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# Integrated analysis of flow, form, and function for river management and design testing

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#### 16 Abstract

17 The extent and timing of many river ecosystem functions is controlled by the interplay of 18 streamflow dynamics with the river corridor shape and structure. However, most river 19 management studies evaluate the role of either flow or form without regard to their dynamic 20 interactions. This study develops an integrated modeling approach to assess changes in 21 ecosystem functions resulting from different river flow and form configurations. Moreover, it 22 investigates the role of temporal variability in such flow-form-function tradeoffs. The use of 23 synthetic, archetypal channel forms in lieu of high-resolution topographic data reduces time 24 and financial requirements, overcomes site-specific topographic features, and allows for 25 evaluation of any morphological structure of interest. In an application to California's 26 Mediterranean-montane streams, the interacting roles of channel form, water year type, and 27 hydrologic impairment were evaluated across a suite of ecosystem functions related to 28 hydrogeomorphic processes and aquatic habitat. Channel form acted as the dominant control 29 on hydrogeomorphic processes, while water year type controlled salmonid habitat functions. 30 Streamflow alteration for hydropower increased redd dewatering risk and altered aquatic 31 habitat availability. Study results highlight critical tradeoffs in ecosystem function performance and emphasize the significance of spatiotemporal diversity of flow and form at 32 33 multiple scales for maintaining river ecosystem integrity. The proposed approach is broadly 34 applicable and extensible to other systems and ecosystem functions, where findings can be 35 used to inform river management and design testing.

36

## 37 **1 Introduction**

38 Rivers are complex, dynamic systems that support many natural ecosystem functions, 39 including hydrogeomorphic processes and the creation and maintenance of aquatic and riparian 40 habitat (Doyle et al. 2005). The extent and timing of these functions is largely controlled by the 41 interplay of *flow*, described by streamflow magnitude, timing, duration, frequency, and rate-of-42 change (Poff 1997), and form, described by the shape and composition of the river corridor (Small et al. 2008; Pasternack et al. 2008; Worthington et al. 2014; Wohl et al. 2015; Vanzo et 43 44 al. 2016). In spite of this complex interplay, most environmental river management studies 45 evaluate the role of either flow or form without regard for these interactions.

46 The few studies that have effectively examined flow - form interactions related to ecosystem 47 functions highlight the scientific and management value of such analyses. For instance, by 48 evaluating the potential for shallow water habitat in both the historic and current lower Missouri 49 River under alternative flow regimes, Jacobson and Galat (2006) informed restoration priorities 50 for the river. However, this and similar studies (Brown and Pasternack 2008; Price et al. 2013; 51 Gostner, Parasiewicz, et al. 2013) are site specific, limiting their applicability to the range of 52 flow and form settings exhibited by a given hydroscape, each combination supporting distinct 53 ecosystem functions. Vanzo et al. (2016) offer an exception in their evaluation of ecohydraulic 54 responses to hydropeaking over a spectrum of existing and proposed flows and forms. See

55 Supplementary materials section 1 for more details on the flow-form-function conceptual 56 framework.

57 Utilizing archetypal channel forms in lieu of detailed, site-specific datasets allows for the 58 evaluation of a larger range of flow-form settings with limited data and financial requirements. 59 An archetype refers to a simple example exhibiting typical qualities of a particular group without 60 the full local variability distinguishing members of the same group (Cullum et al. 2017). An 61 archetype-based analysis of the Yuba River, California, was employed by Escobar-Arias and 62 Pasternack (2011), who evaluated salmonid habitat conditions across archetypal 1D cross-63 sections. An emerging technique for synthesizing digital terrain models (DTMs) of river 64 corridors using mathematical functions (Brown et al. 2014) provides an opportunity to expand on 65 the work of Escobar-Arias and Pasternack (2011) to evaluate 2D hydraulic response across any 66 channel or floodplain morphology of interest without a major increase in data requirements. 67 The application of synthetic DTMs to the evaluation of river ecosystem performance 68 bypasses data constraints of previous studies through the ability to directly generate

69 representations of historic, existing, or proposed morphologies with user-defined geomorphic 70 attributes. Synthetic river corridors have been used to evaluate controls on riffle-pool salmonid 71 habitat quality and erosion potential as well as to test the occurrence of the hydrogeomorphic 72 mechanism of flow convergence routing across a range of archetypal morphologies (Brown et al. 73 2015), but have not yet been applied to the development of ecohydraulic design criteria. At the 74 rapid rate of river habitat change and biodiversity loss (Magilligan and Nislow 2005), the ability 75 to design and compare the ecohydraulic performance of distinct morphologies with relevance 76 beyond an individual study site to an entire watershed or region would offer a powerful tool to 77 support the design of functional large-scale river rehabilitation measures.

78

## 79 **1.1 Study objectives**

This study applied synthetic DTMs of archetypal river morphologies, developed at the 80 81 regional scale based on an existing channel classification, to the evaluation of regional flow-82 form-function linkages. The authors investigate the common notions of flow- and form-process 83 linkages, in which different flow regimes and morphologies are assumed to support distinct 84 hydrogeomorphic processes (Montgomery 1997; Poff 1997; Kasprak et al. 2016). The overall 85 goal of the study is to test whether archetypal combinations of flow and form attributes generate 86 quantifiable hydraulic patterns that support distinct ecosystem functions. The study objectives 87 are to (1) generate synthetic DTMs of distinct river corridor archetypes mindful of patterns of 88 topographic variability necessary to ecogeomorphic dynamics, (2) evaluate the spatiotemporal 89 patterns of depth and velocity across the archetypal DTMs from objective one, and (3) quantify 90 the performance of a suite of critical ecosystem functions across alternative flow-form scenarios. 91 The specific scientific questions addressed through these objectives are as follows: (i) Do

- 92 archetypal river corridor morphologies support distinct ecosystem functions or is more or
- 93 different local topographic variation within archetypes needed? (ii) What is the significance of
- subreach-scale topographic variability in river ecosystem functioning? (iii) What is the role of
- 95 water year type and hydrologic impairment? (iv) What ecosystem performance tradeoffs can be
- 96 identified with relevance for environmental water management?

## 97 **1.2** Case study setting: Mediterranean-montane rivers

98 Mediterranean-montane river systems, which exhibit cold wet winters and warm dry 99 summers, provide a useful setting for evaluating flow-form interactions because they exhibit

- significant variability to test system sensitivity to different drivers (Gasith and Resh 1999). In the
   California Sierra Nevada, USA, native aquatic and riparian species are adapted to the biotic
- stresses (e.g., reduced water quality in summer) and abiotic stresses (e.g., high shear stress
- 103 during winter floods) associated with the highly seasonal flow regimes that depend on flow-form
- 104 interactions. Salmonid eggs, for example, require sufficient inundation depths and intragravel
- 105 flows in certain channel locations during biologically significant periods to survive (USFWS
- 106 2010a). Sierra Nevada rivers have been highly altered by dams and reservoir operations for water
- 107 supply, flood control, and hydropower (Hanak et al. 2011), driving dramatic declines in native
- 108 aquatic populations (Yoshiyama et al. 1998; Moyle and Randall 1998; Yarnell et al. 2012). See
- 109 Supplementary materials section 1.2 for more details.

# 110 2 Methods

111 The methodology can be summarized by three steps (Figure 1). First, a set of synthetic river

- 112 corridor DTMs is generated to represent channel types from an existing channel classification
- 113 (Section 2.1) and a set of hydrologic scenarios is selected for evaluation (Section 2.2). Next, a
- 114 2D hydrodynamic model [SRH-2D (Lai 2008)] is used to simulate ecologically relevant
- 115 hydraulic parameters [ERHPs, *sensu* Vanzo et al. (2016)] for each flow-form scenario (Section
- 116 2.3, 2.4). Finally, spatiotemporal patterns in ERHPs are used to evaluate the performance or
- 117 occurrence of a suite of ecosystem functions (Section 2.5) under each scenario. Each of these
- 118 steps is described in depth in the following paragraphs.



Figure 1. Major steps used to quantify ecosystem function performance across archetypal channel forms
 and hydrologic scenarios, with step numbers associated with text above. Key inputs and outputs are
 bolded and modeling tools are blue parallelograms.

