

1 **What are the drivers of riparian vegetation recruitment and**
2 **establishment in an alpine embanked river?**

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16

17 **Abstract**

18 1 – Rejuvenation operations are often conducted in rivers to manage vegetation encroachment and
19 balance biodiversity benefits and flood risks. Understanding the drivers of vegetation recruitment and
20 establishment is necessary to implement these operations but requires monitoring at spatial and
21 temporal scales that are suited to pioneer riparian environment.

22 2 – Bi-yearly drone surveys were used to retrieve the trajectory of pioneer vegetation in a six
23 kilometres reach of the Isère River (France) following rejuvenation operations. Visible images acquired
24 at the beginning and the end of six consecutive vegetation periods were classified using a convolutional
25 neural network and changes in vegetation cover through time were characterized using spatial
26 operations across dates.

27 3 – Vegetation encroachment in the reach was driven by patch expansion and presented a high inter-
28 annual variability. Provided environmental conditions were favourable, vegetation could cover 20% of
29 bar surfaces in a two-year period following rejuvenation actions. Vegetation destruction in the reach
30 was on the other hand mainly driven by vegetation clearing operations.

31 4 – Vegetation recruitment was found to be mainly driven by the hydrological conditions during the
32 seed dispersal and germination period in May and June. Prolonged bar submersion for flow levels
33 starting at $125 \text{ m}^3 \cdot \text{s}^{-1}$ led to lower recruitment while prolonged dry periods were instead favourable to
34 vegetation encroachment. Under favourable hydrological conditions, recruitment mainly occurred in
35 areas that were covered by fine sediments or water at the beginning of the vegetative period.

36 5 – Spring and winter floods did not result in significant vegetation destruction when plants had at least
37 a year to establish. The surfaces of vegetation destroyed by the Isère River was always lower than the
38 net gain in vegetation cover for each period, and the only spike in vegetation destruction occurred
39 when most vegetation in the reach was less than one year old.

40 6 – Synthesis: Summer hydrology played a key role in vegetation encroachment in the reach by
41 providing either favourable (*i.e.*, prolonged dry periods and no high flows late June) or unfavourable
42 conditions for vegetation recruitment and establishment. Timing eco-morphogenic releases to sustain
43 submersion could help prevent encroachment.

44

45 **Keywords.**

46 Anthropized rivers; Pioneer riparian vegetation; Recruitment processes; Remote Sensing;

47 Biogeomorphology; Ecohydrology

48 **1. Introduction.**

49 River functioning has been altered worldwide by a wide range of pressures such as land use change,
50 damming, and channelization (Belletti et al., 2020; Entwistle et al., 2019; Grill et al., 2019). By modifying
51 the flow and sediment regimes of rivers, human activities have resulted in widespread changes in river
52 geomorphology (Liébault and Piégay, 2002; Surian and Rinaldi, 2003), with impacts on riparian zones
53 and related plants communities (Corenblit et al., 2007; Gurnell et al., 2012). In many regulated rivers,
54 loss in lateral and vertical connectivity led to changes in species assemblages (Breton et al., 2023;
55 Janssen et al., 2021), in plant health (Godfroy et al., 2023) and in the dynamic of riparian habitats
56 (Comiti et al., 2011; Serlet et al., 2018). While riparian vegetation can provide ecosystem services that
57 benefit biodiversity and human activities (Riis et al., 2020), the development of vegetation in river
58 corridors can also lead to changes in roughness, facilitating fine sediment deposition that increase
59 flooding risk, notably in embanked reaches (Vargas-Luna et al., 2015). In such a context, managing
60 riparian vegetation through rejuvenation operations is required. These operations comprise
61 vegetation clearing and bar reprofiling, by lowering topographic levels and exporting fine sediments.
62 While such operations can benefit pioneer riparian plants communities (Janssen et al., 2023),
63 questions remain about the sustainability of these operations and the habitats that have been
64 recreated. This is especially true for highly-anthropized rivers where changes in flow and sediment
65 regimes led to rapid vegetation recruitment and establishment (Comiti et al., 2011; Serlet et al., 2018).

66 Pioneer stages of riparian vegetation are shaped by hydrological and geomorphological drivers
67 (Corenblit et al., 2014; Karrenberg et al., 2002; Mahoney and Rood, 1998; Stella et al., 2006). For
68 example, flow pulses during the growing season have been shown to lead to seedling mortality and
69 reduced survival for the *Salicaceae* (Carter Johnson, 2000; Dixon, 2003; Stokes, 2008). However, these
70 drivers often interact at multiple spatial and temporal scales, and it remains challenging to understand
71 their relative importance and threshold effects associated with them (but see Corenblit et al., 2024a;
72 Pasquale et al., 2014; Stokes, 2008). This requires monitoring and assessing the trajectory of riparian

73 communities following rejuvenation operations over large spatial and temporal extend. Nevertheless,
74 such high frequency monitoring, covering entire river reaches, has very rarely been possible using field-
75 based approaches (González et al., 2015). Furthermore, although *ex situ* experiments have successfully
76 identified the drivers influencing seed germination, such as air temperature and substrate moisture
77 (Chen et al., 2013; Van Splunder et al., 1995), they cannot incorporate the effects of flow regime and
78 geomorphic changes. Advances in remote sensing over the past decades are promising for high
79 frequency monitoring of vegetation recruitment and establishment over large and continuous spatial
80 extents (Corenblit et al., 2024b; Huylenbroeck et al., 2020; Piégay et al., 2020; Viles, 2016).

81 Components of riparian vegetation has been extracted from imagery for three decades now, using
82 increasingly-advanced image processing techniques (Carboneau et al., 2020; Dufour et al., 2012).
83 Satellite time-series (*i.e.*, with a coarse spatial resolution but a dense temporal resolution) have been
84 used to investigate the links between vegetation dynamics and hydrology. Combining field monitoring
85 and Landsat data on the Yangtze River (China) for 20 years, Mei et al. (2025) demonstrated a rise in
86 vegetation cover following reduced flow conditions and shortened submergence periods associated
87 with active restoration efforts to promote vegetation development. On the Po River (Italy), Nones et
88 al. (2024) also used Landsat data to show that the trend toward vegetation encroachment was linked
89 to interactions between past human pressures and climate change, resulting in longer and more
90 frequent dry periods. The role of floods in vegetation destruction was furthermore highlighted using
91 ASTER satellite data on a reach of the Tagliamento River (Italy), where contrasting responses of riparian
92 vegetation to flooding, based on differences in vegetation vigour, were highlighted (Bertoldi et al.,
93 2011a). However, while satellite data provides critical insights on how vegetation responds to changes
94 in flow regimes, they are limited by their low spatial resolution. This may induce a mismatch between
95 the year of recruitment and a year of detection and can prevent a finer scale understanding of the
96 ecological processes driving these trends from year to year.

97 At a finer spatial resolution, studies were conducted using aerial images from airplanes and drones.

98 Time-series of historic aerial images were used to retrieve similar trends as satellite observations, for

99 example with studies on vegetation development on the Isère River being able to go back to the 1930s

100 (Serlet et al., 2018; Vautier et al., 2002). While the temporal depth of these datasets is appealing, their

101 temporal resolution is often poor with gaps of multiple years between surveys. On the other hand,

102 yearly surveys over smaller time-periods were able to shed light on the ecological and hydrological

103 processes behind vegetation recruitment and establishment. On the Drome River (France), Räpple et

104 al. (2017) used yearly aerial images to link recruitment with hydro-climatic drivers such as flow timing

105 and intensity. Low flows and high temperatures following a reset of bare surfaces by floods were

106 favourable to patch recruitment and expansion as there were no scouring floods both following seed

107 dispersal and the winter. In addition, Maeda and Miyamoto (2025) used vegetation maps from UAV

108 surveys acquired for 5 years on the Kinu River (Japan) to build a model predicting vegetation

109 recruitment following a flood. They tested biological, hydrological and geomorphological predictors

110 and showed that the presence of vegetation before the flood was one of the best predictors of

111 vegetation recruitment, highlighting the role of biological processes related to resprout from plant

112 fragments. However, there is still often a mismatch between the temporal and spatial resolution of

113 the data used to monitor riparian vegetation and the ecological processes driving recruitment and

114 establishment, as surveys using one image per year or less are very sensitive to acquisition timing.

