

1 **Morphometry of widespread tidal meander cutoffs discloses similarity to fluvial**  
2 **morphodynamics**

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21 **Key Points:**

- 22
- Tidal meander cutoffs are far more common than typically thought and share remarkable morphometric similarities with fluvial counterparts.
- 23
- Similar mechanisms trigger cutoffs in both tidal and fluvial landscapes, with differences arising only during post-cutoff evolution.
- 24
- Tidal cutoffs seldom disconnect from parent channels and form oxbow lakes due to the enhanced hydrological connectivity of tidal wetlands.
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- 27

28 **Abstract**

29 Similarities in planform dynamics between tidal and fluvial meandering channels contrast with  
30 the apparent lack in coastal wetlands of lateral-migration features like meander cutoffs and  
31 oxbow lakes, which led to the broad interpretation that tidal and fluvial meanders differ  
32 morphodynamically. We analyzed meander neck cutoffs from diverse tidal and fluvial  
33 environments worldwide, and show that tidal cutoffs are more widespread than believed. Their  
34 perceived paucity is a figment of coastal-wetland pronounced drainage density and hydrological  
35 connectivity, coupled with reduced size of most tidal channels and dense vegetation cover.  
36 Whereas these factors do not efface tidal meander cutoffs, they collectively inhibit oxbow-lake  
37 formation and render tidal cutoffs ephemeral, hardly detectable geomorphic features. We thus  
38 argue that similar morphodynamic processes drive cutoff formation in tidal and fluvial  
39 landscapes, with differences arising only during post-cutoff evolution. This bears important  
40 implications for understanding the ecomorphodynamics of coastal wetlands and predicting their  
41 long-term evolution.

42 **Plain Language Summary**

43 The sinuous channels that wander through tidal coastal wetlands look like meandering rivers.  
44 However, features of alluvial floodplains that indicate active river meandering over time, such as  
45 oxbow lakes and meander cutoffs, are difficult to find in tidal realms. Their apparent absence has  
46 led researchers to infer that tidal and river meanders differ in their evolutions. We re-examined  
47 this conclusion by identifying, measuring, and compiling examples of meander cutoffs from a  
48 variety of tidal coastal wetlands and fluvial floodplains worldwide. Our analysis suggests that the  
49 shapes and geometric properties of tidal and river cutoffs are indeed remarkably similar. This  
50 indicates that while tidal and fluvial environments differ in many ways, they nevertheless share  
51 the same physical mechanism affecting meander morphodynamical evolution. Differences do  
52 arise, however, after meanders have cut off. We observe that tidal meanders remain  
53 preferentially connected to the channel from which they originated, preventing the formation of  
54 crescent-shaped oxbow lakes and thus making tidal cutoffs more difficult to detect. Our results  
55 support a close similarity in meandering channel behavior across tidal and fluvial systems, thus  
56 opening new opportunities for how researchers model tidal wetlands numerically, with important  
57 implications for the effective conservation and restoration of these critical ecosystems.  
58

## 59 **1 Introduction**

60 Sinuous meandering channels are common in fluvial and coastal landscapes (Leopold et al.,  
61 1964). Meandering channels migrate laterally through erosion and deposition of sediment along  
62 the outer and inner banks, respectively, of individual meander bends. As meanders evolve,  
63 channels frequently shortcut themselves through cutoffs and form oxbow lakes (hereinafter  
64 "oxbows"; Dunne & Aalto, 2013; Ielpi et al., 2023; Mason & Mohrig, 2019; Schwenk et al.,  
65 2015; Stølum, 1996). Cutoffs play a critical role in channel and floodplain evolution by reducing  
66 channel sinuosity, modifying rates of lateral migration, and affecting floodplain sedimentology,  
67 stratigraphy, and sediment residence times (Ielpi et al., 2021, 2023; Schwenk et al., 2015; Zinger  
68 et al., 2011). Oxbows are important not only from ecological perspectives (Dieras et al., 2013;  
69 Thomas et al., 2022) but also because they retain signatures of the flow characteristics that  
70 shaped them (Guo et al., 2019). Collectively, the dynamics of meander cutoffs, of which oxbows  
71 are one consequence, have broad implications for the flux, storage, and sequestration of  
72 terrestrial organic carbon (Torres et al., 2017; Walcker et al., 2021).

73 While meandering river floodplains feature visible evidence of meander migration such as scroll  
74 bars and cutoffs (Constantine & Dunne, 2008; Dunne & Aalto, 2013; Hooke, 2013; Ielpi et al.,  
75 2023), channels in tidal coastal floodplains have been thought to lack meander cutoffs and,  
76 therefore, morphological evidence of active meandering (Gabet, 1998; Johnson & Campbell,  
77 1929) (Figure 1). The apparent tendency for sinuous tidal channels to be fixed in place – or at  
78 least the relative subtlety of their meandering dynamics – has been variously ascribed to  
79 ecomorphodynamics peculiar to coastal settings (Fagherazzi et al., 2004; Gabet, 1998; Garofalo,  
80 1980; Hughes, 2012; Kleinhans et al., 2009; Solari et al., 2002). However, recent studies  
81 highlighted common morphodynamics in fluvial and tidal meanders, with similar planform  
82 dynamics, width-adjusted migration rates, and morphodynamic regimes in high-amplitude bends  
83 (Finotello et al., 2018, 2022; Gao, Finotello, & Wang, 2022; Shaari et al., 2020). This motivated  
84 us to question the perceived paucity of tidal meander cutoffs, and to further demonstrate the  
85 parallels between tidal and fluvial meandering channels. Here, we analyzed the planform  
86 geometry of  $N_t=600$  tidal meander cutoffs identified in high-resolution satellite images from  
87 settings around the world, characterized by different tidal regimes, vegetation cover, and  
88 geomorphological backgrounds. Direct comparisons with cutoffs in meandering rivers ( $N_f=158$ )  
89 highlight geometric similarities that - with supporting evidence from theoretical, numerical, and  
90 field studies – are shared by morphodynamic processes in both tidal and fluvial realms.