119

123 Specifically, selected streamflow time series (flows) and river corridor DTMs (forms) are 124 input to a 2D hydrodynamic model to produce a continuum of hydraulic rasters [i.e., depth (d), 125 velocity (v), shear stress  $(\tau)$ ] for a modeled river corridor at each modeled flow stage. For each 126 model run, a set of ERHP rasters [e.g., Shields stress  $(\tau_0^*)$ ), indices incorporating both depth and 127 velocity  $(d \cdot v)$ ] is calculated from hydraulic model raster outputs. Finally, spatial and temporal 128 statistics characterizing ERHP outputs are used first to evaluate model results in terms of depth 129 and velocity at baseflow (0.2 x bankfull), 50% exceedance, and bankfull flows, and then to 130 quantify the performance of distinct ecosystem functions. Bankfull discharge, defined as the flow 131 that just reaches the transition between the channel and its floodplain, was estimated from the 132 channel geometry as described in Section 2.4. Temporal dynamics are evaluated by integrating 133 ERHP spatial statistics over each hydrologic scenario (Parasiewicz 2007) such that not only the 134 magnitude but also the timing, duration, and frequency of ecosystem functions can be evaluated 135 depending on the particular temporal requirements. The resulting annual time series represent the 136 temporal pattern of the 2D hydraulic response in a specific DTM for a single hydrologic 137 scenario. This process is detailed in Supplementary materials section 2.

138 The experimental design involved a series of 16 hydraulic model runs under steady flow

139 conditions, simulating two river corridor morphologies across eight discharges spanning

140 baseflow (0.2x bankfull) to twice bankfull flow stages. The decision to evaluate proportions of

141 bankfull flow was driven by the established geomorphic and ecological significance of bankfull

142 flow in the literature (Wolman and Miller 1960; Doyle et al. 2005; Richter and Richter 2000).

143 Further, scaling flows by a common non-dimensional metric allows for readers worldwide to

evaluate these results relative to the setting in their locality. These eight discharges discretized

- the daily flow regimes evaluated to simplify temporal analysis. All simulated combinations were
- 146 designed to reproduce realistic archetypal flow and form conditions in Mediterranean-montane
- river systems for two channel types of interest, plane bed and pool-riffle (see section 2.1). A
- rigorous scaling approach to compare the full range of possible configurations was outside the
- scope of the current study. The following sections describe the flow regimes, river corridor
- 150 morphologies, hydraulic modeling approach, and ecosystem functions considered.

# 151 **2.1** Synthetic river corridor morphologies

152 Two archetypal river corridor morphologies distinguished in the Sacramento Basin channel 153 classification by Lane, Pasternack, et al. (2017) were considered in this study as a proof-of-154 concept: semi-confined plane bed and semi-confined pool-riffle. These morphologies were 155 selected for their common occurrence in mid-elevation montane environments (Montgomery and 156 Buffington 1997; Wohl and Merritt 2005) and their similar channel dimensions and slopes 157 contrasted by major differences in subreach-scale topographic variability. An existing field data 158 driven channel classification for the Sacramento Basin (Lane, Pasternack, et al. 2017) provided 159 the parameter values needed to synthesize the two archetypal morphologies, quantified as the 160 median field-surveyed values for each channel type.

- DTMs of the investigated channel types were generated using the synthetic rivers model developed by Brown et al. (2014). Below, we briefly provide the equations vital to understanding the DTMs created in this study. The goal of the design process was to capture the essential organized features of each channel type so that their functionalities can be evaluated in a reductionist approach without the random details of real river corridors that cause highly
- 166 localized effects.

# 167 2.1.1 Reach-average parameters

168 The synthetic rivers approach first creates a reach-averaged river corridor that is scaled by 169 reach-averaged bankfull width  $(w_{BF})$  and bankfull depth  $(h_{BF})$ , with median sediment size  $(D_{50})$ , 170 slope (*S*), sinuosity, and floodplain width and lateral slope as user-defined input variables. For 171 each synthetic river scenario, 140 longitudinal nodes were spaced at 1 m (~1/10 bankfull channel 172 widths).

# 173 2.1.2 Channel variability functions

174 Next, this approach incorporates subreach-scale (<10 channel widths frequency) topographic

variability because many hydrogeomorphic processes of ecological significance depend on

176 specific patterns of topographic variability and associated habitat heterogeneity (MacWilliams et

- al. 2006; Poff and Ward 1990; Scown et al. 2015). The local bankfull width at each location  $x_i$
- along the channel,  $w_{BF}(x_i)$ , is given by Eq. 1 as a function of reach-averaged bankfull width  $w_{BF}$

and a variability control function  $f(x_i)$ , with a similar equation used to characterize vertical bed undulation that incorporates *S*:

181 
$$w_{BF}(x_i) = w_{BF} * f(x_i) + w_{BF}$$
[1]

182 There are many available mathematical and statistical control functions that may be used to 183 describe archetypal river variability (Brown and Pasternack 2016). For this study, the variability 184 of  $w_{BF}$  and  $h_{BF}$  about the reach-averaged values was determined by a sinusoidal function, as

$$f(x_i) = a_s \sin(b_s x_r + h_s)$$

186 where  $a_s$ ,  $b_s$ , and  $h_s$  are the amplitude, angular frequency, and phase shift alignment parameters

[2]

187 for the sinusoidal component, respectively, and  $x_r$  is the Cartesian stationing in radians. The

188 Cartesian stationing was scaled by  $w_{BF}$  so that the actual distance was given by  $x_i = x_r * w_{BF}$ .

189 The sinusoidal function alignment parameters were adjusted iteratively to achieve desired values

190 for  $h_{BF}$ , width-to-depth ratio ( $w/h_{BF}$ ), sinuosity, and the coefficient of variation (CV) of  $w_{BF}$ 

and  $h_{BF}$  based on plane bed and pool-riffle channel classification archetypes (Lane, Pasternack,

et al. 2017). Floodplain confinement, the bankfull to floodplain width ratio, was used to set

193 valley width and overbank topography.

194 Because river classifications traditionally aim to capture the central tendency of river types at 195 the reach scale, they contain little to no information on subreach-scale topographic variability 196 and landform patterning (Lane, Pasternack, et al. 2017). This study used outputs from a channel 197 classification that was unique in including statistical characterization of subreach width and 198 depth variability using the metric of CV, based on the average and standard deviation of field-199 derived values (Lane, Pasternack, et al. 2017). However, there remained numerous landform 200 patterning permutations using the control function parameters of Eq. 2 that could yield those CV 201 values, many associated with profoundly different geomorphic processes. In these cases, field 202 experience and judgment informed the design of topographies capable of supporting the 203 dominant geomorphic processes of each channel type as outlined in the classification study. For 204 example, for the pool-riffle system, minimum depth and maximum width were made to

205 positively covary in the DTM to represent this patterning (Brown and Pasternack 2016).

206

#### 207 **2.2 Flow regimes**

Four hydrologic scenarios characteristic of the mixed snowmelt and rain flow regime typical of Mediterranean-montane systems were evaluated (Lane, Dahlke, et al. 2017): unimpaired wet, unimpaired dry, altered wet, and altered dry annual flow regimes (Figure 2). Daily streamflow time series for two mid-elevation gauge stations in the western Sierra Nevada, California, were chosen to represent these archetypal flow regimes under unimpaired (North Yuba River below Goodyears Bar) and altered (New Colgate Powerhouse) conditions (see Supplementary materials

- section 2.2 for map of gage locations). These gauges lie within similar physioclimatic and
- 215 geologic settings and provide daily streamflow time series for both an extremely wet (WY 2010;
- 216 >75<sup>th</sup> percentile annual streamflow) and an extremely dry (WY 2014; <25<sup>th</sup> percentile annual
- 217 streamflow) water year. The New Colgate Powerhouse gage captures typical hydropeaking
- 218 patterns of Sierra Nevada streams. The 50% exceedance flows for each hydrologic scenario are
- 219 23.3, 5.0, 19.2, and 18.5  $m^3/s$  for the wet unimpaired, dry unimpaired, wet altered, and dry
- altered scenarios, respectively.



Figure 2. Four hydrologic scenarios were considered: unimpaired wet, unimpaired dry, altered wet, and
altered dry. Graphs illustrate unimpaired and altered daily time series of (a) streamflow and (b)
discretized proportions of bankfull flow based on stage-discharge thresholds from Table 1.