115 Following multiple studies documenting rapid vegetation encroachment in the Isère River due to past

116 and ongoing human pressures (Corenblit et al., 2020; Serlet et al., 2018; Vautier et al., 2002), the local

117 stakeholders conducted rejuvenation operations over a 6 km reach, and acquired bi-yearly data to

118 monitor vegetation response over a period of 7 years. The goal of this monitoring was to further our

119 understanding of the processes behind vegetation development in the reach and to objectify when

120 maintenance work is needed. This provides a unique framework to investigate the response of riparian

121 vegetation to these actions and the hydrological drivers of vegetation dynamics in this reach.

122 Therefore, in this paper, we (i) train a convoluted neural network to classify a bi-yearly time-series of

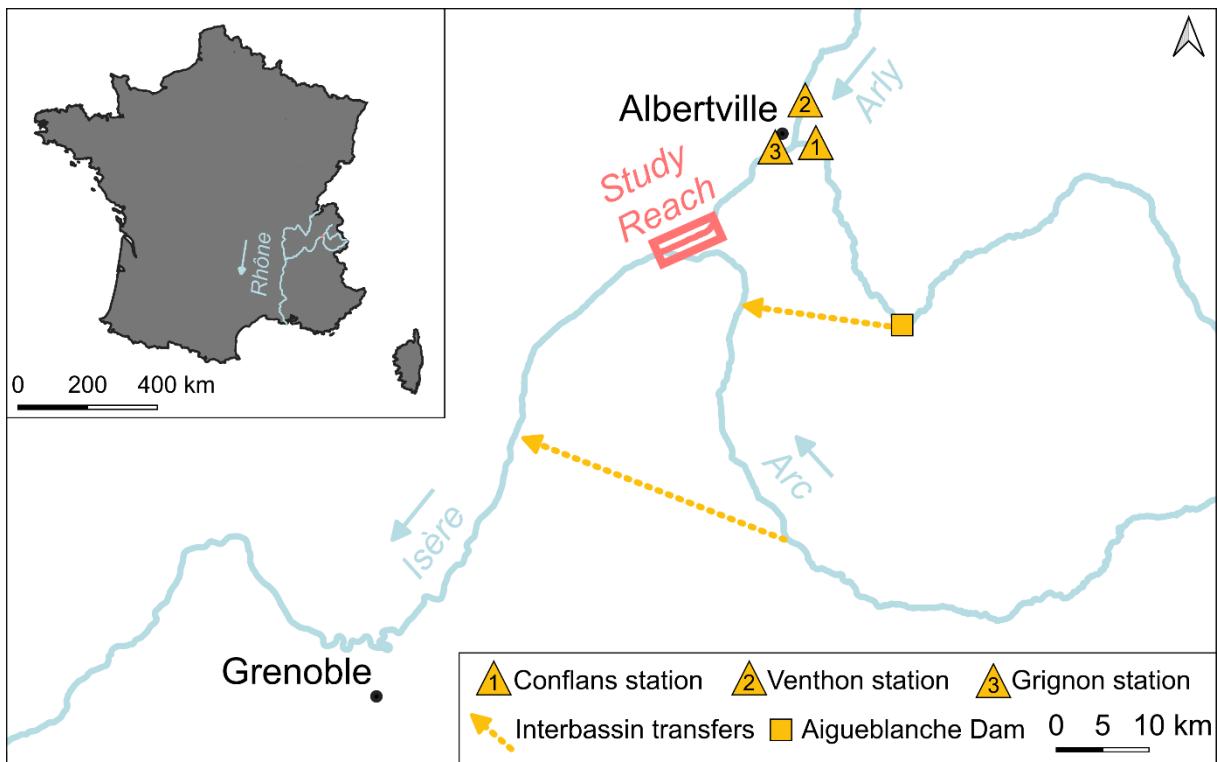
123 drone aerial images acquired over this reach, (ii) document the trajectory of vegetation patches (i.e.,
124 recruitment, lateral growth, destruction, resprouting) during six consecutive vegetative seasons
125 following rejuvenation operations and (iii) test the effects of surface grain size, bar submersion and
126 flooding on vegetation recruitment, establishment and destruction.

127

128 **2. Study site.**

129 The study was conducted on the Isère River, which is located in south-eastern France and is one of the
130 main tributaries to the Rhône River (the mean annual flow at Grenoble is $178 \text{ m}^3.\text{s}^{-1}$, the total length
131 is 286 km and the catchment area is 11.890 km^2) [Figure 1]. The natural hydrological regime of the
132 river was governed by snow and glacier melts with high flows from May to June (Vivian, 1969). The
133 flow regime of the Isère River has been altered since the 1950s, increasing low flows during winter and
134 reducing high flows during summer (Serlet et al., 2018; Vivian, 1994).

135 The floodplain of the river was strongly modified since the end of the 19th century. The initial braided
136 river was first channelized and artificially confined in an embanked channel along its entire length.
137 Most of the floodplain has been converted into agricultural and urban areas. In the latter half of the
138 20th century, sediments were intensively extracted in the main channel and the construction of dams
139 and water bypasses modified the flow regime.



140 **Figure 1.** – Location of the study reach in France and on the Isère River.

141 Following channelization and until the 1950s, the fluvial style of the Isère River was characterized by
 142 alternate bare gravel bars. During high flow periods, those bars were eroded upstream and sediments
 143 were deposited downstream, leading to downstream bar migration that stopped plant succession at
 144 early pioneer stages (Didier, 1994). Geomorphic activity then progressively slowed following sediment
 145 extraction and damming, and vegetation encroachment and bar accretion was observed. Several
 146 studies were previously conducted to understand the drivers of vegetation encroachment. Vautier et
 147 al. (2002) noted an increase in the area of the vegetation in a reach of the Isère river from 5% in 1970
 148 to 20% in 1996, which they attributed to changes in surface grain size and sediment transport. Serlet
 149 et al. (2018) analysed the timing of vegetation encroachment and the associated changes in bar
 150 morphology and mobility. They concluded that encroachment occurred following hydropower
 151 development and sediment mining in the studied reach and resulted in the formation of elongated and
 152 non-migrating bars. More recently, Corenblit et al. (2020) investigated the statistical relationship
 153 between vegetation successions and geomorphological variables at the bar level such as bar planform,

154 bar topography and fine sediment depth, obtained using time-series aerial images coupled with a field
155 survey.

156 Our study covers a reach of about 6 km located in the Combe de Savoie, between the Grésy bridge and
157 the confluence of the Isère and Arc rivers [Figure 1]. In this reach, the local river manager (SISARC) has
158 carried out massive vegetation clearing and sediment removal operations to mitigate flood risk and
159 recreate pioneer environments. Specifically, the reach was rejuvenated by removing woody vegetation
160 and fine sediments from the bars and reshaping them to an elevation that favours frequent
161 submersion. The rejuvenation work was conducted in three phases: (i) in the downstream portion of
162 the study site in 2017, (ii) the upstream portion of the study site in 2018 and (iii) on two vegetated bars
163 that had not yet been rejuvenated in winter 2021–2022 [Appendix F1A]. Since then, maintenance work
164 (i.e., removal of vegetation and reshaping of bars without exporting fine sediments) has been carried
165 out on two bars downstream the study reach in April 2021 [Appendix F1B] and on the bar located at
166 the Grésy bridge in winter 2021–2022 [Appendix F1C].