## 91 **2 Materials and Methods**

### 92 *2.1 Data collection*

93 We used high-resolution satellite images, freely available from Google Earth Pro, to detect  
94 instances of meander cutoffs undisturbed by anthropic activities. The selected cutoffs encompass  
95 a wide variety of geographical locations, including coastal areas and inland alluvial plains, as  
96 well as a diversity of climatic and geological regions. Consequently, the sampled cutoffs reflect a  
97 range of hydrological and tidal regimes, sediment grain sizes, vegetation types, and land cover  
98 (Figure 1a-g). Our full dataset includes over 1200 examples of tidal cutoffs. Of these 1200  
99 examples, 600 tidal cutoffs with clearly discernable boundaries were manually digitized as  
100 polygons using Google Earth Pro. The remainder lacked sufficient detail to be digitized due to  
101 poor preservation, dense vegetation canopy, low image resolution, complex morphology  
102 resulting from multiple cutoffs, or combinations of these factors, and were categorized as

103 “unanalyzed cases” (Gao & Finotello, 2023). Furthermore, we obtained an additional set of 158  
 104 fluvial cutoffs specifically digitized for comparative analyses. These cutoffs were extracted from  
 105 rivers located in various regions, including the Amazon Basin, the conterminous USA and  
 106 Alaska, Russia, Canada, Kazakhstan, and New Zealand. The selection was made to ensure a  
 107 diverse range of channel sizes, with river widths spanning approximately four orders of  
 108 magnitude (Figure 2).

109 Tidal cutoffs were also further classified based on tidal regime (microtidal  $n=315$ ; mesotidal  
 110  $n=249$ ; macrotidal  $n=36$ ), vegetation cover (mangroves  $n=118$ ; salt marshes  $n=433$ ; tidal flats  
 111  $n=49$ ), and geomorphological setting (bays  $n=164$ ; back-barrier lagoons  $n=219$ ; open coasts  
 112  $n=105$ ; estuaries  $n=112$ ) (Figure S1 in Supporting Information). The mean tidal range ( $MTR$ ) for  
 113 each study site was determined by analyzing tidal gauge data from Dong (2020) and the National  
 114 Oceanic and Atmospheric Administration (<https://tidesandcurrents.noaa.gov/>), and individual  
 115 study cases were classified as macro-tidal ( $MTR > 4$  m), meso-tidal ( $2 < MTR < 4$  m), and  
 116 microtidal ( $MTR < 2$  m).

117 We focus only on ‘neck’ cutoffs, formed when a high-amplitude loop gets isolated by the  
 118 pinching connection of two adjacent bends. In the tidal settings we examined, we found no  
 119 examples of ‘chute’ cutoffs, which are formed when a river bend is shortcutted by a new channel  
 120 cutting through meander point bars.

## 121 2.2 Data analysis

122 To calculate their morphometric parameters, cutoff polygons were projected into appropriate  
 123 UTM coordinates and converted to binary images. The channel centerline was computed based  
 124 on a standard skeletonization procedure and then resampled using standard cubic spline-fit  
 125 polylines. Cutoff endpoints were determined as the two branchpoints of the polygon skeleton  
 126 (Figure 11). To further characterize cutoff planform features, we computed the curvature  $\mathcal{C}$  ( $[m^{-1}]$ )  
 127 of the channel centerline as  $\mathcal{C} = -d\theta/ds$ , where  $\theta$  is the angle between the tangent to the  
 128 channel axis and an arbitrarily selected reference direction,  $x(s)$  and  $y(s)$  are the Cartesian  
 129 coordinates of a given centerline point, and  $s$  is the intrinsic (i.e., along-channel) coordinate,  
 130 assumed to be positive in the upstream (i.e., landward) direction. Because flow orientation within  
 131 tidal meanders changes with tidal phases, we hereinafter assume a river-like reference system in  
 132 which the terms ‘upstream’ and ‘downstream’ refer to landward and seaward directions,  
 133 respectively.

134 After computing curvature, a Savitzky–Golay low-pass filter was applied to smooth noise in the  
 135 original signal. Then, the apex of any individual cutoff was identified as the locus of maximum  
 136 curvature (Figure 11), and the cutoff asymmetry index was computed as  $\mathcal{A} = (\ell_u - \ell_d)/(\ell_u + \ell_d)$  ([-])  
 137 where  $\ell_u$  and  $\ell_d$  are the distances between the cutoff apex and its upstream and  
 138 downstream endpoints, respectively (Figure 11). Negative values of  $\mathcal{A}$  correspond to upstream-  
 139 skewed cutoffs, and positive values of  $\mathcal{A}$  to downstream-skewed cutoffs. Other morphometric  
 140 parameters were also calculated, including: average channel width  $W$  ([-]); cutoff intrinsic length  
 141  $\ell = \ell_u + \ell_d$  ([m]); cutoff cartesian length  $L$  ([m]), which is the planar distance between cutoff  
 142 endpoints; cutoff sinuosity  $\chi = \ell/L$  ([-]); cutoff amplitude  $A$  ([-]), computed as the maximum  
 143 point-line distance between the cutoff centerline and the line connecting the two cutoff  
 144 endpoints; cutoff radius of curvature  $R$  ([m]), defined as the radius of the best-fitting circle  
 145 through all cutoff axis points; and flow-diversion angle  $\Phi$  between the cutoff and its parent  
 146 channel (Figure 11). Because of bidirectional flow through tidal channels, morphodynamically  
 147 meaningful flow-diversion angles can be identified at both the cutoff upstream ( $\Phi_u$ ) and

148 downstream ( $\Phi_d$ ) ends. By comparison, because of the unidirectional flow through river  
 149 channels, only the upstream flow-diversion angle ( $\Phi_u$ ) is morphodynamically meaningful for  
 150 fluvial cutoffs (Dieras et al., 2013).