225

#### 226 2.3 Hydraulic modeling

227 The surface-water modeling system (SMS; Aquaveo, LLC, Provo, UT) user interface and 228 Sedimentation and River Hydraulics- Two Dimensional (SRH-2D) algorithm (Lai 2008) were 229 used to produce exploratory hydrodynamic models for each flow-form scenario. SRH-2D is a 230 finite-volume numerical model that solves the Saint Venant equations for the spatial distribution 231 of water surface elevation, water depth, velocity, and bed shear stress at each computational 232 node. It can handle wetting/drying and supercritical flows, among other features. The parametric 233 eddy viscosity equation was used for turbulence closure. A coefficient value of 0.1 suitable for 234 shallow rivers with coarse bed sediment was used in that equation. A computational mesh with 235 internodal mesh spacing of 1 m (relative to a channel width of 10 m) was generated for each 236 synthetic DTM. Because this study was purely exploratory, using numerical models of 237 theoretical river archetypes, no calibration of bed roughness or eddy viscosity was possible. 238 Similarly, no validation of model results was possible.

239 The model required inputs of discharge and downstream flow stage as well as boundary

240 conditions of bed topography and roughness. Eight model runs for each morphology capture the

241 discharge range of 0.2 - 2.0 *x* bankfull flow stage (Table 1), where bankfull flow stage is the

242 water surface elevation at which flows spill onto the floodplain. The specific simulated discharge

- 243 values associated with these stages were estimated for each morphology using Manning's
- 244 equation based on representative cross-sections of the synthetic DTMs. Bankfull stage and
- 245 wetted perimeter were determined manually from the cross-sections, and cross-sectional area
- was calculated using the trapezoidal approximation. Manning's n was set at 0.04 to represent a
- 247 typical unvegetated gravel/cobble surface roughness (Abu-Aly et al. 2014).

248Table 1. Simulated channel archetype discharge values for 0.2 - 2.0 times bankfull flow stage

calculated from Manning's equation, and associated stage - discharge threshold estimates for the NorthYuba River.

	Simulated	d discharge	N. Yuba River discharge
Fraction of bankfull flow	Plane Bed (m3/s)	Pool-Riffle (m3/s)	Stage - discharge threshold (m3/s)
0.2	1.3	1.2	2.8
0.4	6.8	4.5	14.2
0.6	17.7	9.7	22.7
0.8	28.7	17.8	28.3
1.0	58.2	27.7	56.6
1.2	95.5	64.3	85.0
1.5	164.4	139.9	113.3
2.0	310.3	338.1	141.6

#### 251

#### 252 2.4 Hydrological scaling

253 Finally, in order to scale the real streamflow time series to the synthetic DTMs, stage -254 discharge relationships are needed to associate each of the eight flow stages simulated in the 255 hydraulic model (Table 1) with the actual discharge required to fill the North Fork Yuba river 256 channel to that flow stage. In the absence of local stage-discharge relationships, these thresholds 257 were instead estimated manually (Table 1, final column) with the aim of retaining archetypal 258 hydrologic characteristics of interest. Specifically, stage thresholds were set so, in the wet year, 259 the flow stage time series remained at or above bankfull during winter storms and throughout the 260 spring snowmelt recession while, in the dry year, flow stage rarely exceeded bankfull and spent 261 the majority of summer at base flow. The estimated stage - discharge thresholds were justified by 262 the ability of the discretized flow regimes (Figure 2b) to retain these hydrologic patterns 263 exhibited by the un-discretized flow regimes (Figure 2a). However, these thresholds are 264 estimates and should not be considered as ultimate targets to inform river management. A major 265 assumption of this approach is that the flow stage discretization captures all significant spatial hydraulic patterns in the river corridor relative to the functions under consideration in this study 266 (see Supplementary materials for more details). 267

#### 268 2.5 River ecosystem functions

269 Five Mediterranean-montane ecosystem functions were considered, associated with two 270 major components of river ecosystem integrity: hydrogeomorphic processes and aquatic habitat 271 (Table 2). The performance of these functions was tested based on the following criteria: (1) a 272 longitudinal shift in the location of peak shear stress at high flows from topographic highs to 273 topographic lows was used to test the occurrence of flow convergence routing, a dominant 274 geomorphic formation and maintenance process in certain channels (MacWilliams et al. 2006); 275 (2) a measure of hydrogeomorphic variability was used to quantify overall habitat heterogeneity 276 in the river corridor (Gostner, Alp, et al. 2013); and (3) fall-run Chinook salmon habitat was 277 evaluated with respect to (a) bed preparation and (b) bed occupation functions based on 278 established shear stress thresholds and biologically significant timing thresholds (Escobar-Arias 279 and Pasternack 2010) as well as (c) redd dewatering risk during bed occupation. These functions 280 were evaluated using a Python script that enabled rapid evaluation of model outputs over the 281 specific spatio-temporal constraints.

Table 1. The five river ecosystem functions evaluated in this study and their associated ecologically
 relevant hydraulic parameters (ERHPs), biologically relevant periods, and spatial extents.

Ecosystem Function	ERHP(s)	Biological Period	Spatial Extent	Citations
Hydrogeomorphic processes				
Flow convergence routing	shear stress		bankfull channel	MacWilliams et al. 2006
Hydrogeomorphic diversity	velocity, depth		river corridor	Gostner et al. 2013a
Aquatic habitat				
Salmonid bed preparation	shear stress	Oct – Mar	bankfull channel	Escobar-Arias and
Salmonid bed occupation	shear stress	Apr – Sep	bankfull channel	Pasternack 2010
Redd dewatering	velocity, depth	Oct – Mar	spawning channel	USFWS, 2010b

#### 284 2.5.1 Flow convergence routing mechanism

Flow convergence routing, the periodic reversal of peak shear stress location, is often considered critical to pool-riffle maintenance (White et al. 2010). The Caamaño criterion (Caamaño et al. 2009) was used to estimate the minimum riffle depth needed for a reversal to occur in each archetypal morphology. This mechanism was further evaluated based on the presence of a shift in peak shear stress from topographic wide-highs (riffles) to narrow-lows (pools), which indicates that the locations of scour and deposition are periodically shifted in the channel to maintain the relief between riffles and pools (Brown and Pasternack 2014).

#### 292 2.5.2 Hydrogeomorphic diversity

The hydro-morphological index of diversity (HMID) (Gostner, Alp, et al. 2013) was used to quantify overall physical heterogeneity of the river corridor as follows, where the coefficient of variation (CV) is the standard deviation of depth or velocity divided by its mean:

$$HMID_{reach} = (1 + CV_{\nu})^{2} * (1 + CV_{d})^{2}$$
[3]

Three tiers of HMID were delineated as follows: HMID <5 indicates simple uniform or channelized reaches; 5< HMID <9 indicates a transitional range from uniform to variable reaches; HMID >9 indicates morphologically complex reaches (Gostner, Parasiewicz, et al. 2013). To date, no studies have applied this index to archetypal terrains, so this is a novel application to further understand its value in quantifying ecosystem functions. Percent exceedance curves of HMID provided graphical representations of the temporal patterns of hydraulic diversity under alternative flow-form scenarios.

#### 304 2.5.3 Salmonid bed occupation and preparation

Ecosystem functions related to salmonid habitat can be split into bed occupation functions, which occur while the fish are directly interacting with the channel bed (i.e. spawning, incubation and emergence), and (2) bed preparation functions, which occur between occupation periods during migration (Escobar-Arias and Pasternack 2011). A stable bed, indicated by low shear stress ( $\tau_o^* < \tau_{c \ 50}$ ), is needed to minimize scour during bed occupation (Oct – Mar), while high shear stress capable of mobilizing the active layer ( $\tau_o^* > \tau_{c \ 50}$ ) is necessary to rejuvenate the sediment during bed preparation (Apr – Sep) (Soulsby et al. 2001; Konrad et al. 2002). See

312 Supplementary materials section 2.5.4 for more details.

Bed mobility transport stages delimited per-pixel by nondimensional boundary shear stress or Shields stress ( $\tau_o^*$ ) thresholds (Jackson et al. 2015) were used to quantify these bed occupation and preparation functions according to the following equation:

- 315 and preparation functions according to the following equation:
- 316

296

$$\tau_o^* = \frac{\tau_b}{g(\rho_s - \rho)D_{50}} \tag{4}$$

317 where  $\tau_b$  is bed shear stress:

318

$$\tau_b = \rho_w(u^*)^2 \tag{5}$$

based on water density  $(\rho_w)$  and shear velocity  $u^* = U\sqrt{C_d}$ , where U is depth-averaged velocity 319 for an individual pixel and  $C_d$  is the depth-based drag coefficient (Pasternack 2011).  $\tau_0^*$  therefore 320 321 varies spatially and with discharge as a function of depth and velocity. For the present 322 application, a stable bed is assumed when  $\tau_o^* < 0.01$ , intermittent transport when  $0.01 < \tau_o^* < 0.03$ , 323 partial transport when  $0.03 < \tau_o < 0.06$  and full mobility when  $0.06 < \tau_o^* < 0.10$  (Buffington and 324 Montgomery 1997). At each discharge, the areal proportion of each bed mobility tier occurring in the river corridor region of interest can be calculated. Function performance is then quantified 325 326 through time as the cumulative proportion of the channel providing functional bed mobility 327 conditions during biologically relevant periods. Results are then binned such that low, mid, and 328 high performances are associated with 0-25%, 25-75%, and 75-100% performance. For example, 329 at least 75% of the channel area must exhibit partial or full bed mobility on average over the bed 330 preparation period to achieve high bed preparation performance.