167

168 **3. Materials & Methods.**

169 *3.1. Remote sensing surveys and hydrological data.*

170 Aerial visible images were acquired over the study reach each year between 2018 and 2023 by local
171 stakeholders in order to investigate the processes behind vegetation development in the reach and to
172 objectify when maintenance work is needed. For each year, two surveys were mobilized: one close to
173 the beginning and one close to the end of the vegetation period [Appendix T1]. For each survey, a DJI
174 Phantom 4 drone was flown at \approx 95 meters height over the active channel and sampled light in three
175 spectral bands: red, green and blue. After the flight, all images were processed by the stakeholders
176 using Agisoft Metashape in order to create an orthomosaic of the study site. The planimetric accuracy
177 of the images was assessed both visually and using 46 permanent ground control points located near
178 the roads on the embankments. We used these post-processed images for the analysis.

179 Hydrological data was retrieved using three different gauging stations upstream the study reach (data
180 available at <https://hydro.eaufrance.fr/>). Due to multiple gaps in data availability, two different time
181 series were created: 1) the data from the Grignon station on the Isère River 14 km upstream the study
182 reach was extracted for the 2021–2023 period; 2) the data from the Conflans station on the Isère River
183 and from the Venthon station on the Arly River were used to reconstruct a hydrological series for the
184 2018–2023 period. The data from each of these stations was aggregated on an hourly timestep and
185 the daily mean discharge was calculated. The hyrogram for the aggregated time-series, along with
186 the dates of all aerial images is presented on Appendix F2.

187 *3.2. Image classification.*

188 Images were first classified using a simple convolutional neural network (CNN) with seven hidden
189 layers with ReLu as the activation function and of descending spatial size, and a final fully-convolutional
190 layer used to predict the classes.

191 A training dataset was built by manually digitizing 125 regions of interest (ROI) for five classes (i.e., 25
192 per class) on each image: vegetation, fine sediments, coarse sediments, water and shadows. All classes
193 were identified visually using photo-interpretation. Training and validation points were randomly
194 sampled in each ROI with an 80-20 split using the *st_sample* function of the *sf* package for the R
195 processing software (Pebesma, 2018; R Core Team, 2023) to obtain 10 000 training points and 2 500
196 validation points per image. Images were normalized and 16x16 training and validation patches were
197 extracted using the Orfeo Toolbox meets TensorFlow (OTBTF) module built on Orfeo Toolbox and
198 TensorFlow (Cresson, 2019). Patches from each image were combined to create a single training
199 dataset for all images (totalling 120 000 training patches and 30 000 validation patches) and were used
200 to train and validate a single neural network.

201 To further assess the accuracy of the two sediment classes, we used field measurements that were
202 acquired as a part of the monitoring plan implemented by the stakeholders. Five surveys spanning
203 multiple days were conducted between 2021 and 2023 but were not synchronous with the flights. For

204 each date, the sand-silt cover of six bars was surveyed by using a 5x5 meters grid and a visual
205 assessment of the relative proportion of fine sediments (*i.e.*, sands and silts) deposited above gravel
206 in a 50 cm radius around each point of the grid (*i.e.*, between 2273 and 2845 points per date). The
207 relative proportion of fines was assessed using six discrete classes: 0–5%, 5–25%, 25–50%, 50–75%,
208 75–95% and 95–100%. For each of these points, the relative proportion of the fine sediments class
209 compared to the coarse sediments class (*i.e.*, not counting surfaces that were classified as shadows or
210 vegetation) was calculated in the 50 cm radius using the closest image available. The relative
211 proportion of fine sediments from the classified images and from the in-field surveys were then
212 compared for each field–image couple.

213

214 *3.3. Patch extraction and semantic classification of change.*

215 To create change maps between each date, the vegetation class was vectorized and gravel bars were
216 manually digitized at low flow conditions each year (*i.e.*, at the end of summer). This layer was used as
217 a mask to remove the vegetation on the riverbanks, that overhangs above the active channel, and the
218 biofilm cover which develops in side channels and shallow water areas. Vegetation patches were then
219 extracted by keeping all polygons with an area > 2 meters (Räpple et al. 2017) and trends in vegetation
220 cover at the pixel level and at the patch level were compared.

221 Spatial operations were used on vegetation patches to provide insights on vegetation changes.
222 Intersections and differences across layers from the most recent to the oldest were used to extract the
223 previous geometry of each vegetation patches and assign changes to four states:

224 (a) Recruitment: defined as the appearance of a vegetation patch that did not intersect a
225 vegetation patch at the end of the previous vegetative season;

226 (b) Lateral growth: defined as the difference between the spatial extent of a vegetation patch at
227 the end of the current vegetative season and the spatial extent of all patches it intersects ($n >$
228 1) from the end of the previous vegetative season;
229 (c) Resprouting and annual plants: defined as vegetation that was already present at the end of
230 the previous vegetative season but not at the beginning of the current vegetative season;
231 (d) Removal: defined as vegetation that was present at the end of the previous vegetative season
232 but is not present anymore at the end of the current vegetative season;

233

234 *3.4. Analysis framework.*

235 Vegetation dynamics were assessed at the reach scale. For each year, the surface area corresponding
236 to each vegetation state was calculated in order to assess trends in vegetation cover. To control for
237 the effects of changes in hydrology and overbank fine sediments cover across dates, the area of the
238 digitized gravel bars and the proportion of both vegetation and fine sediments on the bars were also
239 extracted for each year. In addition, vegetation dynamics were further assessed by exploring the
240 longitudinal vegetation encroachment pattern using a spatially-continuous segmentation of the river
241 reach into successive discrete river segments of 100 m long (Alber and Piégay, 2011). This process was
242 achieved using the segmentation algorithm of the Fluvial Corridor Toolbox (see Rousson and Dunesme,
243 2024; Roux et al., 2015).

244 Three variables were then investigated as potential drivers: the initial state of new vegetation pixels,
245 the time since the last rejuvenation operation and hydrology.

246 To identify the initial state of each vegetation pixels (i.e., the class at the beginning of the season),
247 change rasters were produced for each vegetation season. Each time, the vegetation layer was
248 reclassified using the change states to obtain initial information for each state. The difference in bar
249 geometry between dates due to the higher flows at the beginning of the season was accounted for by

250 reclassifying as “water” all pixels that were in the spatial extent of the bar at the end of the season but
251 not at the beginning.

252 To understand the effects of rejuvenation operations on vegetation encroachment, the relative area
253 of vegetation on the bars was assessed each year depending on the time since the last operation.
254 Intersection and difference operations between years were used to obtain a map of surface age from
255 the last clearing operation.

256 To link hydrology and vegetation recruitment, we calculated the number of days during the seed
257 dispersal period for which the daily mean discharge exceeded different ecological thresholds. We
258 started with an initial threshold of $125 \text{ m}^3 \cdot \text{s}^{-1}$ that had been previously identified by Jourdain (2017) as
259 the (critical) discharge for which bar coarse sediments was entrained on sparsely-vegetated bars
260 before the rejuvenation work was conducted in the reach. We iterated on this initial threshold with a
261 $10 \text{ m}^3 \cdot \text{s}^{-1}$ step and across a range of $\pm 30 \text{ m}^3 \cdot \text{s}^{-1}$ and determined the coefficient of determination of the
262 relationship between the area of recruited patches and the number of days the threshold was
263 exceeded. Based on field observations, the May–June period was selected for the recruitment of the
264 Salicaceae species present in the study site.