151 To directly compare meander cutoffs of different sizes, dimensional morphometric variables  
 152 were normalized using channel width ( $W$ ), such that width-adjusted cutoff radius of curvature,  
 153 amplitude, and lengths are defined as  $R^* = R/W$ ,  $A^* = A/W$ ,  $L^* = L/W$ , and  $\ell^* = \ell/W$ .

### 154 **3 Results and Discussions**

155 Dimensional morphometric features of individual cutoffs –  $R$ ,  $A$ , and  $\ell$  – all exhibit statistically  
 156 significant ( $p$ -value  $< 0.01$ ) power-law relationships to cutoff width  $W$  with matching best-fit  
 157 power-law exponents and limited separation in power-law scaling constants (i.e., prefactors;  
 158 Figure 2 and Figure S2 in Supporting Information). We also found a statistically significant  
 159 quasi-linear relation between  $L$  and  $W$  (Figure S3 in Supporting Information), with  $L \cong W$ . The  
 160 latter has been described previously as the condition leading to neck cutoff (Li et al., 2022),  
 161 whereas  $L < W$  thus represents a geometrically impossible configuration (Hayden et al., 2021).  
 162 Similarly, radius of curvature  $R = W/2$  represents a physically meaningful lower bound, since the  
 163 edges of a channel centerline with a radius of curvature smaller than half its width would  
 164 intersect each other (Hayden et al., 2021). Although theoretically there are no physical limits to  
 165 the development of both  $A$  and  $\ell$  (besides the basic requirements that  $A > 0$  and  $\ell > L$  in order for a  
 166 centerline to be sinuous), meandering channel planforms in reality are fractal (Montgomery,  
 167 1996; Nikora, 1991). As a consequence, the prevalence of smaller curves weights the distribution  
 168 of meander features toward the physically meaningful lower bound (Hayden et al., 2021;  
 169 Vermeulen et al., 2016). Therefore, the observed scaling similarities might be due to the finite-  
 170 width nature of the sinuous features we measured. Despite slight differences in the power-law  
 171 scaling constants, the distributions reported in Figure 2 overlap substantially, making these  
 172 dimensional metrics a relatively poor diagnostic with which to distinguish the fluvial or tidal  
 173 nature of meander cutoffs (Figure S2 and Figure S3 in Supporting Information).

174 Characteristic differences do emerge, however, from the analyses of dimensionless cutoff  
 175 morphometrics. Kolmogorov-Smirnov (KS) tests performed at standard 5% significance level  
 176 highlight statistically significant differences between tidal and fluvial cutoffs for distributions of  
 177 width-adjusted radius ( $R^*$ ), amplitude ( $A^*$ ), and intrinsic length ( $\ell^*$ ), along with the width-  
 178 independent measures of sinuosity ( $\chi$ ), asymmetry ( $\mathcal{A}$ ), and flow-diversion angles ( $\Phi_u$  and  $\Phi_d$ ).  
 179 Tidal cutoffs have smaller  $R^*$ ,  $A^*$ ,  $\ell^*$ , and  $\chi$  (Figure 3 and Table S1 in Supporting Information).  
 180 Given that, before cutoff, each of these morphometric parameters is expected to increase with  
 181 time, our findings could be broadly interpreted as showing a lower morphodynamic maturity in  
 182 tidal cutoffs, thus pointing to a faster evolutionary trajectory. However, similar width-adjusted  
 183 meander migration rates in tidal and fluvial settings (Finotello et al., 2018) contrast with such an  
 184 interpretation. Furthermore, similar values of asymmetry ( $\mathcal{A}$ ) and upstream flow-diversion angle  
 185 ( $\Phi_u$ ) in tidal and fluvial cutoffs (Figure 3 and Table S1 in Supporting Information) not only  
 186 reflects similar morphodynamic maturity but also suggests that similar mechanisms trigger cutoff  
 187 formation, likely associated with the planform configuration of the parent channel (Dieras,  
 188 2013). Notably, both fluvial and tidal cutoffs exhibit negative median and peak values of the  
 189 asymmetry index  $\mathcal{A}$  (Figure 3e). That is, both types of cutoffs tend to be upstream-skewed,  
 190 supporting similarity in their dominant morphodynamic regime (*sensu* Seminara et al., 2001).

191 We thus suggest that the smaller size and lower maturity of tidal relative to fluvial cutoffs  
 192 depend on extrinsic factors broadly related to the hydro-ecogeomorphology of tidal wetlands

193 rather than on fundamental differences between tidal and fluvial meander morphodynamics.  
194 Specifically, we hypothesize that the high drainage density that typically characterizes tidal  
195 wetlands explains the comparably smaller size of tidal cutoffs, with enhanced hydrological  
196 connectivity justifying the apparent paucity of cutoff traces in tidal environments.

### 197 *3.1 Drainage density control on meander cutoff formation*

198 Meander migration in densely channeled tidal floodplains might shape the landscape differently  
199 than in fluvial contexts, where rivers can freely migrate laterally without intercepting other  
200 channels and confluences are comparatively infrequent. In contrast, the higher drainage density  
201 of tidal wetlands (*sensu* Marani et al., 2003) limits meander dynamics by preventing channels  
202 from freely migrating and meanders from fully developing without intercepting adjoining  
203 channels (Letzsch & Frey, 1980; Vilas et al., 1999). A similar dynamic is described in multi-  
204 thread, anastomosing rivers with individual meandering anabranches, where meander evolution  
205 is more likely to produce channel avulsions than meander cutoffs, due to enhanced drainage  
206 density compared to single-thread-meandering-river floodplains (Lagasse et al., 2004; Schumm  
207 et al., 1996). Accordingly, evidence from modern and ancient deposits shows that channel piracy  
208 (i.e., capture) in dense tidal networks (Figure S4 in Supporting Information) limits the lateral  
209 accretion of point bar bodies and can modify the network-scale distribution of the tidal prism,  
210 feeding back into the long term ecomorphodynamic evolution of the entire tidal system (Cosma  
211 et al., 2020; Finotello, Ghinassi, et al., 2020). Because tidal channel dynamics depends directly  
212 on the tidal prism (D'Alpaos et al., 2005; Finotello et al., 2018), which in turn responds to water  
213 surface gradients that generally decrease with decreasing distance among individual channels  
214 (Rinaldo et al., 1999), it follows that enhanced drainage density limits meander dynamics.