#### 331 2.5.4 Redd dewatering

332 Salmonid redd dewatering is a major concern in Sierra Nevada streams managed for 333 hydropower (USFWS 2010b). Reductions in flow stage exposing the tailspill and reductions in 334 velocity diminishing intragravel flow through the redd can dramatically reduce the survival of 335 salmonid eggs and pre-emergent fry (Healey 1991; USFWS 2010b). This study focused on fall-336 run Chinook salmon (Oncorhynchus tshawytscha) as an important species in Sierra Nevada 337 streams. Redd dewatering risk was measured as the areal proportion of viable spawning habitat 338 with depth below 0.15 m and/or velocity below 0.09 m/s during the occupation period 339 [incubation and emergence period (Dec – Mar) (USFWS 2010b)]. Viable spawning habitat was 340 defined according to USFWS as the portion of the bankfull channel with velocity from 0.1 - 1.6 m/s and depth from 0.15 - 1.3 m at 0.4x bankfull stage, the most common stage experienced 341 342 under unimpaired conditions during the spawning period (Oct – Dec) (USFWS 2010a).

#### 343 **2.6 Holistic ecosystem functions analysis**

344 There is limited guidance available in the literature regarding how best to evaluate and 345 visually represent the environmental performance of rivers across a suite of ecosystem functions. 346 The question of how best to evaluate environmental performance using individual metrics or 347 functions is well established (CITE). The broader water resources management literature offers 348 several summary performance indices that could be applied to this new setting. For example, the 349 water resources management sustainability index (Sandoval-Solis et al. 2010), which evaluates 350 different water management policies by summarizing across metrics of reliability, resilience, and 351 vulnerability, could be applied as a summary index of environmental performance. In this study, 352 we created a graphic that aligns all of the ecosystem functions in one table to visualize the 353 performance of a suite of temporally varying functions simultaneously.

## 354 **3 Results**

The synthetic DTM results are presented first (study objective 1). Then the hydraulic modeling results are discussed in terms of depth and velocity patterns (study objective 2). Finally, model results are used to interpret the performance of five ecosystem functions (Table 2) across alternative flow-form scenarios (study objective 3).

#### 359 3.1 Synthetic digital terrain models

Two 140 m long synthetic DTMs were generated representing archetypal morphological configurations of semi-confined pool-riffle and plane bed morphologies (Figure 3). These DTMs exhibited distinct reach-averaged attributes (e.g., S,  $w/h_{BF}$ , and  $D_{50}$ ) (Table 3a), subreach-scale topographic variability (e.g., CV), and proportions of the river corridor exhibiting positive and negative geomorphic covariance structures (GCSs) (Table 3b). The sinusoidal function alignment parameters used are listed in Table 3b. The resulting morphologies exhibited major differences in subreach-scale topographic variability as illustrated by the planform and longitudinal topographic patterns in Figure 3. The bankfull channel area was 868  $m^2$  in the poolriffle and 1,041  $m^2$  in the plane bed DTM.

	(a) Geomorphic Attributes			(b) Alignment Parameters			
	Plane Bed	Pool - Riffle		Plane Bed	Pool - Riffle		
Channel			Planform				
w <sub>BF</sub> (m)	10	10	phase shift	0	0		
h <sub>BF</sub> (m)	1	1	amplitude	0.8	0		
S (%)	1	2	frequency	2	2		
w/h <sub>BF</sub>	10	10	Bankfull Widt	h			
D <sub>50</sub> (m)	0.2	0.1	phase shift	$\pi/2$	π		
sinuosity	1.1	1.1	amplitude	0.01	0.5		
CVw <sub>BF</sub>	0.01	0.35	frequency	2	3		
$\mathrm{CVH}_{\mathrm{BF}}$	0.03	0.18	<b>Bed Elevation</b>				
+ GCS (%)	55	86	phase shift	0	2.7		
- GCS (%)	45	14	amplitude	0.04	0.35		
Floodplain			frequency	2	3		
confinement	0.5	0.5	Floodplain Ou	tline			
lateral slope	0.80%	0.80%	phase shift	0	0		
width (m)	16	16	amplitude	0	0.5		
D <sub>50</sub> (m)	0.03	0.03	frequency	1.5	1.5		
	Č						

Table 2. (a) Channel and floodplain geomorphic attributes and (b) control function alignment
 parameters used in the design of synthetic DTMs of plane bed and pool-riffle channel morphologies



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Figure 3. Two archetypal river corridor morphologies were evaluated, plane bed (A - D) and pool-riffle (E
- H). The synthetic DTMs (B and G) are shown overlaid by their bankfull channel boundaries, where dark
red is high elevation and dark blue is low elevation, followed by their longitudinal profiles (C and H).

#### 375 **3.2** Spatial and temporal distribution of hydraulic variables

376 Depth and velocity values fell within typical ranges for gravel-bed montane streams across 377 base, 50% exceedance, and bankfull flows, supporting the archetypal specifications used in this 378 study (Richards 1976; Jowett 1993). Water depths ranged from 0.0 to 2.4 m, with higher average 379 depths in the plane bed than the pool-riffle across all three flows. The pool-riffle had lower 380 minimum and higher maximum depths across all flow levels, resulting in a larger depth range 381 and CV. Flow velocities ranged from 0.0 to 5.5 m/s, exhibiting a similar pattern to depth between 382 morphologies, with higher average and minimum velocities in the plane bed across all 8 383 discharge stages. In contrast with depth, however, at bankfull flow maximum velocity was 384 significantly higher in the plane bed than the pool-riffle, resulting in a higher velocity CV. The

HMID was substantially higher at base flow than higher flows, and was more than twice as highin the pool-riffle as in the plane bed at base flow.

387 Time series plots of hydraulic summary statistics illustrate the daily temporal variability of

depth and velocity over the four hydrologic scenarios (Figure 4). A reversal in the maximum CV

of velocity from the pool-riffle to the plane bed is evident during spring in the wet unimpaired

390 scenario and during summer in the wet altered scenario, corresponding with a very high

391 maximum velocity in the plane bed (5.5 m/s). The remainder of seasons and water year types

exhibit higher hydraulic variability in the pool-riffle, with the largest differences in CV occurring

at low flows.



394

Figure 4. Annual time series plots of maximum, average, and minimum (a) flow velocity and (b) waterdepth in plane bed and pool-riffle morphologies over four hydrologic scenarios.

397 Water depth was more sensitive to low flow variations in terms of rate of change, while 398 velocity was more sensitive to changes in high flows. This likely occurs because, in parabolic 399 channel geometries, the channel fills rapidly from low to bankfull flow, whereas, once the 400 bankfull channel is overtopped, a larger flow increase is required to engender the same increase in water depth over the wider floodplain so high flow changes translate more directly to velocity. 401 402 With regards to channel type, the pool-riffle morphology demonstrated an approximately linear 403 increase in depth with flow, while the plane bed morphology demonstrated a rapid increase in 404 depth from low flow to 0.8x bankfull and a reduced rate of increase at higher flows. Conversely, 405 velocity in both morphologies increased at a slow linear rate from low flow to 0.8x bankfull flow 406 and then increased much faster in the plane bed at higher flows. Only at high flows (>1.5x 407 bankfull) did pool-riffle velocity exhibit a strong sensitivity to flow variability. These findings demonstrate that changes in the hydraulic environment due to variations in discharge were 408

- 409 stronger in the plane bed than the pool-riffle, indicating that pool-riffle hydraulics are less
- 410 sensitive to changes in flow on average but instead exhibit more complex spatial patterns.