265 To link hydrology and vegetation destruction, the values of the “Removal” vegetation class were first
266 adjusted by removing polygons that were overlapping with the spatial extent of rejuvenation and
267 maintenance operations in the reach. For each hydrological period (P_i) corresponding to an
268 observation of vegetation destruction (*i.e.*, between two end-of-summer acquisitions), we calculated
269 the excess flow power of flood peaks (Arnaud et al., 2017; Bagnold, 1966). For this, we selected a
270 critical discharge value of $200 \text{ m}^3 \cdot \text{s}^{-1}$ based on the current discharge range targeted by stakeholders for
271 dam flushing and releases ($200\text{--}250 \text{ m}^3 \cdot \text{s}^{-1}$) and whose current aim is to promote erosion and
272 sedimentation processes in the reach. We also summed the number of hours where $Q_{\text{mean}} \geq 200 \text{ m}^3 \cdot \text{s}^{-1}$
273 ¹ for each hydrological period. Two variables were calculated for the excess flow power:

274 - (i) The excess flow power per unit area ($\omega_{max} - \omega_{200}$, in W.m^{-2}) defined as $\frac{\rho g(Q_{max} - Q_{200})S}{w}$,
 275 where ρ is the water density equal to 1000 kg.m^{-3} , g is the acceleration due to gravity equal to
 276 9.81 m.s^{-2} , Q_{max} is the maximum hourly discharge within P_i , Q_{200} is the discharge equal to 200
 277 $\text{m}^3.\text{s}^{-1}$, S is the channel slope of the reach equal to 0.0019 m.m^{-1} , and w is the median active
 278 channel width equal to 95 m;

279 - (ii) The sum of the unit excess flow power for flood peaks ($\sum^{P_i}(\omega_{max} - \omega_{200})$, in W.m^{-2})
 280 defined as $\sum^{P_i} \frac{\rho g(Q_f - Q_{200})S}{w}$, where Q_f are the hourly peak discharges of each event summed
 281 within P_i .

282

283 **4. Results.**

284 *4.1. Classification accuracy.*

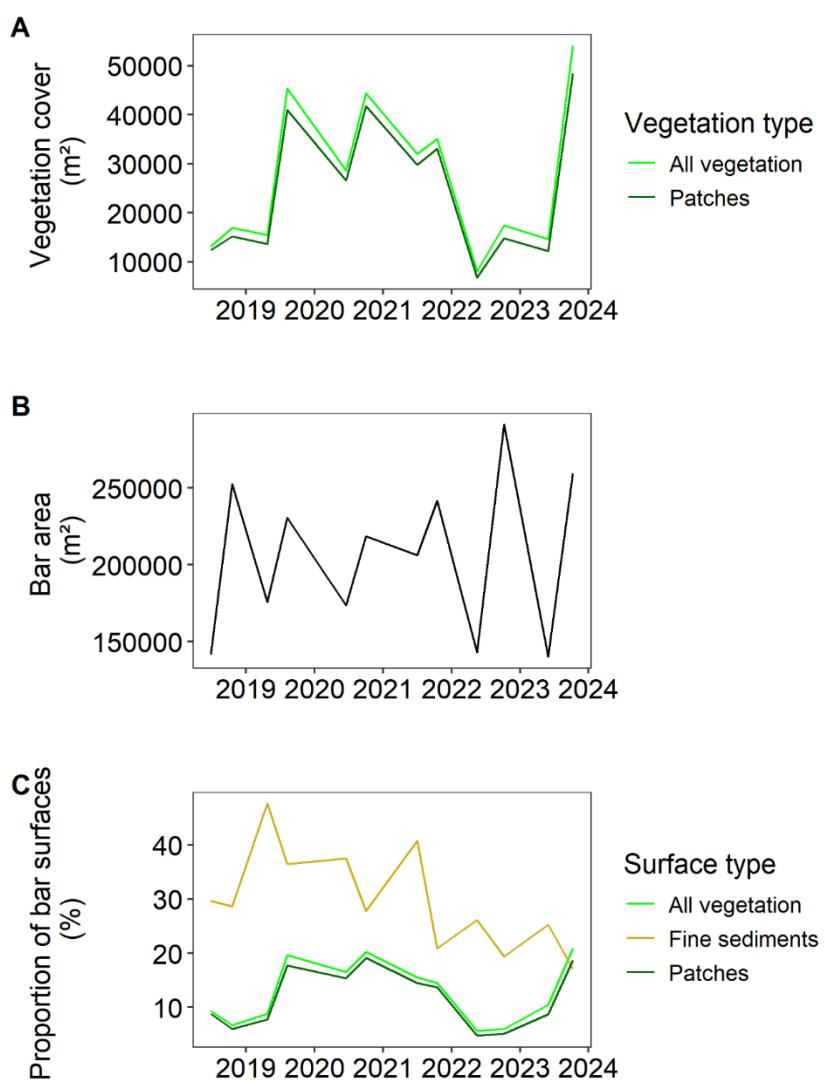
285 A single classifier was trained to process all images in the time-series and reached an overall accuracy
 286 of 0.97 and a kappa index of 0.97 on the validation data. The confusion matrix [Appendix T2] suggests
 287 that confusion between classes occurs mainly between the fine sediments class and either the coarse
 288 sediments class or the water class. Visual inspection of the classified images shows the limitations of
 289 the trained classifier, which is sensitive to changes in the properties of the water column [see Appendix
 290 F3 for details].

291 Comparison of the fine sediments class with in-field observations shows that the proportion of fine
 292 sediments on bar surfaces tends to be underpredicted by classifier [Appendix F4]. For two of the five
 293 surveys (A and C), a 75–95% cover in the field was estimated at around 50%, while it was estimated at
 294 around 75% for the three other surveys (B, D, E). On average, when fine sediments cover was lower
 295 than 50% in the field, the proportion of pixels classified as fine sediments was lower than 10%.
 296 Therefore, pixels classified as fine sediments are mostly coherent with in-field observation of surfaces
 297 dominated by fine sediments and not with coarser materials or a mix of fine and coarse material.

298

299 *4.2. Riparian vegetation dynamics at the reach scale.*

300 Vegetation cover was initially low following the two years of clearing operations and started to increase
301 in 2019 [Figure 2a]. Maintenance work during winter 2020–2021 and winter 2021–2022 led to a
302 reduction in vegetation cover but the system recovered to pre-maintenance state in a two-year
303 timeframe. During the study period, vegetation cover increased by a factor of five, reaching its highest
304 value at the end of summer 2023.