215 Our hypothesis is further corroborated by the evidence that distributions of  $R^*$ ,  $A^*$ ,  $L^*$ , and  $\chi$   
216 show only subtle variations across a range of tidal environments, with significant differences  
217 associated only with vegetation cover (Figure 3). Tidal cutoffs in salt marshes are smaller and  
218 less sinuous than those found in mangrove forests and tidal flats (Figure 3). This aligns with  
219 observations of tidal channel networks in salt marshes being more dense and hydrologically  
220 efficient (Kearney & Fagherazzi, 2016; Schwarz et al., 2022), and corroborates the idea that  
221 drainage density limits the development of tidal meandering cutoffs. Similar cutoff asymmetries  
222 ( $\mathcal{A}$ ) and flow-diversion angles ( $\Phi_u$ ,  $\Phi_d$ ) among distinct tidal settings also suggests similarity in  
223 the morphodynamic processes responsible for cutoff development. Kolmogorov-Smirnov tests  
224 reveal significant differences in distributions of  $\Phi_u$ ,  $\Phi_d$ , and  $\mathcal{A}$  only based on geomorphological  
225 setting (Tables S2 to S13 in Supporting Information), but we find no differences in these  
226 morphometrics as a function of tidal range and vegetation cover despite the potential influence  
227 that both controls can exert on channel bank erosion (Gao, Finotello, D'Alpaos, et al., 2022;  
228 Gasparotto et al., 2022; Zhao et al., 2022).

### 229 *3.2 Hydrological connectivity control on post-cutoff development*

230 To further substantiate that differences in tidal and fluvial cutoff morphology stem from extrinsic  
231 factors and not intrinsic meander morphodynamics, we also examined the connection state of  
232 individual cutoffs with their parent channels. Once a river meander cutoffs, a plug bar forms in  
233 response to flow separation and reduced energetic conditions, leading to the gradual blockage of  
234 both cutoff entrance (Toonen et al., 2012). Eventually the cutoff becomes completely  
235 disconnected from the parent channel and forms an oxbow lake. Based on the presence and  
236 position of plug bars in our tidal and fluvial examples, we classified cutoffs into four groups:

237 completely connected, upstream connected, downstream connected, and disconnected (Figure 4).  
238 The upstream- and downstream-connected cases can also be merged into a broader category of  
239 partially connected cutoffs. Whereas more than 43% of fluvial cutoffs in our dataset are entirely  
240 disconnected and only 28% are completely connected (Figure 4a), tidal cutoffs tend to remain  
241 connected to their parent channels, with 87% of examples completely connected, 9% partially  
242 connected, and only 4% entirely disconnected (Figure 4a).

243 This marked difference in the connection state of tidal versus fluvial cutoffs is not an apparent  
244 function of tidal range, vegetation cover, and geomorphological setting (Figure 4). Moreover,  
245 similar flow-diversion angles are observed in all our study cases, with median values consistently  
246 ranging between  $105^\circ$  and  $108^\circ$  (Figure 4b,c) and further pointing to similar cutoff-triggering  
247 mechanisms in fluvial and tidal landscapes. Morphological differences thus can be expected to  
248 emerge once cutoffs have formed. Remarkably, the percentage of completely connected fluvial  
249 cutoffs decreases as the flow-diversion angle increases, implying that larger  $\Phi_u$  promote the  
250 formation of plug bars and oxbows (Figure 4d). In contrast, tidal cutoffs tend to remain  
251 connected to their parent channel irrespective of flow-diversion angles, whether upstream or  
252 downstream (Figure 4d and Figure S5 in Supporting Information). Therefore, unlike fluvial  
253 analogs, most tidal cutoffs remain hydrodynamically active to some extent, as they continue to  
254 drain water from the adjoining tidal floodplains. Indeed, in stark contrast to meandering rivers,  
255 the rate of lateral flow injections in tidal channels can greatly exceed the baseline flow rates,  
256 resulting in a peculiar landward reduction of channel width (i.e., funneling; Lanzoni &  
257 D'Alpaos, 2015). Notably, some tidal cutoffs may also remain connected to other active parts of  
258 the network through minor lateral tributaries flowing directly into the cutoff (Figure 1a-I and  
259 Figure S6 in Supporting Information). Hence, pronounced hydrological connectivity in tidal  
260 wetlands prevents the formation of a plug bar and the subsequent evolution of tidal cutoffs into  
261 oxbows. Such an evolutionary trajectory clearly differs from fluvial cutoffs, which are typically  
262 abandoned and receive water and sediment input almost exclusively during major flood events  
263 (Leopold et al., 1964; Shen et al., 2021) either through minor tie channels carved through the  
264 plug bar (cf. Rowland et al., 2009) or as the entire alluvial plain floods (Shen et al., 2021).

265 Among the partially connected cutoffs in our dataset, the fluvial ones are preferentially  
266 connected with their parent channels at the upstream end: plug bars tend to form at the cutoff  
267 downstream end where flow separations and recirculation create a zone of dead velocity that  
268 hinders mixing and promotes the deposition of fine sediment (Richards et al., 2018; Turnipseed  
269 et al., 2021). In contrast, the few partially connected tidal cutoffs on record, tend to maintain  
270 connectivity at the downstream end, aligned with the direction of ebb flows (Figure 4a). This  
271 predominance can be attributed to the typically ebb-dominated character of tidal flows  
272 (Fagherazzi et al., 2008; Finotello, Ghinassi, et al., 2020), which keeps the cutoff downstream  
273 end periodically flushed.