#### 411 **3.3 Ecosystem function performance results**

- 412 All six Mediterranean-montane river ecosystem functions were found to be controlled by
- both flow and form attributes to varying extents, as illustrated in Figure 5 for the unimpaired
- 414 flow regime and in Figure 6 for the altered flow regime.



415

Figure 5. Summary of temporally varying ecosystem function performance under an unimpaired flow regime across four flow-form scenarios: wet – pool-riffle, wet – plane bed, dry – pool-riffle, dry – plane bed. The five ecosystem functions evaluated are: 1. Flow convergence routing (VR), 2. hydrogeomorphic diversity (HG), 3. redd dewatering risk (RD), 4. salmonid bed preparation (BP), and 5. salmonid bed occupation (BO). Tiered performance is indicated in the key by increasingly dark shading and bimodal performance (VR and RD) is either colored or empty. Greyed regions indicate periods of the year that functions are not biologically relevant. Base flow = 0.2x, bankfull flow = 1.0x, and flood flow = 1.5x bankfull flow as defined in Table 1.





Figure 6. Summary of temporally varying ecosystem function performance under an altered flow regime across four flow-form scenarios: wet – pool-riffle, wet – plane bed, dry – pool-riffle, dry – plane bed (key in Fig. 5).

#### 426 3.3.1 Flow convergence routing

427 The pool-riffle morphology demonstrated a shear stress reversal from low to high flow, as 428 indicated by a Caamaño criterion riffle depth threshold for reversal of 0.21 m (approximately 429 0.4x bankfull stage) and a shift in the location of peak shear stress from the riffle crest to the pool 430 trough from base to bankfull flow (see Supplementary materials for more details). The existence 431 of a dominant flow convergence routing mechanism is further indicated by 86% of the pool-riffle 432 morphology exhibiting a positive GCS (i.e., primarily wide shallow riffles and narrow deep 433 pools). Alternatively, the plane bed morphology did not exhibit a shear stress reversal based on 434 either the Caamaño criterion or a peak shear stress location shift, and only 55% of the river XV 435 corridor exhibited positive GCS.

#### 436 3.3.2 Hydrogeomorphic diversity (HMID)

437 HMID was higher in the pool-riffle than the plane bed morphology at flows up to 1.2x438 bankfull, beyond which they were nearly equivalent. That is, for a given hydrologic scenario, the 439 cumulative HMID over the year was higher in the pool-riffle. The highest index values and the 440 greatest difference between the two morphologies occurred at the lowest flow stage (0.2x 441 bankfull discharge), when HMID was twice as high in the pool-riffle. The rapid decrease in 442 HMID for in-channel flows in both morphologies with increasing discharge illustrates the limited 443 temporal persistence of high diversity hydraulic habitats in all but the lowest flow conditions. In 444 natural conditions, once flows spill over the banks, there should be a significant increase in 445 HMID as topographically heterogeneous floodplains inundate. However, in the absence of 446 detailed floodplain attributes from the channel classification, this study considered simple 447 floodplain morphologies in both archetypes.

448 HMID exceedance curves for each of the eight flow-form scenarios provided insight into 449 hydraulic diversity patterns (Figure 7). As low flows produce higher HMID values in general, it 450 is unsurprising that in a very dry year both morphologies exhibited high HMID for most of the 451 year. Under dry conditions, the unimpaired flow regime provided nearly twice as many days with 452 high HMID in both morphologies. Under the altered flow regime, HMID was slightly higher in 453 the wet pool-riffle than the dry plane bed for all flows above 17% exceedance. The highest 454 HMID was exhibited by the pool-riffle under dry unimpaired conditions (HMID=5.9), 455 presumably due to the combination of high topographic variability and extended summer low 456 flows. At the 50% exceedance flows of each hydrologic scenario, hydraulic diversity was more 457 sensitive to water year type than hydrologic alteration, and appeared to be most controlled by 458 channel morphology. Alternatively, at the 10% exceedance flows, water year type played a more 459 significant role, with the dry year exhibiting much higher HMID across both morphologies and 460 impairment conditions. More significantly, the temporal analysis of HMID revealed that, unlike 461 the dry altered flow regime, the dry unimpaired flow regime exhibited high HMID during the 462 fall-run Chinook bed occupation period, as detailed in the Supplementary materials.



463

Figure 7. Hydromorphic index of diversity (HMID) exceedance curves for (a) unimpaired and (b) altered flow regimes under different channel morphologies and water year types.

#### 466 3.3.3 Salmonid bed preparation and occupation

467 Significant differences in salmonid habitat performance across flow-form scenarios were 468 identified from shear stress based sediment mobility patterns (Figure 8). Under unimpaired 469 conditions, the wet year exhibited high bed preparation performance and low bed occupation 470 performance, while the dry year exhibited mid performance in both functions with reduced bed 471 preparation but increased bed occupation performance. Under streamflow alteration, bed 472 preparation performed well across water years while bed occupation performed poorly across 473 water years and morphologies due to increased sediment mobility under elevated low flows 474 during the occupation period. Spatially, in the pool-riffle channel, higher sediment mobility 475 occurred over the riffle crests while the pools remained less mobile at all but flood flows. 476 Conversely, sediment mobility was nearly uniform in the plane bed channel across all flows.

477



Figure 8. Daily time series plots of the proportion of the bankfull channel exhibiting different tiers of
sediment mobility illustrate the performance of salmonid bed preparation (boxed, partial/high mobility
from Apr-Sep) and occupation (no/low mobility from Oct-Mar) functions.

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478

#### 483 3.3.4 Redd dewatering

Viable spawning habitat, based on depth and velocity requirements, varied between the channel forms. Nearly 50% of the bankfull channel provided viable spawning habitat in the poolriffle compared to only 31% in the plane bed. Pool-riffle spawning habitat was extensive and patchy, excluding only excessively high velocity zones on the riffle crests. Alternatively, plane bed spawning habitat only occurred in 1-2 meter bands along the wetted channel margins with sufficiently low velocity to meet predefined spawning requirements.

Redd dewatering risk within viable spawning habitat areas also varied significantly across
flow-form scenarios. Redd dewatering risk was greater in the plane bed than the pool-riffle at
base flow (100% vs 57% of spawning habitat) but risk was maintained across a greater range of
flows (0.2 - 0.4x bankfull flow) in the pool-riffle. This is because the pool-riffle morphology has
more gradual side slopes and the total available spawning habitat is greater. High dewatering risk
(>30% of spawning habitat) occurred only in the dry altered scenario, with very low flows
occurring throughout the redd incubation (Oct – Dec) and emergence (Jan – Mar) periods.

497

# 498 **4 Discussion**

# 499 **4.1 The utility of synthetic river archetypes**

This study demonstrated the ability to synthesize DTMs from channel classification
 archetypes exhibiting distinct ecosystem function performance, offering a scientifically
 transparent, repeatable, and adjustable framework for flow-form-function inquiry. Specific

503 geomorphic attribute values were accurately represented by the synthetic morphologies,

- 504 including channel dimensions, cross-sectional geometry, depth and width variability, sinuosity,
- and slope (Fig. 3). The flow convergence routing mechanism was shown to occur in the pool-
- 506 riffle archetype but not in the plane bed one, confirming that the two morphologies were
- 507 capturing distinct geomorphic maintenance processes as distinguished by the Sacramento Basin
- 508 channel classification (Lane, Pasternack, et al. 2017). Once a person understands how to produce 509 a synthetic DTM with the required subreach-scale variability to drive geomorphic processes and
- 50 ecological functions, then the software implements that very quickly. As a result, time and
- 511 financial requirements are dramatically reduced compared to doing a field-based campaign
- 512 involving meter-resolution topographic surveying, parameter calibration, and quality assurance
- 513 procedures. This approach therefore liberates future research to explore and isolate a larger range
- 514 of flow and form characteristics than those considered in the present study. This is an important
- 515 first step for evaluating how different kinds of rivers function, but care should be used in
- 516 extrapolating the specific results to any specific actual channel topography for real river
- 517 management. Archetypal studies provide useful guidelines, and then local studies should be
- 518 conducted to ascertain how the mechanisms play out in their details in the local setting, possibly
- 519 including validation efforts, to the extent feasible.

