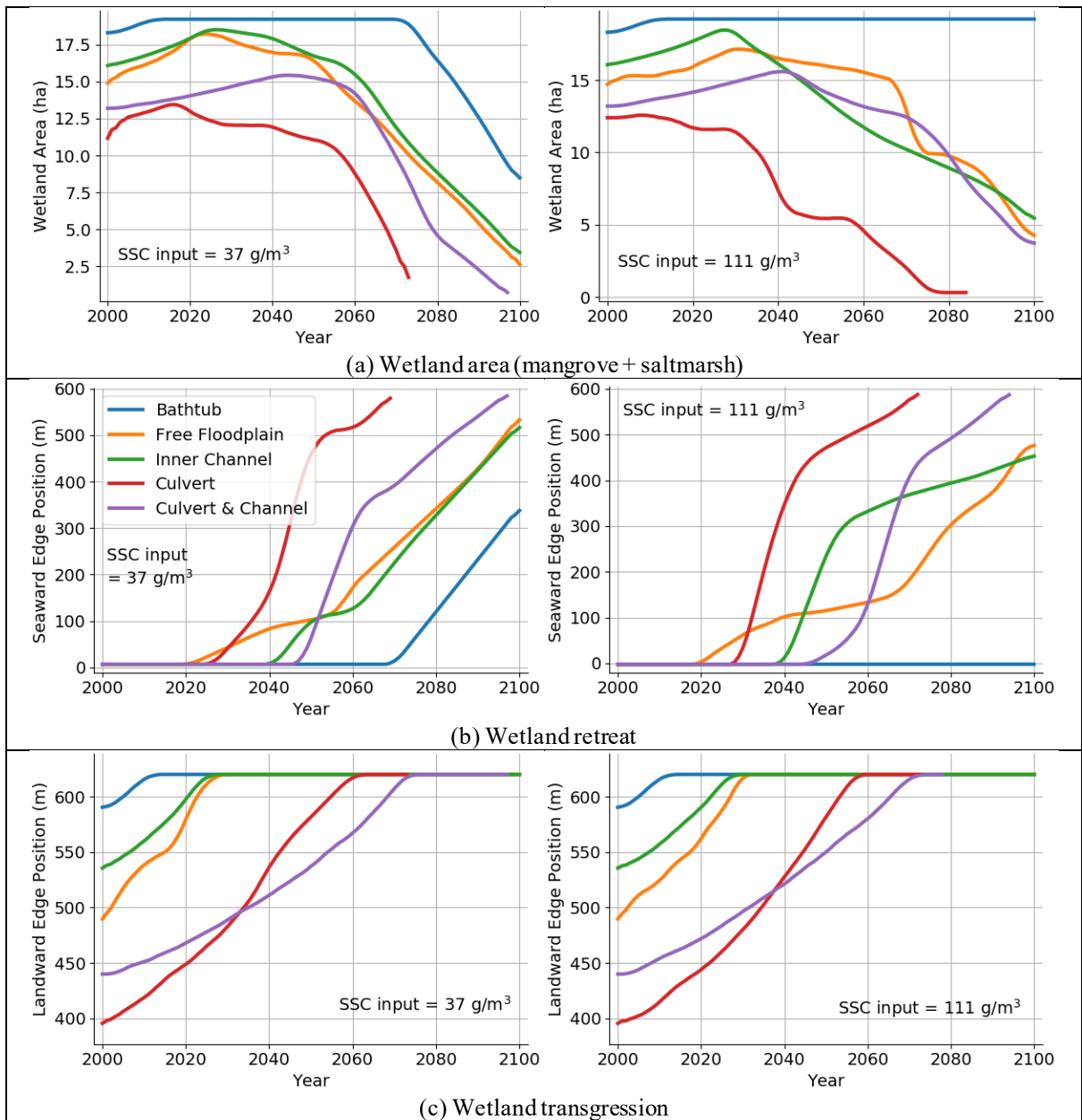


815 **Figure 6. Predominant position occupied by each vegetation type in the tidal flat from 2000 to 2100. Simulations for**  
 816 **high sediment input,  $SSC = 111 \text{ g/m}^3$ . Experiments: 1 Bathtub, 2 Free tidal flat, 3 Inner channel, 4 Embankment with**  
 817 **culvert and 5 Embankment and inner channel.**

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821 **Figure 7. Time evolution of wetland in poor ( $SSC=37 \text{ g/m}^3$ ) and rich ( $SSC=111 \text{ g/m}^3$ ) sediment environments under**  
 822 **SLR. a) wetland area; b) wetland retreat and; c) wetland transgression**

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