### 274 *3.3 Meander cutoffs in tidal coastal landscapes: rare or everywhere?*

275 Abundant tidal cutoffs akin to oxbow-rich alluvial floodplains can be found in some tidal settings  
276 with possibly lower drainage density and/or sediment supply that limits cutoff infill and  
277 vegetation encroachment (Figure 1f-i and Figures S7 and S8 in Supporting Information). This  
278 further corroborates the observation that tidal and fluvial meandering channels not only evolve  
279 through similar morphodynamic processes, but also that tidal meanders are as prone to form  
280 cutoffs as their fluvial counterparts given conducive environmental conditions. One thus  
281 wonders, given the apparent ubiquity of cutoffs across a variety of tidal environments, why the

282 long-held notion prevailed that sinuous tidal channel bends are inherently unlikely to cut  
283 themselves off.

284 Our take is that, first, the characteristic width and amplitude of fluvial cutoffs may not vary  
285 significantly along a given reach of a meandering river between major tributaries, whereas  
286 meander cutoffs within a given tidal wetland can occur across a broad range of meander  
287 wavelengths and widths (Finotello, D'Alpaos, et al., 2020). Low-order, narrow tidal creeks are  
288 more frequently found than higher-order, wide channels and are thus the most likely to express  
289 cutoff development (Figure 1a-i and Figures S7 and S8 in Supporting Information). Yet small  
290 channels produce small cutoffs, which are especially challenging to observe from a broader  
291 spatial vantage particularly when the vegetation canopy is dense (e.g., in mangrove forests,  
292 Figure S9 in Supporting Information).

293 Another consideration is the sustained rate of vertical accretion that characterizes tidal wetlands,  
294 coupled with halophytic vegetation that can tolerate significant waterlogging stress. These  
295 factors may becloud cutoff traces (Figure 1b,d,f-I and Figure S9 in Supporting Information)  
296 through rapid sedimentation in the less hydrodynamically active portions of the cutoff, and the  
297 subsequent encroachment of vegetation. This levels out cutoff geomorphic expressions and  
298 further hinders their identification from aerial images. Although similar reasoning could apply to  
299 fluvial floodplains, reduced overbank sediment supply and slower rates of riparian vegetation  
300 growth in permanently waterlogged areas may prolong the time required to fill oxbows, making  
301 large river-cutoff scars identifiable from aerial photos for much longer periods (Figure 1j,k).

302 The apparent absence of tidal cutoffs is thus more an artifact of observations than a consequence  
303 of physical mechanisms. High drainage densities in tidal wetlands surely constrain the freely  
304 meandering of tidal channels (Figure S10 in Supporting Information). Yet the relatively small  
305 size of most tidal channels, along with the distinctive hydrological characteristics of tidal  
306 wetlands, contribute to the transient nature of tidal cutoffs and make them challenging to record.  
307 That is, unlike other features of meandering channels that might jump out at the observer, to find  
308 tidal cutoffs one has to go carefully looking for them.

309 The implied morphodynamic similarity between tidal and fluvial meanders is by no means  
310 diminished by the absence of prominent scroll bars in tidal wetlands, standing in stark contrast to  
311 river floodplains that often showcase intricate arrangements of sub-parallel scrolls indicative of  
312 previous channel locations (Figure 1k) (Dunne & Aalto, 2013; Strick et al., 2018). While there is  
313 no consensus on what drives the formation of scroll bars (van de Lageweg et al., 2014; Zen et al.,  
314 2017), we offer two possible, not mutually exclusive explanations for the absence of scroll bars  
315 in tidal meanders. One possibility is that tidal meanders undergo small incremental migrations  
316 during each tidal cycle, unlike fluvial meanders, which tend to migrate more episodically after  
317 the height of incrementally formed inner-bank levee-like features reaches a threshold condition  
318 (Mason & Mohrig, 2019; Wu et al., 2016). Another hypothesis is that sustained rates of vertical  
319 aggradation relative to lateral channel migration in tidal wetlands (cf. Brivio et al., 2016; Cosma  
320 et al., 2019) prevent scroll bars by systematically overshadowing any topographic irregularities.  
321 This explanation aligns with the lack of scroll bars in meandering streamflows evolving through  
322 curvature-driven fluvial-like mechanisms in aggradational settings such as coastal backwater  
323 areas (Swartz et al., 2020), peatlands (Candel et al., 2017; Guo et al., 2020), and submarine  
324 turbidity-current channels (Jobe et al., 2016; Morris et al., 2024)

#### 325 **4 Implications and Conclusions**

326 Our findings demonstrate that meandering channels in tidal wetlands are as capable of forming  
327 meander cutoffs as their fluvial counterparts. From the morphometric evidence we have  
328 compiled, we suggest that the morphodynamic processes that drive tidal and fluvial cutoff  
329 development are fundamentally similar, with substantial differences arising only after cutoffs  
330 have formed. Rather than forming oxbows, tidal cutoffs remain preferentially connected to their  
331 parent channel owing to the pronounced hydrological connectivity that characterizes tidal  
332 wetlands. In this way, tidal meander cutoffs continue to drain (and help flood) the surrounding  
333 wetlands, remaining active and integral parts of the overall tidal channel network. Coupled with  
334 earlier theoretical, numerical, and field observations (Finotello et al., 2018, 2022; Gao, Finotello,  
335 & Wang, 2022), our results point to a complete morphodynamic analogy between tidal and  
336 fluvial meandering channels from meander inception to cutoff. Unified tidal and fluvial meander  
337 morphodynamics enable extending classical techniques employed for modeling meandering  
338 rivers numerically (Bogoni et al., 2017; Howard & Knutson, 1984; Parker et al., 2011; Seminara  
339 et al., 2001) to ecomorphodynamic models of tidal wetlands, where meandering is ubiquitous  
340 and yet routinely omitted in common practices. Such an advance in numerical modeling would  
341 have important implications for the conservation and restoration of critically endangered tidal  
342 wetlands, for example by helping improve assessments and estimations of past, present, and  
343 future blue carbon fluxes.