520 To choose the correct permutation of depth and width parameters to the synthetic 521 morphologies, expert judgment was used based on field experience and understanding of how to 522 interpret the processes associated with different patterns of topographic variability. However, 523 some attributes required to generate representative topographic attributes, such as floodplain 524 width variability and floodplain lateral slope, were not available in the channel classification of 525 Lane, Pasternack, et al. (2017). This represents an important limitation of the proposed method, 526 because useful results for certain ecosystem functions (e.g., riparian recruitment) require better 527 information than is currently available. More datasets focusing on different aspects of 528 geomorphic variability at different scales would enable more informed metric and parameter

529 choices (Brown and Pasternack 2017).

# 530 **4.2** Ecological significance of specific patterns of topographic variability

531 The spatial and temporal distributions of depth and velocity across channel forms illustrate 532 differences in sensitivity to flow changes, with major implications for ecosystem functioning and 533 aquatic biodiversity (Dyer and Thoms 2006). The pool-riffle morphology was less sensitive to 534 temporal changes in flow in terms of associated changes in depth and velocity, but more spatially 535 variable, exhibiting a larger range and CV of depth and velocity values for a given discharge. 536 This indicates that the pool-riffle has more sustained persistence of hydraulic patterns, making 537 many ecosystem functions less prone to temporal fluctuations with flow as long as the discharges 538 fall below the threshold for particular processes (Gostner, Parasiewicz, et al. 2013).

539 Study results support emerging scientific understanding that many river ecosystem functions 540 are controlled by subreach-scale topographic variability (White et al. 2010; Brown and 541 Pasternack 2016; Thompson 1986; Murray et al. 2006) by quantifying the occurrence of distinct

- 542 ecosystem functions in reaches of high versus low topographic variability. Specifically, results
- 543 emphasize that it is not enough to just obtain random topographic variability or any arbitrary
- 544 coherent permutation of variability, but rather the pattern of organized variability must meet the
- 545 requirements of the appropriate GCS and dominant geomorphic processes for that channel
- 546 archetype (Brown and Pasternack 2014; Brown et al. 2015).

547 Distinct spatial and temporal hydraulic patterns identified in this study but not explicitly 548 incorporated into performance metrics highlight important future directions for this 549 methodology. For example, changes in spatial patterns of sediment mobility exhibited across 550 flow-form combinations likely influence biological suitability for bed occupation in addition to 551 the magnitude- and timing-based performance metrics considered here. The temporal patterns of 552 bed mobility also varied substantially within the bed occupation and preparation periods (Fig. 7), 553 which was not captured by the selected performance metrics. More information about the spatio-554 temporal hydraulic requirements for particular species and life-stages and improved metrics for 555 quantifying these characteristics would refine performance estimates within the proposed

556 framework.

## 557 **4.3** Flow and form controls on ecosystem functioning

558 Five Mediterranean-montane river ecosystem functions related to geomorphic variability and 559 aquatic habitat were evaluated in the context of interacting flow (i.e., water year type and 560 hydrologic impairment) and form (i.e., morphology type) controls on ecohydraulic response 561 (Figure 5 and 6). Flow convergence routing was controlled primarily by channel form, as it only 562 occurred in the pool-riffle morphology. However, sufficiently high flows were also needed for a 563 shear stress reversal to occur in support of the mechanism. Hydrogeomorphic diversity was 564 controlled primarily by channel form, and specifically topographic variability, as expected. More 565 surprisingly, HMID was also influenced by flow attributes, with water year type, hydrologic 566 impairment, and morphology type all playing significant and interacting roles in the ecohydraulic 567 response. Salmonid bed preparation and occupation illustrate trade-offs in all three controlling 568 variables, with bed preparation performing best in the wet, altered, plane bed scenario while bed 569 occupation performed best in the dry, unimpaired pool-riffle morphology. The duration and 570 timing of redd dewatering risk were controlled by water year type and hydrologic impairment, 571 while the magnitude of dewatering risk, based on the proportion of spawning habitat exhibiting 572 sufficiently low depth or velocity, was controlled solely by channel form. These results 573 emphasize the complex interacting flow and form controls on key ecosystem functions and the 574 differences in dominant controls between ecosystem functions.

575 HMID performance tradeoffs in particular provide insight for environmental water
576 management, given the common conception that increased habitat heterogeneity promotes
577 biodiversity (Dyer and Thoms 2006). The highest HMID was exhibited by the pool-riffle under

578 dry unimpaired conditions. However, under hydrologic impairment, HMID was higher under the

- 579 wet pool-riffle than the dry plane bed scenario for all but the lowest flows. This finding indicates
- 580 a tradeoff between flow and form with respect to diversity whereby either increasing topographic
- variability (i.e., plane bed to pool-riffle) or increasing the number of low flow days in the flow
- regime (i.e., wet to dry water year type) was capable of increasing overall spatiotemporal
- 583 diversity. In such instances, knowledge of flow-form interactions could be used to guide more
- nuanced, targeted management efforts to promote ecological end goals such as increased
- 585 biodiversity.

In general, bed occupation performed poorly across all flow and form scenarios. This finding may be due to the coarse bankfull stage discretization used in the study (eight discharges from 0.2 - 2x bankfull stage, Table 1), allowing lower daily discharge values to be associated with higher sediment mobility than occurs in reality. Results such as these can inform future studies by promoting iterative modification of decisions such as the bankfull stage discretization and the range of discharges considered to improve representation of ecosystem functions within the proposed methodology.

592 proposed methodology.

#### 593 **4.4 Implications for environmental management**

594 The quantitative metrics of relative performance across a suite of ecosystem functions 595 highlighted critical performance tradeoffs, emphasizing the significance of spatiotemporal 596 diversity of flow and form at multiple scales for maintaining river ecosystem integrity. For 597 example, the pool-riffle morphology supported flow convergence routing and promoted high 598 hydraulic diversity and salmonid bed occupation, while the plane bed morphology supported 599 salmonid bed preparation and provided habitats of reduced dewatering stress for salmonid redds 600 during dry years. These results indicate that restoring or designing a pool-riffle dominated stream 601 network to provide interspersed plane bed reaches may support higher overall ecosystem 602 integrity by promoting distinct and complementary functions in different locations during 603 biologically significant periods. Such findings support the emerging recognition of spatial and 604 temporal heterogeneity as fundamental characteristics of fluvial systems and the need for a 605 flexible framework within which natural processes, such as sediment transport and nutrient 606 dynamics, can occur (Clarke et al. 2003; Gostner, Parasiewicz, et al. 2013; Vanzo et al. 2016; 607 Escobar-Arias and Pasternack 2010).

608 With respect to hydrologic variability, only wet years supported high performance of 609 salmonid bed preparation and shear stress reversals, while dry years significantly increased 610 hydraulic diversity and availability of fall-run Chinook spawning habitat. A range of wet to dry 611 years is required to support the full suite of ecosystem functions considered here. Inter-annual 612 variability plays a key role (in concert with spatial variability of form and bed substrate) in 613 maintaining river ecosystem integrity. This finding also indicates the potential for changes or 614 losses in function under a changing climate in which the spectrum or the ratio of wet to dry years 615 is significantly altered from that to which native riverine species are adapted (Null and Viers 616 2013). For example, fewer sufficiently wet years to generate shear stress reversals in pool-riffle

617 reaches may compromise their ability to maintain high topographic variability, thus shifting the

618 suite of ecosystem functions supported in these reaches towards those already supported by plane

- 619 bed reaches. This would reduce ecological variability and thus overall ecological resilience of the
- 620 stream network.

621 This application of synthetic datasets to flow-form-function inquiry provides a foundation for 622 transitioning from expressing ecosystem impacts and responses in terms of fixed flow or form 623 features to spatiotemporally varying hydrogeomorphic dynamics along a spectrum of alterations 624 of the synthetic datasets. The simple, process-based framework proposed here is expected to 625 elucidate key processes and thresholds underlying spatial and temporal dynamics of river 626 ecosystems through future applications. For instance, the functional role and alteration thresholds 627 of individual geomorphic attributes (e.g., confinement, channel bed undulations) could be 628 isolated through iterative generation and evaluation of numerous synthetic channel forms. This 629 information is expected to improve understanding of ecosystem resilience and the potential for 630 rehabilitation projects under current and future hydrogeomorphic alterations.