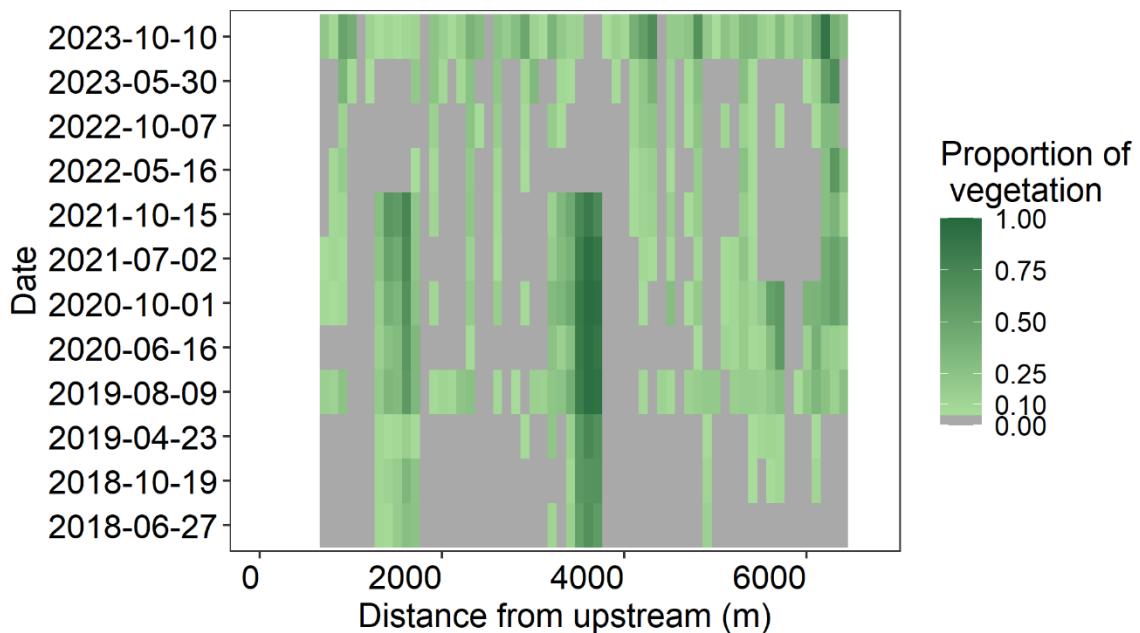
305 **Figure 2.** – Evolution of A) vegetation cover in the study site for all pixels classified as vegetation on
306 the bars and for vegetation patches, B) bar surface area (i.e., bare and vegetated) and C) the
307 proportion of bar surface area occupied by vegetation, vegetation patches or fine sediments.

308

309 The bi-yearly trends in vegetation cover are similar between all pixels classified as vegetation and
310 vegetation patches (i.e., area $\geq 2\text{m}^2$). Each year, vegetation cover increases due to growth and
311 recruitment during the vegetative period and decreases during the off-season, either due to the death
312 of plant with an annual life cycle, destruction, submersion, burial, or a lack of detection at the
313 beginning of the vegetative period.

314 At the same time, bar area varied seasonally with higher area ($\simeq 250\,000\,\text{m}^2$ ($\pm 10\%$)) at the end of
315 vegetative periods, when the flow level is lower [Figure 2b]. It was highest following rejuvenation and
316 maintenance operations in 2022. Overall, vegetation encroachment covered 15% to 20% of bar
317 surfaces starting from the end of the 2019 vegetative season to the maintenance operations that
318 cleared vegetation on targeted bars [Figure 2c]. Those operations resulted in a drop of encroachment
319 to around 5% but it rose again to 20% of bar surfaces at the end of summer 2023.

320 Fine sediments surfaces covered between 20% and 40% of the total area of bars in the reach [Figure
321 2c]. Fine sediments cover was higher at the beginning of the vegetative season than at the end. They
322 occupied around 35% of bar surfaces on average between 2018 and 2021 and dropped between 20%
323 and 25% of bar surfaces starting the end of the 2021 vegetative season to the end of the survey period.
324 Fine sediments cover reached a maximum of near 50% at the beginning of the 2019 vegetative season.
325 Spatializing the distribution of vegetation encroachment along the study reach [Figure 3] shows a
326 coherent increase in the proportion of vegetation on the bars for years where increases in overall
327 vegetation cover occurred.

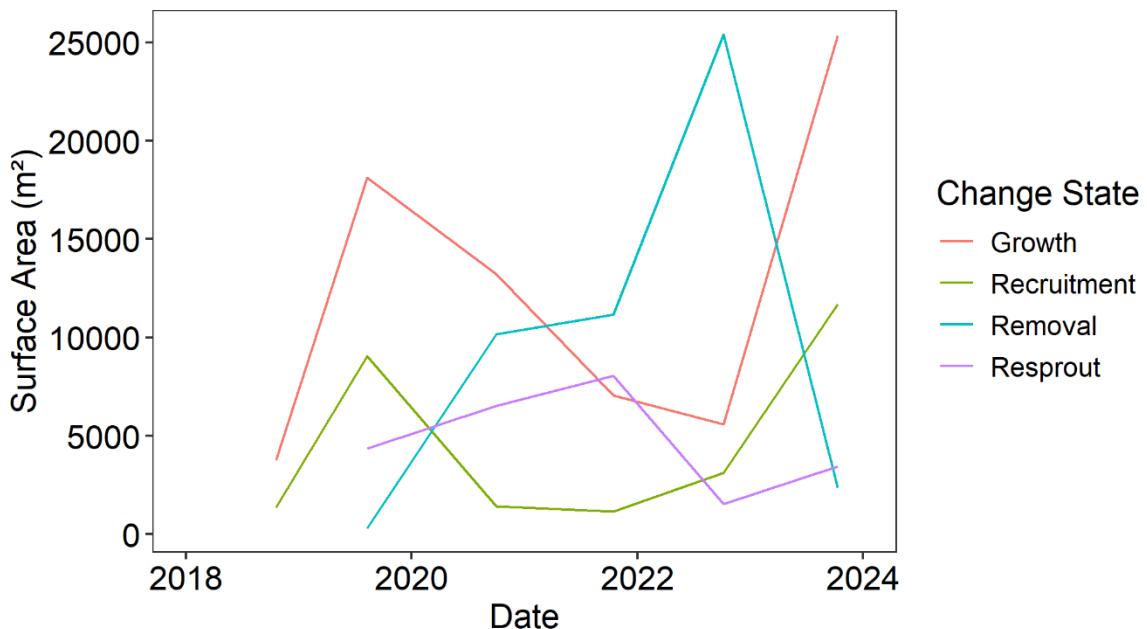


328 **Figure 3.** – Vegetation encroachment on bars along 100 meters segments. When the proportion of
 329 vegetation on the bars in a DGO is between 0% and 5%, values are shown in dark grey instead.

330 At the beginning of the study period, vegetation was located on two vegetated bars that had not been
 331 cleared during the initial rejuvenation work, and on an island downstream our study site [Figure 3].
 332 Vegetation recruitment in 2018 occurred mainly in the downstream-most section – the first one to
 333 undergo clearing work. Vegetation encroachment then increased on all gravel bars during summer
 334 2019. The effects of targeted maintenance work to remove vegetation are visible near the confluence
 335 in 2021 (near 5500 m in Figure 3) and on the two initially vegetated bars in 2022 (near 1500 m and
 336 3500 m in Figure 3). At the end of the study period, most of the river reach is covered by vegetation,
 337 with vegetation encroachment reaching more than 50% of the immersed bar surface in some of the
 338 river segments.

339 The temporal resolution of the data allowed for the distinction of changes of vegetation into different
 340 states [Figure 4]. Recruitment of new vegetation patches was the highest during 2023 and 2019. The
 341 third highest year for patch recruitment was 2022, for which recruitment was still 2 to 3 times higher
 342 than in 2018, 2020 and 2021. Patch growth was the highest in 2023 and 2019 and was also high the
 343 years following high patch recruitment. The area from the lateral expansion of vegetation patches was

344 always higher than the one from appearing patches. The resprouting state reached a higher surface
 345 area than recruitment in 2020 and 2021. Partial or total destruction of vegetation patches was the
 346 highest during winter in 2019–2020, 2020–2021 and 2021–2022. The removal of vegetation due to
 347 maintenance work in 2021 and 2022 accounted for 75% and 91% of the loss in vegetation area
 348 respectively.