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#### 351 **Conflict of Interest Statement**

352 The authors declare no conflict of interest.

#### 353 **Open Research**

354 All the data presented and analyzed in this paper are freely available from a public Zenodo folder  
355 (Gao & Finotello, 2023)

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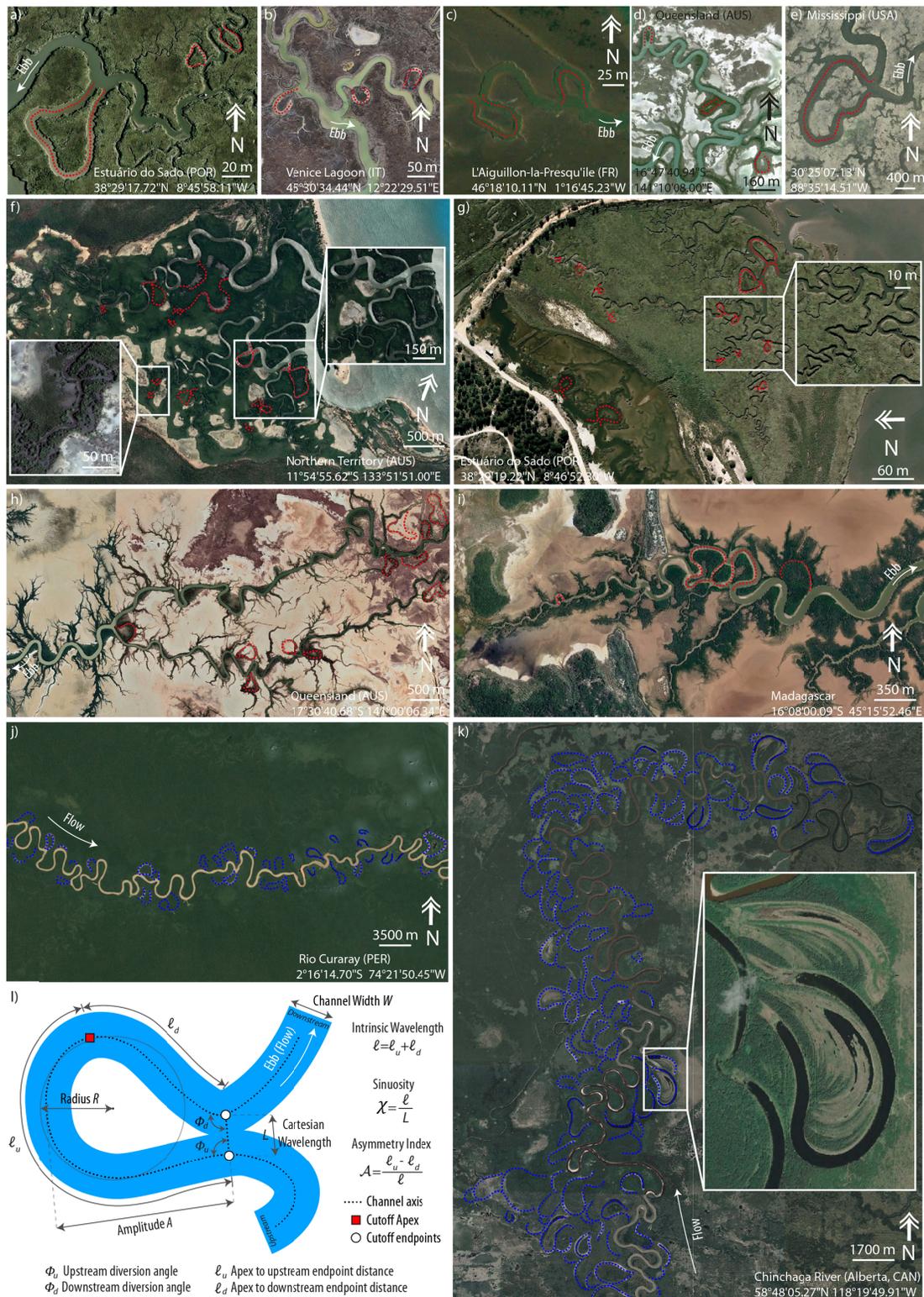
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573 **Figures and Tables**