#### 631 **4.5 Study uncertainty**

632 Uncertainties in the ecosystem functions model developed here include uncertainty in model 633 completeness, parameters, and data inputs. With respect to model completeness, this study 634 explicitly incorporated attributes of key hydrologic and geomorphic processes controlling river 635 ecosystem functions for more complete evaluation of controlling variables and their dynamic 636 interactions. However, several critical aspects of river ecosystems including water quality, 637 temperature, population dynamics, and morphodynamics are not considered in the scope of the 638 current study.

Model parameter uncertainties derive from parameter and equation selection. For example, 639 640 the depth slope product shear stress equation assumes steady uniform flow, which is appropriate 641 for the geomorphic archetypes considered here under steady discharges but should be assessed 642 on a case-by-case basis for application to real channel morphologies (Brown and Pasternack 643 2008; Pasternack et al. 2008). The use of Shields parameter thresholds to delimit sediment 644 transport stages provided a simple approach to explore flow - hydrogeomorphic process 645 relationships, but there is uncertainty associated with these thresholds and others could be 646 selected depending on the application or with more information regarding bed composition. The 647 spatial and temporal thresholds of ERHPs constraining the ecosystem functions are also 648 uncertain. For instance, the requirement of seven consecutive days of flooding for riparian 649 recruitment is an estimate based on field studies across the Sierra Nevada that exhibit high 650 variability between sites.

Data input uncertainties originate from the streamflow time series and river corridor
 morphologies. In the current application, stage-discharge relationships were the main source of
 hydrologic uncertainty, as they were manually estimated for the Yuba River in the absence of
 established rating curves. Rating curves derived from field measurements would substantially
 reduce this source of uncertainty. The use of real streamflow time series minimized uncertainty

associate with hydrologic inputs. However, the use of modeled streamflow or hydrologic

archetypes, as proposed for future applications, would create additional uncertainty.

658 Uncertainties arising from the use of synthetic river valleys morphologies include field

659 measurements of reach-averaged geomorphic attributes including the CV of width and depth.

660 The frequency and distribution of width and depth measurements used in these calculations will

influence variability estimates, and as a result, the synthesized topographies. More research is

662 needed to evaluate the influence of different sampling schemes and measures of topographic

variability on the synthesized DTMs and dependent hydrogeomorphic processes.

664

# 665 **5 Conclusions**

666 This study tackles key questions regarding the utility of synthetic DTMs for ecohydraulic 667 analysis, the ecological significance of topographic variability, how to evaluate the ecological impacts of different flow-form settings or types of river restoration efforts, and whether 668 (re)instatement of key flow or form attributes is likely to restore ecological processes (Council 669 670 2007). The development and application of simple, quantitative ecosystem performance metrics 671 enabled evaluation of the ecohydraulic response to changes in flow and/or form settings typical 672 of Mediterranean-montane rivers. By comparing these performance metrics across individual and 673 combined adjustments to flow and form attributes, this study provides a novel framework for 674 assessing and comparing ecosystem function performance under natural and human altered flow 675 regimes and river corridor morphologies. Moreover, this research demonstrates the significance 676 of spatiotemporal diversity of *flow* (seasonal and inter-annual) and *form* (channel shape and bed 677 substrate) and their interactions for supporting distinct ecosystem *functions* that maintain river 678 ecosystem integrity.

679

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#### 690 7 Citations

691

- Abu-Aly, TR, Gregory B. Pasternack, Joshua R. Wyrick, R Barker, D Massa, and T Johnson. 2014. 'Effects
   of LiDAR-derived, spatially distributed vegetation roughness on two-dimensional hydraulics in a
   gravel-cobble river at flows of 0.2 to 20 times bankfull', Geomorphology, 206: 468-82.
- Brown, Rocko A., and Gregory B. Pasternack. 2008. 'Engineered channel controls limiting spawning
   habitat rehabilitation success on regulated gravel-bed rivers', Geomorphology, 97: 631-54.
- Brown, Rocko A., and Gregory B. Pasternack. 2014. 'Hydrologic and topographic variability modulate
   channel change in mountain rivers', Journal of Hydrology, 510: 551-64.
- Brown, Rocko A., and Gregory B. Pasternack. 2016. 'Analyzing bed and width oscillations in a self maintained gravel-cobble bedded river using geomorphic covariance structures', Earth Surface
   Dynamics Discussions: 1-48.
- Brown, Rocko A., and Gregory B. Pasternack. 2017. 'Bed and width oscillations form coherent patterns in
   a partially confined, regulated gravel–cobble-bedded river adjusting to anthropogenic disturbances',
   Earth Surf. Dynam., 5: 1-20.
- Brown, Rocko A., Gregory B. Pasternack, and T. Lin. 2015. 'The Topographic Design of River Channels
   for Form-Process Linkages', Environmental Management, 57: 929-42.
- Brown, Rocko A., Gregory B. Pasternack, and W. W. Wallender. 2014. 'Synthetic river valleys: Creating
   prescribed topography for form–process inquiry and river rehabilitation design', Geomorphology, 214:
   40-55.
- Buffington, John M, and David R Montgomery. 1997. 'A systematic analysis of eight decades of incipient
   motion studies, with special reference to gravel-bedded rivers', Water Resources Research, 33: 1993 2029.
- Caamaño, Diego, Peter Goodwin, John M. Buffington, Jim C. Liou, and Steve Daley-Laursen. 2009.
  'Unifying criterion for the velocity reversal hypothesis in gravel-bed rivers', Journal of Hydraulic
  Engineering, 135: 66-70.
- Clarke, Stewart J., Lydia Bruce-Burgess, and Geraldene Wharton. 2003. 'Linking form and function:
   towards an eco-hydromorphic approach to sustainable river restoration', Aquatic Conservation: Marine
   and Freshwater Ecosystems, 13: 439-50.
- Council, National Research. 2007. "River science at the U.S. Geological Survey." In, edited by Committee
   on River Science at the U.S. Geological Survey, 206. Washington, D.C.
- Cullum, Carola, Gary Brierley, George LW Perry, and Ed TF Witkowski. 2017. 'Landscape archetypes for
   ecological classification and mapping', Progress in Physical Geography, 41: 95-123.
- Doyle, Martin W., Emily H. Stanley, David L. Strayer, Robert B. Jacobson, and John C. Schmidt. 2005.
   'Effective discharge analysis of ecological processes in streams', Water Resources Research, 41.
- Dyer, Fiona J., and Martin C. Thoms. 2006. 'Managing river flows for hydraulic diversity: an example of
   an upland regulated gravel-bed river', River Research and Applications, 22: 257-67.