349 **Figure 4.**—Evolution of the surface area of all four different vegetation states each year in the study
 350 site: vegetation growth, vegetation recruitment, vegetation removal and vegetation resprout. The
 351 data shown is extracted from the spatial and temporal analysis of the vegetation patches.

352 *4.3. Drivers of riparian vegetation recruitment, establishment and destruction.*

353 The initial state of each pixel at the beginning of the vegetation period was investigated for all pixels
 354 identified as patch recruitment or patch growth [Table 1]. Although the total changes in vegetation
 355 area were different between years, recruitment was the highest (40%–60%) on the fine sediments
 356 class, with two exceptions: during the first and the last years of the study period. In 2018, recruitment
 357 occurred (73%) where sparse vegetation was already present at the end of June, whereas in 2023
 358 recruitment mainly occurred (63%) in areas that were initially submerged. The trend was similar for
 359 patch growth, but with a lower relative proportion of pixels from the fine sediments class and a higher

360 relative proportion from sparse vegetation pixels (> 20% for all years besides 2023). Pixels classified as
 361 fine sediments were also the ones that became vegetated the most. Around 15% to 40% of fine
 362 sediments pixels in the spatial extent of the bars turned into vegetation at the end of the season, a
 363 higher value than for both the coarse sediments pixels and water pixels combined. Following the
 364 previously observed trends, inundated bar areas in spring 2023 were the ones that turned into
 365 vegetation the most. While up to 20% of vegetation was recruited on the coarse sediments class in
 366 2022, less than 5% of the coarse sediments surfaces became vegetated at the end of summer.
 367 Recruitment on already vegetated pixels during spring occurred mostly in 2018, 2021 and 2022, years
 368 for which the first available image was later during the season: near the end of June.

		2018	2019	2020	2021	2022	2023
Recruitment from	<i>Fines</i>	1%	63%	58%	49%	58%	19%
	<i>Coarse</i>	0%	16%	9%	7%	22%	11%
	<i>Vegetation</i>	73%	5%	16%	38%	9%	6%
	<i>Water</i>	26%	15%	15%	5%	11%	63%
Growth from	<i>Fines</i>	1%	58%	48%	42%	53%	35%
	<i>Coarse</i>	0%	6%	5%	3%	12%	8%
	<i>Vegetation</i>	96%	21%	36%	47%	26%	15%
	<i>Water</i>	3%	14%	7%	3%	8%	38%
Fines to	<i>Vegetation patch</i>	0%	24%	17%	7%	15%	38%
	<i>Sparse vegetation</i>	0%	3%	2%	2%	3%	3%
Coarse to	<i>Vegetation patch</i>	0%	4%	2%	1%	2%	5%
	<i>Sparse vegetation</i>	0%	2%	1%	0%	1%	2%

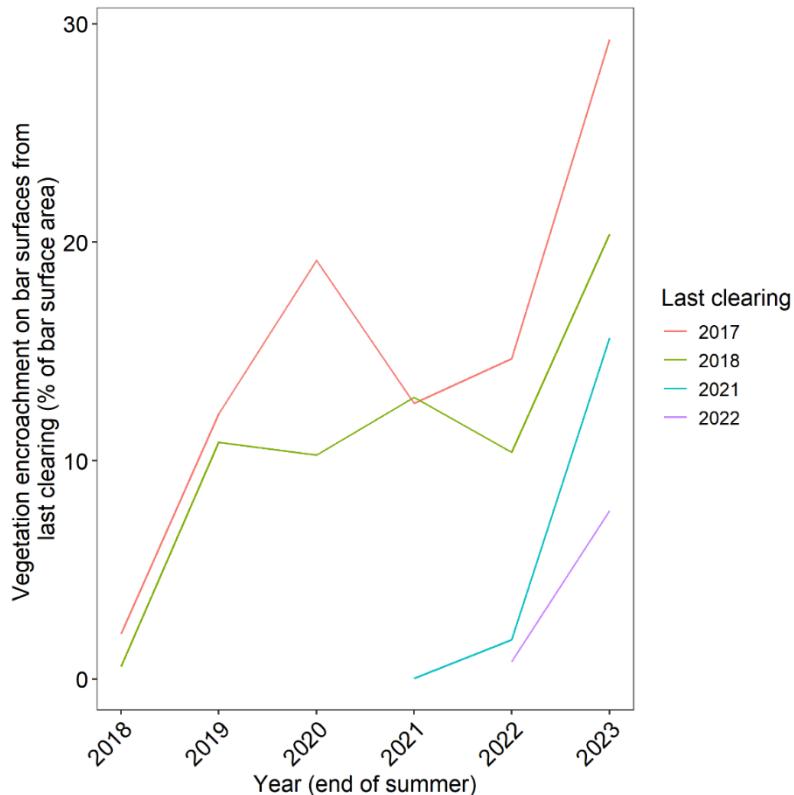
Water to	Vegetation patch	0%	7%	4%	1%	1%	14%
<i>Sparse vegetation</i>	1%	2%	2%	2%	1%	5%	

369 **Table 1.** – Class transitions for each vegetative period: initial state of new vegetation from patch

370 recruitment and patch growth, and proportion of the fine sediments, coarse sediments and water

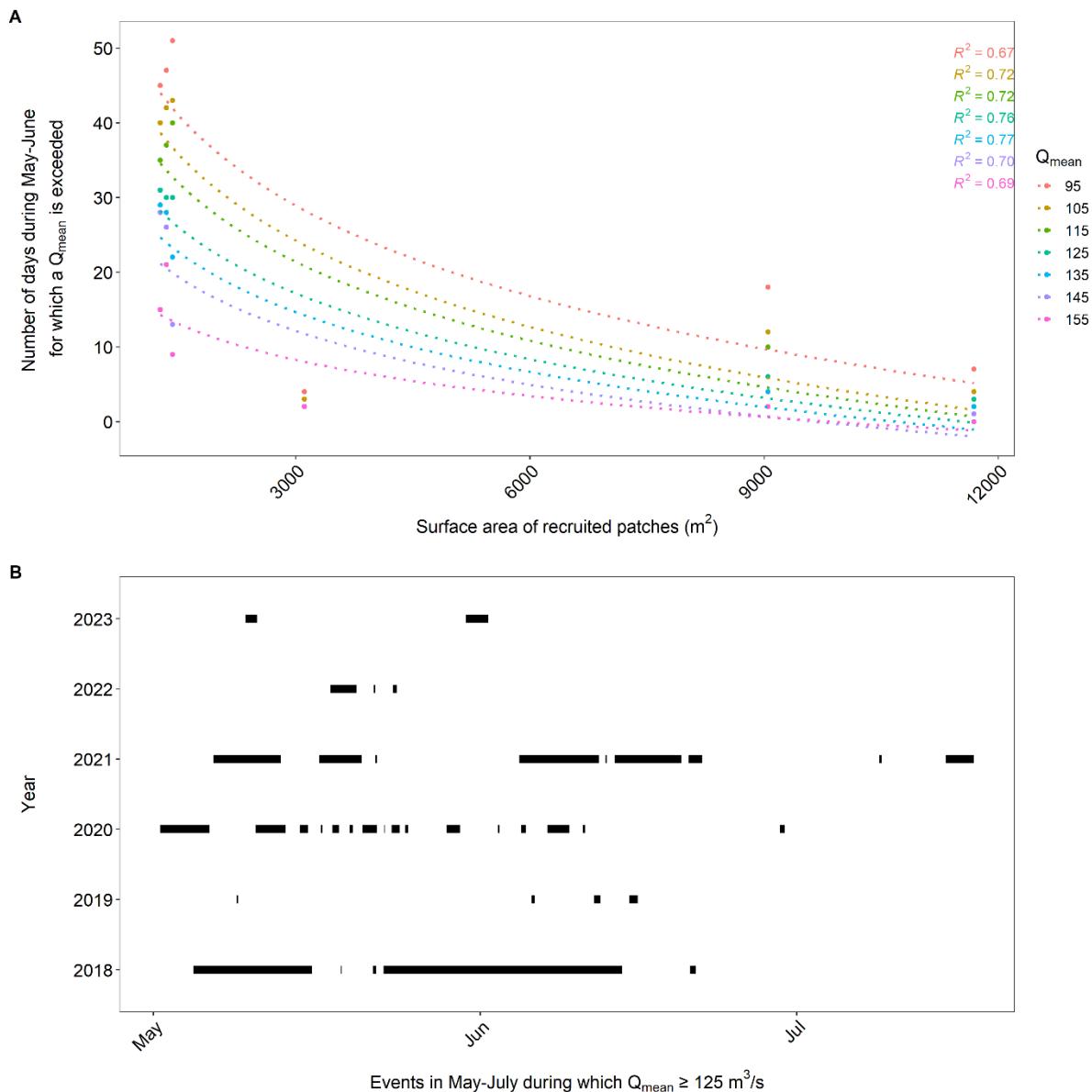
371 classes that becomes vegetation. Transitions from and into shadow are not shown.