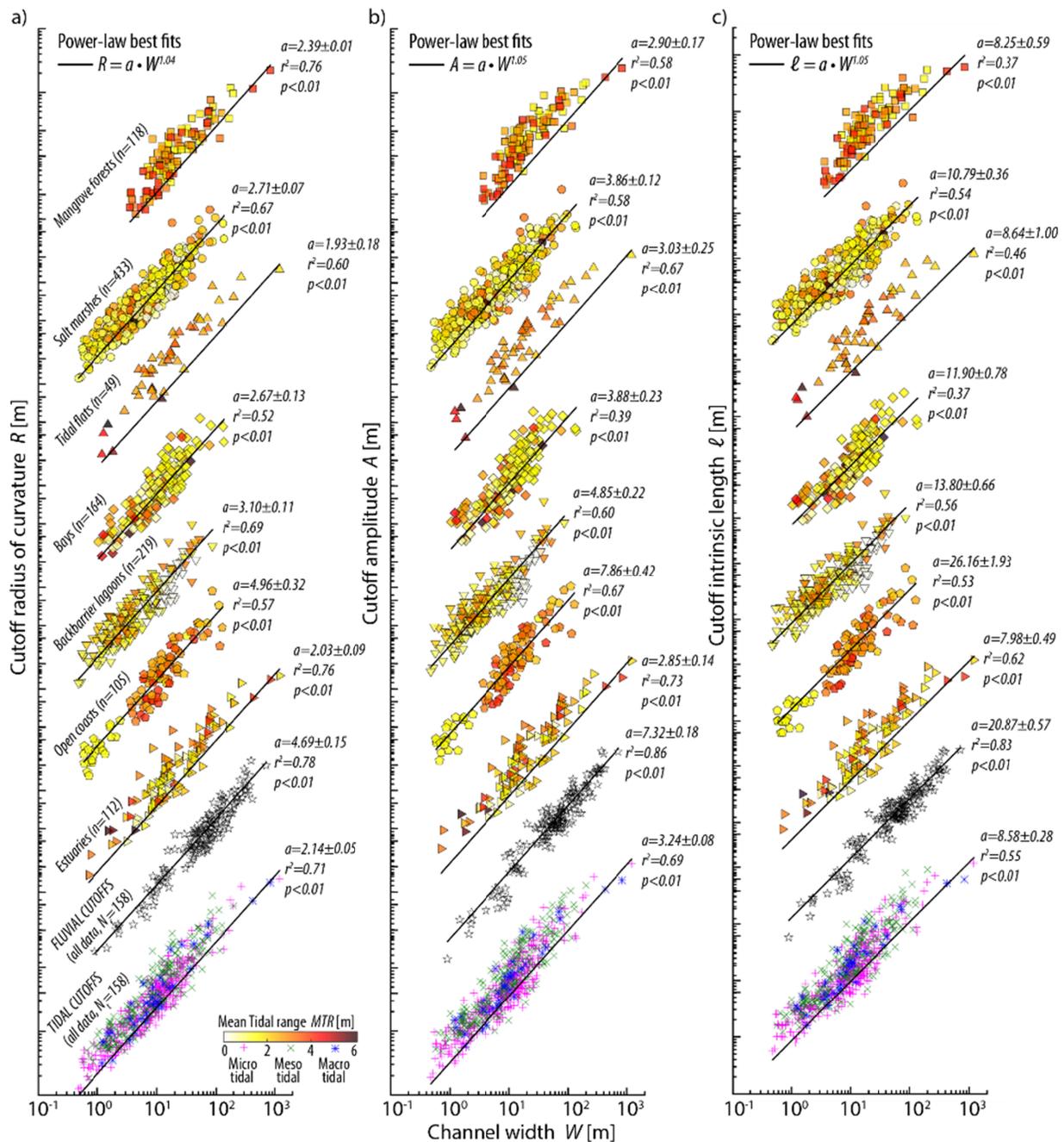


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**Figure 1. Meander cutoffs in tidal and fluvial landscapes.** (a,b,c,d) Examples of individual tidal meander cutoffs from distinct coastal settings worldwide (image© Google: Maxar Technologies and USDA/FPAC/GEO). (f,g,h,i) Examples of tidal environments characterized by widespread meander

578 cutoffs (image©Google: TerraMetrics, CNES/Airbus, Maxar Technologies, Landsat/Copernicus). (j,k)  
579 Examples of river floodplains littered by oxbow lakes and cutoff traces (image©Google: Maxar  
580 Technologies). Geographic coordinates are reported in each panel. Dotted red and blue lines highlight  
581 discernable traces of meander cutoffs in tidal and fluvial landscapes, respectively. l) Sketch illustrating  
582 the main morphometric features of meander cutoffs analyzed in this study.

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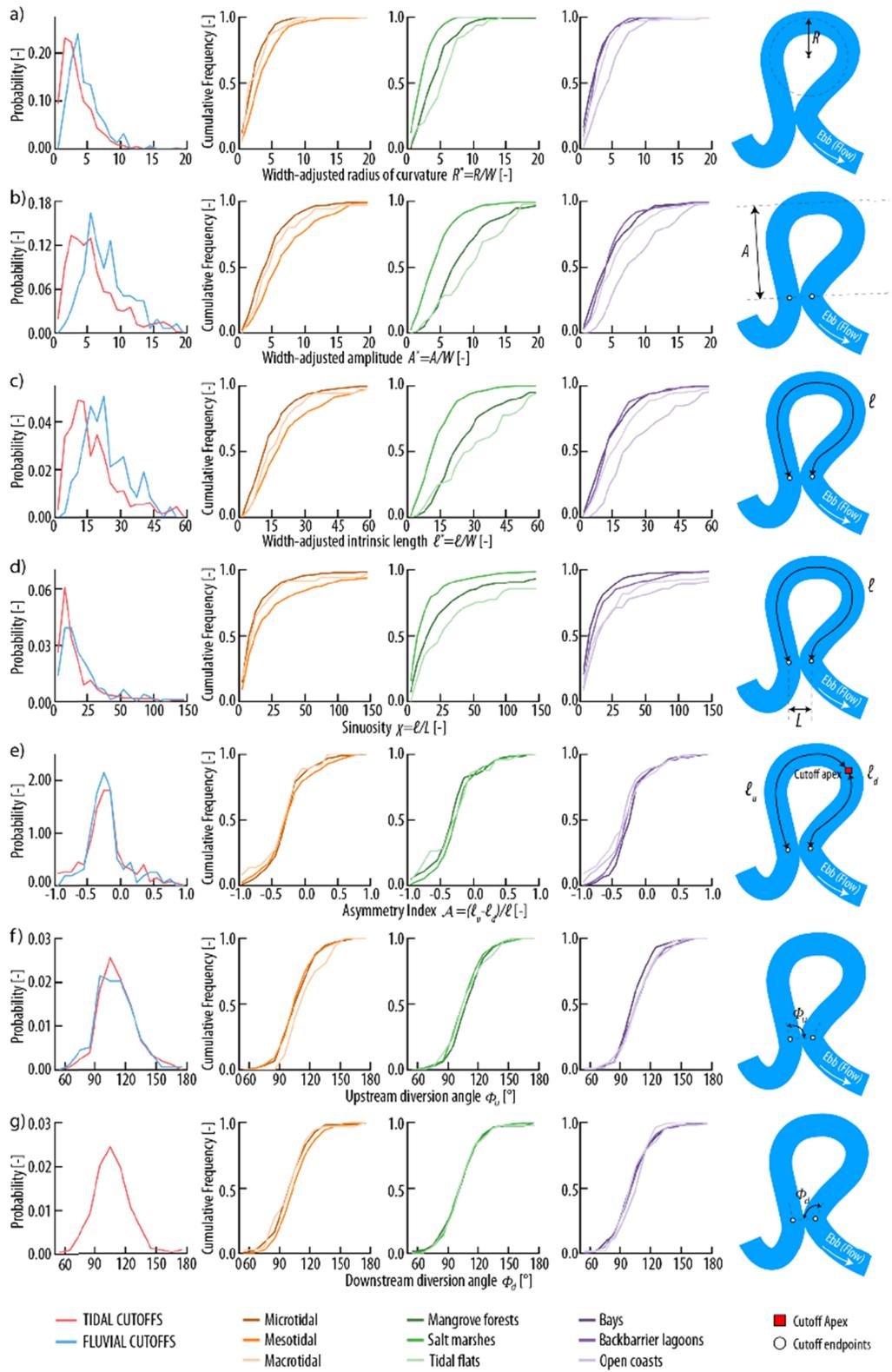
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**Figure 2. Cutoff morphometrics.** Cutoff radius of curvature ( $R$ ), Amplitude ( $A$ ), and intrinsic length ( $\ell$ ) are plotted against channel width ( $W$ ) both separately for all tidal and fluvial cutoffs on record and for different tidal-cutoff ensembles based on geomorphological settings and vegetation cover color-coded based on tidal ranges. Note the arbitrary vertical offset. Continuous black lines represent best-fit power law regressions obtained for different data ensembles, using a common exponent derived from all data and applied to calculate scaling coefficients for each ensemble.