- Escobar-Arias, M. I., and Gregory B. Pasternack. 2010. 'A hydrogeomorphic dynamics approach to assess
   in-stream ecological functionality using the functional flows model, part 1-model characteristics', River
   Research and Applications, 26: 1103-28.
- Escobar-Arias, M. I., and Gregory B. Pasternack. 2011. 'Differences in river ecological functions due to
   rapid channel alteration processes in two California rivers using the functional flows model, part 2 model applications', River Research and Applications, 27: 1-22.
- Gasith, A., and B. Resh. 1999. 'Streams in Mediterranean Regions: Abiotic Influences and Biotic Responses
   to Predictable Seasonal Event', Annual Review of Ecological Systems.
- Gostner, Walter, Maria Alp, Anton J. Schleiss, and Christopher T. Robinson. 2013. 'The hydro morphological index of diversity: a tool for describing habitat heterogeneity in river engineering
   projects', Hydrobiologia, 712: 43-60.
- Gostner, Walter, Peter Parasiewicz, and Anton J. Schleiss. 2013. 'A case study on spatial and temporal
   hydraulic variability in an alpine gravel-bed stream based on the hydromorphological index of
   diversity', Ecohydrology, 6: 652-67.
- Hanak, Ellen, Jay Lund, Ariel Dinar, Brian Gray, Richard Howitt, Jeffery Mount, Peter Moyle, and Barton
   Thompson. 2011. "Managing California's Water: From Conflict to Reconciliation." In. San Francisco,
   CA: Public Policy Institute of California.
- Healey, M.C. 1991. 'Life history of chinook salmon (Oncorhynchus tshawytscha).' in C. Groot and L.
  Margolis (eds.), Pacific salmon life histories (University of British Columbia Press: Vancouver, British Columbia).
- Jackson, J. R., G. B. Pasternack, and J. M. Wheaton. 2015. 'Virtual manipulation of topography to test
   potential pool-riffle maintenance mechanisms', Geomorphology, 228: 617-27.
- Jacobson, Robert B., and David L. Galat. 2006. 'Flow and form in rehabilitation of large-river ecosystems:
   An example from the Lower Missouri River', Geomorphology, 77: 249-69.
- Jowett, Ian G. 1993. 'A method for objectively identifying pool, run, and riffle habitats from physical
   measurements', New Zealand journal of marine and freshwater research, 27: 241-48.
- Kasprak, Alan, Nate Hough-Snee, Tim Beechie, Nicolaas Bouwes, Gary Brierley, Reid Camp, Kirstie
  Fryirs, Hiroo Imaki, Martha Jensen, Gary O'Brien, David Rosgen, and Joseph Wheaton. 2016. 'The
  Blurred Line between Form and Process: A Comparison of Stream Channel Classification
  Frameworks', PLoS One, 11: e0150293.
- Konrad, Christopher P, Derek B Booth, Stephen J Burges, and David R Montgomery. 2002. 'Partial
   entrainment of gravel bars during floods', Water Resources Research, 38.
- Lai, Young G. 2008. "SRH-2D Version 2: Theory and User's Manual." In. Denver, CO: U.S. Department of the Interior.
- Lane, Belize A., Helen E. Dahlke, Gregory B. Pasternack, and Samuel Sandoval-Solis. 2017. 'Revealing
   the diversity of natural hydrologic regimes in California with relevance for environmental flows
   applications', Journal of American Water Resources Association (JAWRA).
- Lane, Belize A., Gregory B. Pasternack, Helen E. Dahlke, and Samuel Sandoval-Solis. 2017. 'The role of
   topographic variability in river channel classification', Physical Progress in Geography.

- MacWilliams, Michael L., Joseph M. Wheaton, Gregory B. Pasternack, Robert L. Street, and Peter K.
   Kitanidis. 2006. 'Flow convergence routing hypothesis for pool-riffle maintenance in alluvial rivers',
   Water Resources Research, 42.
- Magilligan, Francis J., and K.H. Nislow. 2005. 'Changes in hydrologic regime by dams', Geomorphology,
   770 71: 61-78.
- Montgomery, D. R., and J. M. Buffington. 1997. 'Channel reach morphology in mountain basins', GSA
   Bulletin.
- 773 Montgomery, D. R., Buffington, J. 1997. 'Channel-reach morphology in mountain rivers'.
- Moyle, Peter B., and Paul J. Randall. 1998. 'Evaluating the Biotic Integrity of Watersheds in the Sierra
   Nevada, California', Conservation Biology, 12: 1318-26.
- Murray, Orla, Martin Thoms, and Scott Rayburg. 2006. "The diversity of inundated areas in semiarid flood
   plain ecosystems." In Sediment Dynamics and the Hydromorphology of Fluvial Systems. Dundee, UK:
   IAHS Publication.
- Null, Sarah E., and Joshua H. Viers. 2013. 'In bad waters: Water year classification in nonstationary climates', Water Resources Research, 49: 1137-48.
- Parasiewicz, Piotr. 2007. 'Using MesoHABSIM to develop reference habitat template and ecological
   management scenarios', River Research and Applications, 23: 924-32.
- Pasternack, Gregory B., Michael K. Bounrisavong, and Kaushal K. Parikh. 2008. 'Backwater control on riffle–pool hydraulics, fish habitat quality, and sediment transport regime in gravel-bed rivers', Journal of Hydrology, 357: 125-39.
- Pasternack, Gregory Brian. 2011. 2D modeling and ecohydraulic analysis (University of California at Davis).
- Poff, N. LeRoy, and J. V. Ward. 1990. 'Physical habitat template of lotic systems: Recovery in the context
   of historical pattern of spatiotemporal heterogeneity', Environmental Management, 14: 629.
- Poff, N. Leroy; Allan, J. David; Bain, Mark B.; Karr, James R.; Prestegaard, Karen L.; Richter, Brian D.;
  Sparks, Richard E.; Stromberg, Julie C. 1997. 'The natural flow regime: a paradigm for river conservation and restoration', BioScience, 47: 769-84.
- Price, Amina E., Paul Humphries, Ben Gawne, Martin C. Thoms, and John Richardson. 2013. 'Effects of
   discharge regulation on slackwater characteristics at multiple scales in a lowland river', Canadian
   Journal of Fisheries and Aquatic Sciences, 70: 253-62.
- Richards, KS. 1976. 'The morphology of riffle-pool sequences', Earth Surface Processes, 1: 71-88.
- Richter, Brian D., and Holly E. Richter. 2000. 'Prescribing Flood Regimes to Sustain Riparian Ecosystems
   along Meandering Rivers
- Prescripción de Regímenes de Inundación para Mantener Ecosistemas Riparios a lo Largo de Ríos
  Sinuosos', Conservation Biology, 14: 1467-78.
- Sandoval-Solis, S, DC McKinney, and DP Loucks. 2010. 'Sustainability index for water resources planning
   and management', Journal of Water Resources Planning and Management, 137: 381-90.

- Scown, MW, MC Thoms, and NR De Jager. 2015. 'An index of floodplain surface complexity', Hydrology
   and Earth System Sciences Discussions, 12: 4507-40.
- Small, Melanie J., Martin W. Doyle, Randall L. Fuller, and Rebecca B. Manners. 2008. 'Hydrologic versus
   geomorphic limitation on CPOM storage in stream ecosystems', Freshwater Biology, 53: 1618-31.
- Soulsby, C, AF Youngson, HJ Moir, and IA Malcolm. 2001. 'Fine sediment influence on salmonid spawning habitat in a lowland agricultural stream: a preliminary assessment', Science of the Total Environment, 265: 295-307.
- Thompson, Alan. 1986. 'Secondary flows and the pool-riffle unit: A case study of the processes of meander
   development', Earth Surface Processes and Landforms, 11: 631-41.
- USFWS. 2010a. "Flow-habitat relationships for spring and fall-run Chinook salmon and steelhead/rainbow
   trout spawning in the Yuba River." In, 127. Sacramento, CA: The Energy Planning and Instream Flow
   Branch.
- USFWS. 2010b. "Yuba River Redd Dewatering and Juvenile Stranding Report " In, 60. Sacramento, CA:
   Energy Planning and Instream Flow Branch, .
- 817 Vanzo, Davide, Guido Zolezzi, and Annunziato Siviglia. 2016. 'Eco-hydraulic modelling of the interactions
  818 between hydropeaking and river morphology', Ecohydrology, 9: 421-37.
- White, Jason Q., Gregory B. Pasternack, and Hamish J. Moir. 2010. 'Valley width variation influences
  riffle–pool location and persistence on a rapidly incising gravel-bed river', Geomorphology, 121: 20621.
- Wohl, Ellen, Brian P. Bledsoe, Robert B. Jacobson, N. Leroy Poff, Sarah L. Rathburn, D. M. Walters, and
   Andrew C. Wilcox. 2015. 'The Natural Sediment Regime in Rivers: Broadening the Foundation for
   Ecosystem Management', BioScience, 65: 358-71.
- Wohl, Ellen, and David Merritt. 2005. 'Prediction of mountain stream morphology', Water Resources
   Research, 41.
- Wolman, M. Gordon, and John P. Miller. 1960. 'Magnitude and frequency of forces in geomorphic processes'.
- Worthington, T. A., S. K. Brewer, N. Farless, T. B. Grabowski, and M. S. Gregory. 2014. 'Interacting effects
   of discharge and channel morphology on transport of semibuoyant fish eggs in large, altered river
   systems', PLoS One, 9: e96599.
- Yarnell, S. M., A. J. Lind, and J. F. Mount. 2012. 'Dynamic flow modelling of riverine amphibian habitat
  with application to regulated flow management', River Research and Applications, 28: 177-91.
- Yoshiyama, Ronald M., Frank W. Fisher, and Peter B. Moyle. 1998. 'Historical Abundance and Decline of
   Chinook Salmon in the Central Valley Region of California', North American Journal of Fisheries
   Management, 18: 487-521.
- 837