

Morphometry of widespread tidal meander cutoffs discloses similarity to fluvial morphodynamics

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Key Points:

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

Abstract

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

Plain Language Summary

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

1 Introduction

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

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

2 Materials and Methods

2.1 Data collection

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

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

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

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

2.2 Data analysis

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

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

downstream (Φ_d) ends. By comparison, because of the unidirectional flow through river channels, only the upstream flow-diversion angle (Φ_u) is morphodynamically meaningful for fluvial cutoffs (Dieras et al., 2013).

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

3 Results and Discussions

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

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

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

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

3.1 Drainage density control on meander cutoff formation

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

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

3.2 Hydrological connectivity control on post-cutoff development

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

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

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

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

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

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

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

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

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

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

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

4 Implications and Conclusions

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

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Conflict of Interest Statement

The authors declare no conflict of interest.

Open Research

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

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573 **Figures and Tables**

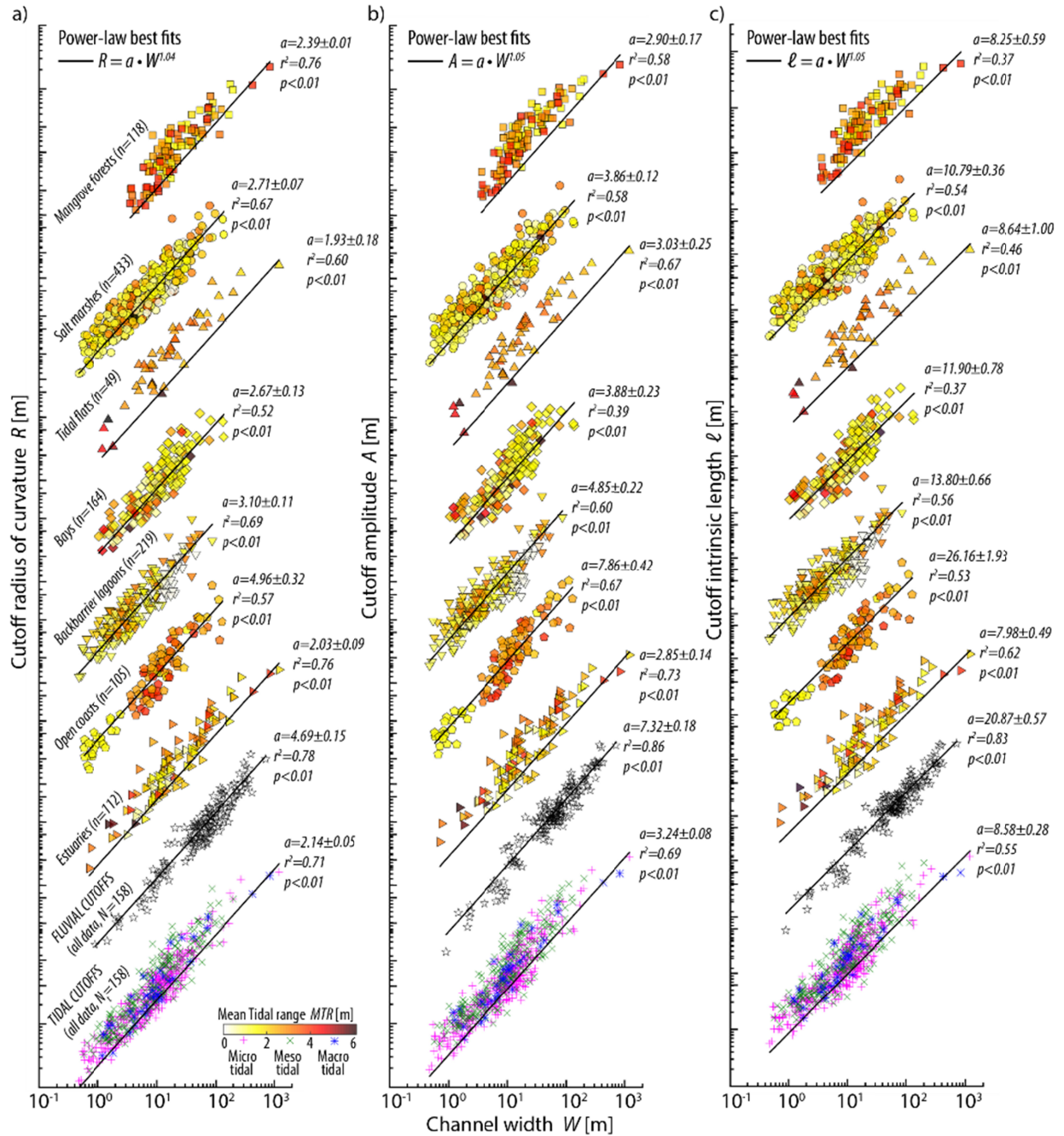


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575 **Figure 1. Meander cutoffs in tidal and fluvial landscapes.** (a,b,c,d) Examples of individual tidal
576 meander cutoffs from distinct coastal settings worldwide (image© Google: Maxar Technologies and
577 USDA/FPAC/GEO). (f,g,h,i) Examples of 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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Figure 2. Cutoff morphometrics. Cutoff radius of curvature (R), Amplitude (A), and intrinsic length (ℓ) 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.