372 The effect of time since the last rejuvenation operation revealed that encroachment was very low (<
 373 2%) at the end of the summer following each operation [Figure 5]. It then increased to between 10%
 374 and 20% in a span of two to three years, with the timing of the increase corresponding to the years for
 375 which recruitment and lateral growth were the highest (*i.e.*, 2019 and 2023). For vegetation that was
 376 initially cleared in 2017, encroachment dropped in 2021 following maintenance operations in April,
 377 and for vegetation that was initially cleared in 2018, it dropped in 2022 following the maintenance
 378 operations during winter 2021–2022. At the end of the study period, encroachment was directly
 379 related to the time since last clearing, with older surfaces being more vegetated than newer surfaces.



380 **Figure 5.** – Vegetation encroachment on bar surface area depending on the date of the last clearing
 381 operation conducted. Changes in bar areas related to hydrology are excluded to keep only the
 382 reworked surfaces.

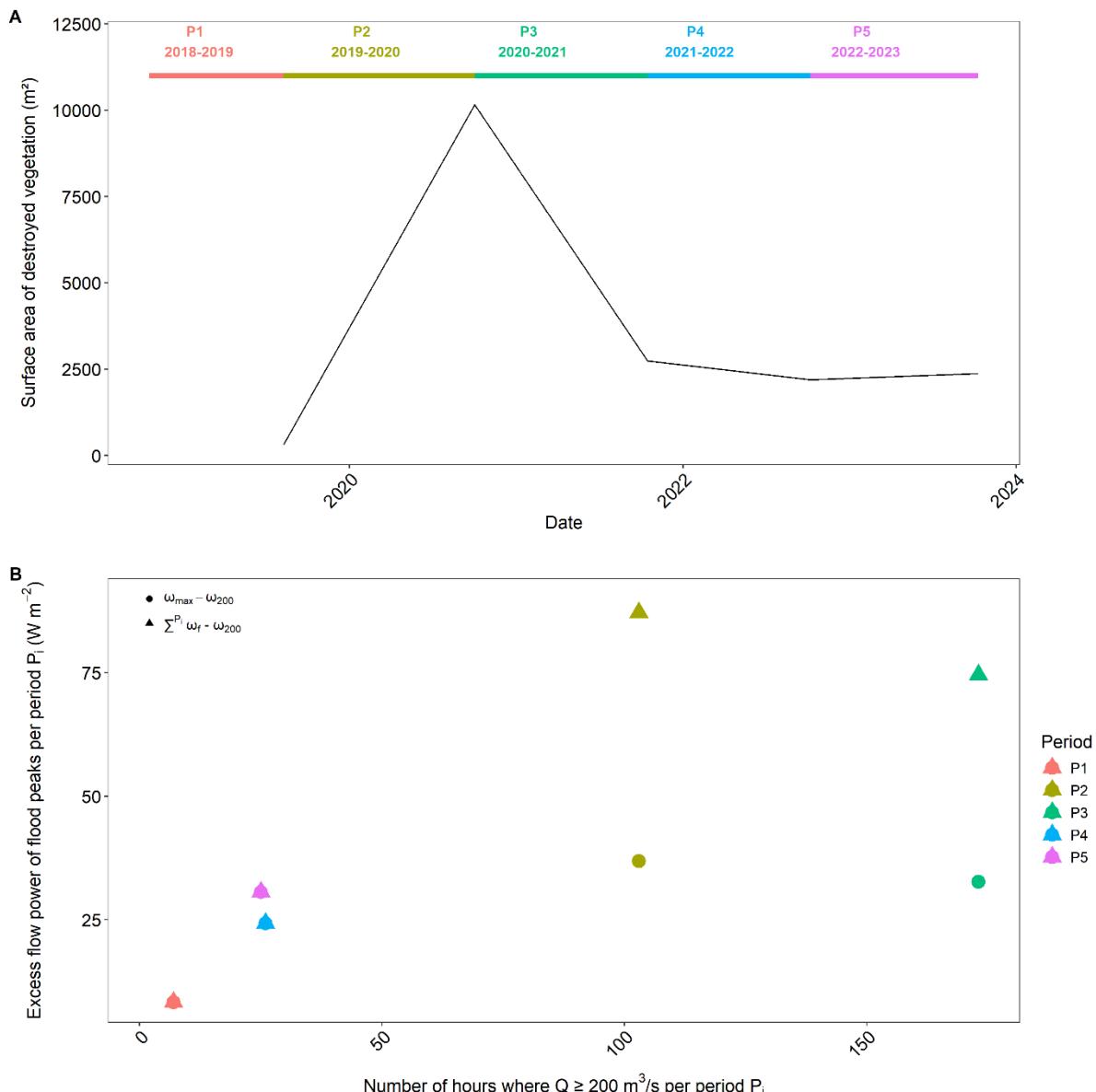
383 Finally, the number of days hydrological thresholds were exceeded was plotted against the area of
 384 recruited patches and the r^2 of the relationship reached its maxima at 0.76 for $125 \text{ m}^3 \cdot \text{s}^{-1}$ and 0.77 for
 385 $135 \text{ m}^3 \cdot \text{s}^{-1}$ [Figure 6A]. Although the number of observations was low, two distinct hydrological
 386 conditions were observed during the vegetative period. The series of years for which patch recruitment
 387 was the lowest (*i.e.*, 2018, 2020 and 2021) were also the ones for which the threshold of bar surface
 388 mobility (*i.e.*, $125 \text{ m}^3 \cdot \text{s}^{-1}$) was exceeded for 30 days (*i.e.*, $\approx 50\%$ of the time). On the other hands, the
 389 years for which patch recruitment was the highest (*i.e.*, 2019, 2022 and 2023) were years for which
 390 this threshold was exceeded for less than 7 days (*i.e.*, $< 10\%$ of the time). The timing and duration of
 391 events where the hourly discharge exceeded this lower threshold were mapped [Figure 6B], which
 392 highlighted that events exceeding $125 \text{ m}^3 \cdot \text{s}^{-1}$ were spread out from the beginning of May to the end of
 393 June during the years with low recruitment.