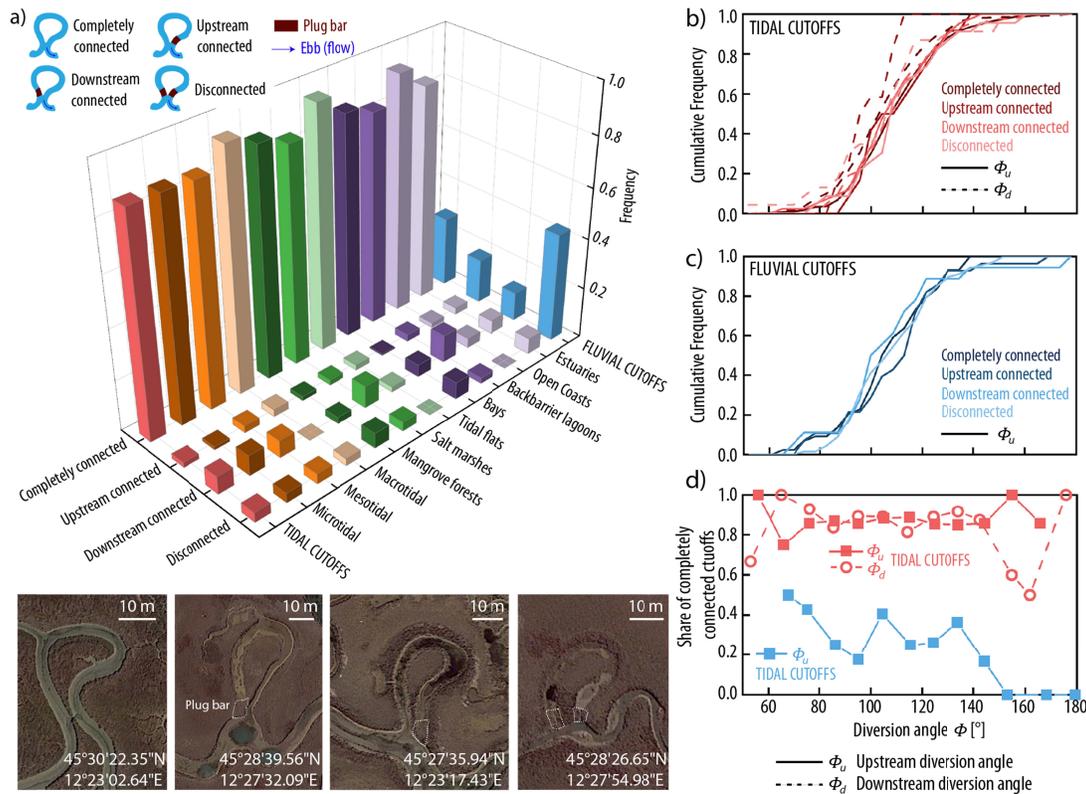
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**Figure 3. Dimensionless cutoff morphometrics.** (a) Width-adjusted radius of curvature; (b) width-adjusted amplitude (c) width-adjusted intrinsic length; (d) sinuosity ( $\chi$ ); (e) asymmetry index ( $\mathcal{A}$ ); (f,g)

595 upstream and downstream flow-diversion angles ( $\Phi_u$  and  $\Phi_d$ ). Panels in the first column show empirical  
596 probability distributions for tidal (red) and fluvial (blue) cutoffs. Panels in the other columns report  
597 empirical cumulative frequency distributions for tidal cutoffs subdivided based on tidal range, vegetation  
598 cover, and geomorphological setting. The fifth column contains sketch-up views for each investigated  
599 morphometric.

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**Figure 4. Cutoff connectivity.** (a) Barplot showing the relative frequency of different connection types between cutoffs and parent channels, differentiating tidal (red) and fluvial (blue) cutoffs, and further segmenting tidal cutoff ensembles based on tidal range (orange), vegetation cover (green), and geomorphological settings (purple). (b,c) Frequency distributions of flow-diversion angles ( $\Phi$ ) for tidal and fluvial cutoffs. Different colors denote different connectivity with the parent channel. Solid and dashed lines denote upstream and downstream diversion angles, respectively. (d) Share of completely connected tidal (red) and fluvial (blue) cutoffs across uniform  $10^\circ$  diversion-angle intervals. Solid squares and empty dots denote upstream and downstream diversion angles, respectively. (e) Tidal cutoffs found in the microtidal lagoon of Venice (Italy) characterized by different connectivity.