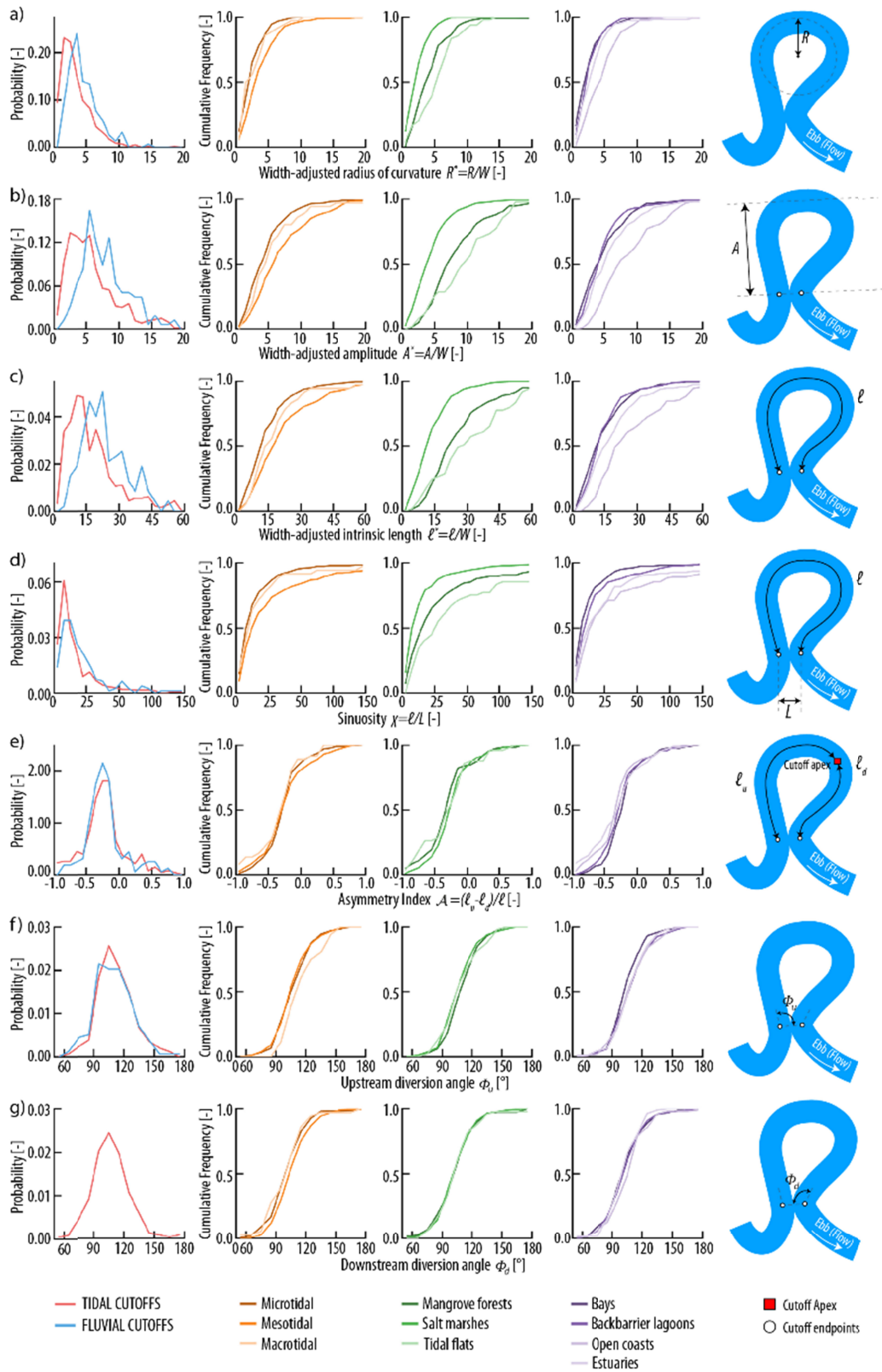
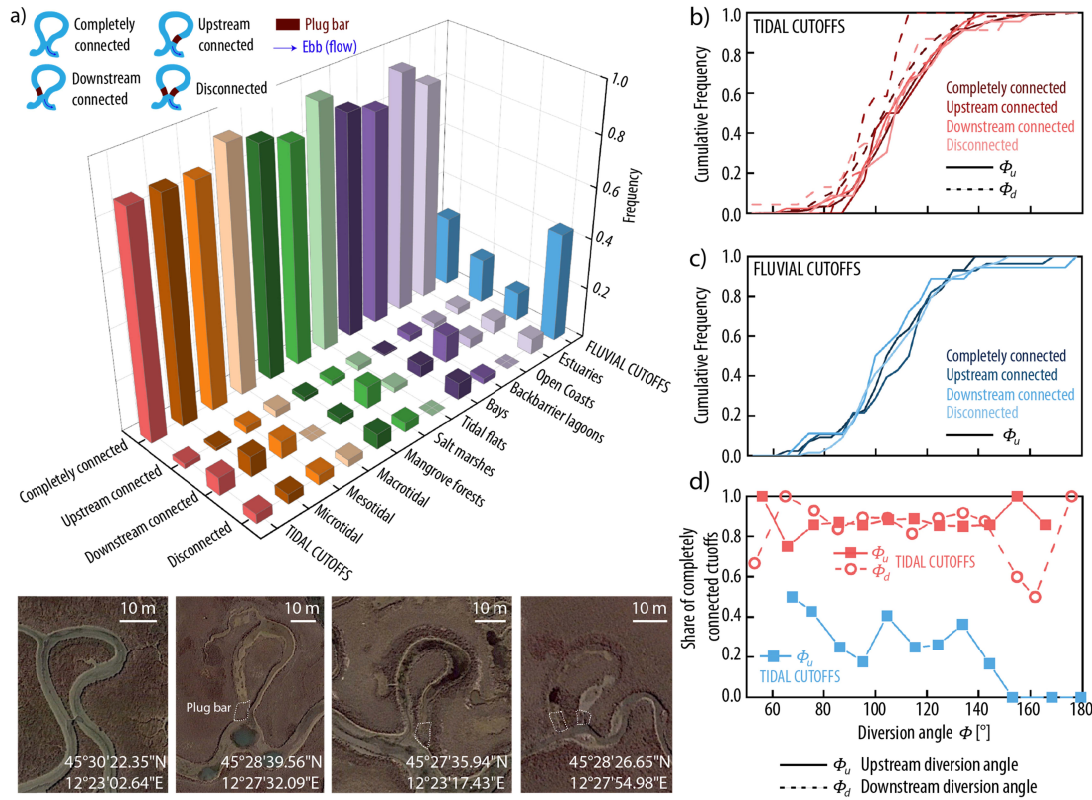


Figure 3. Dimensionless cutoff morphometrics. (a) Width-adjusted radius of curvature; (b) width-adjusted amplitude (c) width-adjusted intrinsic length; (d) sinuosity (χ); (e) asymmetry index (\mathcal{A}); (f,g)

595 upstream and downstream flow-diversion angles (Φ_u and Φ_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 (Φ) 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° 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.