394 **Figure 6.** – A) Relationships between the number of days where different daily mean discharges
395 (Q_{mean}) are exceeded during the seed dispersion and germination period, and the surface area of
396 recruited patches for each year. B) Timing and duration of the events for which the hourly mean
397 discharge is equal to or exceeds $125 \text{ m}^3/\text{s}^{-1}$.

398 After removing the effects of rejuvenation operations, the area of vegetation that was destroyed each
399 year was the highest between 2019 and 2020, reaching $10\,000 \text{ m}^2$ [Figure 7A]. This was also the period
400 for which the sum of the unit excess flow power for flood peaks was the highest [Figure 7B]. Although
401 it reached similar values between 2020 and 2021, this did not result in an increase in vegetation
402 destruction compared to other periods. Destruction was the lowest for the period immediately after

403 the rejuvenation work (*i.e.*, between 2018 and 2019), which was also the period for which excess flow
 404 power was the lowest.



405 **Figure 7.** – A) Surface area of vegetation destroyed by the Isère River for each hydrological period P_i
 406 between two end-of-summer acquisitions. B) Relationship between the unit excess flow power

407 metrics $\omega_{max} - \omega_{200}$, defined as $\frac{\rho g(Q_{max} - Q_{200})S}{w}$, and $\sum P_i (\omega_{max} - \omega_{200})$, defined as

408 $\sum P_i \frac{\rho g(Q_f - Q_{200})S}{w}$, and the number of hours Q_{200} was exceeded for each hydrological period.

409

410 **5. Discussion.**

411 *5.1. Dynamics of riparian vegetation at the reach scale.*

412 Retrieving vegetation from the remote sensing data enabled us to investigate the dynamics of pioneer
413 vegetation on a 6 km reach of the Isère River. Vegetation encroachment fluctuated over time but a
414 general pattern of increasing cover was highlighted during the study period. Vegetation development
415 was rapid, reaching between 10% and 20% of bar surface from an un-vegetated state in a two-year
416 time frame following both the initial rejuvenation work and later maintenance actions. Although it
417 increased during the study period, vegetation encroachment was partly controlled by local
418 rejuvenation actions during the study period that reset \approx 6 km of bar length, as vegetation
419 encroachment kept increasing on older surfaces.

420 Two distinct states were considered to understand increases in vegetation cover: lateral growth and
421 recruitment. Overall, our results highlighted that the increase in area from the lateral extension of
422 vegetation patches was higher than that from the recruitment of new vegetation patches. In this study,
423 lateral growth is determined through spatial operations that may be interpreted as different ecological
424 processes: the growth of individual plants, the creation of new stems through clonal development and
425 new seedling sprouting near established patches, as they create more favourable conditions for
426 seedlings recruitment (Corenblit et al., 2014; Politti et al., 2018). However, patch recruitment is still
427 necessary for patch growth, and large increases area attributed to patch growth followed increases
428 attributed to recruitment in the same year or the year before. These relationships between patch
429 expansion and recruitment are similar to the ones found on the Drome River (Räpple et al., 2017),
430 where the growth of established vegetation patches was also the main driver of encroachment
431 following an initial spike in recruitment after a reset by floods. We also investigated vegetation loss,
432 which was mainly driven by rejuvenation actions conducted by the stakeholders. Destruction of the
433 vegetation by the river flow was always lower than the combined increases in vegetation area from
434 lateral growth and recruitment. Finally, the intra-annual variability inside each vegetative season was
435 assessed using the resprout state, which included both resprout of vegetation buried by winter flows

436 altogether with annual plants that can quickly reach high biomass during one growing season. Resprout
437 was initially higher than recruitment following the initial clearing work and the spike in recruitment
438 during summer 2019 and then fell off when vegetation in the reach became older. This highlights the
439 need to consider the intra-annual dynamics of riparian vegetation when surveying changes in
440 vegetation cover, and that future studies would also benefit from the ability to better distinguish
441 between woody pioneer vegetation and annual plants – for example using higher spectral resolution
442 data that can enable species mapping (da Silva et al., 2022) or using higher temporal resolution to
443 distinguish species by phenological differences (Michez et al., 2016).

444

445 *5.2. Drivers of vegetation patterns*

446 Hydrology played a key role in the recruitment and establishment of vegetation in the studied reach.
447 Increases in vegetation encroachment during low-flow years occurred regardless of the time since the
448 last clearing. Specifically, a window of at least two weeks of low-flow in May was observed the years
449 where recruitment was the most important, while higher flows were consistent throughout the seed
450 dispersal period when recruitment was low. However, amongst the three years for which high
451 recruitment was observed (2019, 2022 and 2023), the driest year (2022) was also associated with the
452 lowest area of recruited patches. As dry periods have been documented to lead to seedling mortality
453 (Guilloy-Froget et al., 2002; Mahoney and Rood, 1998), we can hypothesize that survival rates could
454 have been lower that year. Our overall observations about vegetation recruitment are coherent with
455 the fact that flow pulses during the growing season are known to lead to seedling mortality for woody
456 pioneer species (Carter Johnson, 2000; Dixon, 2003; Guilloy-Froget et al., 2002; Pasquale et al., 2014;
457 Stokes, 2008) and with the observed response of green vegetation to dryer flows across multiple years
458 (Cohen et al., 2022; Nones et al., 2024). Applying a window of opportunity approach (Balke et al., 2011)
459 to our observations is difficult because of the hydrological variability of the Isère River between years
460 and because the end of the period where seed germination might occur each year is not clear and

461 might vary between species (Stella et al., 2006) and between years due to variations in temperatures
462 (Bourgeois and González, 2019). Further research is therefore needed to characterize the seed
463 dispersal period as well as work at a higher temporal resolution and with topographic data in order to
464 better characterize submersion windows and the effects of specific flood events, but also to determine
465 the relative contribution of annual and perennial species to patch recruitment and patch growth.
466 Beside hydrology, we found that fine sediments deposits were more likely to become vegetated at the
467 end of the season, both for patch recruitment and patch expansion. However, for lateral growth, these
468 findings need to additionally consider the feedback between pioneer vegetation and sediments
469 transport that facilitate sediments deposition (Corenblit et al., 2016). In years with higher flows when
470 the initial state images were acquired, submerged areas were also likelier to become vegetated. This
471 suggests that low flow areas where fine sediments are preferentially deposited and where moisture is
472 high are the ones where vegetation establishes preferentially (Díaz-Alba et al., 2023; González et al.,
473 2018).

474 The analysis of vegetation destruction shows that vegetation renewal from hydrological events was
475 low during the study period. The floods that occurred during the period were comprised between two-
476 year return time floods ($Q_2 = 290 \text{ m}^3.\text{s}^{-1}$) and five-year return time floods ($Q_5 = 415 \text{ m}^3.\text{s}^{-1}$) and the
477 effect of floods with a higher instant discharge could not be assessed. The frequent floods that occur
478 each year with a peak discharge of at least $200 \text{ m}^3.\text{s}^{-1}$ are not sufficient to induce a destruction of
479 established patches. While there were two periods for which the sum of the excess flow power for
480 flood peaks was high, only one of the them led to a peak in vegetation destruction (*i.e.*, the 2020–2021
481 period). From a hydrological standpoint, the main differences between 2019–2020 and the 2020–2021
482 hydrological periods were in the number of hours where the critical discharge was exceeded: it was
483 higher during the 2020–2021 due to a higher number of smaller floods, while it was lower during 2019
484 – 2020 since two floods reached a peak discharge $\geq 350 \text{ m}^3.\text{s}^{-1}$. From an ecological standpoint, the state
485 and the nature of riparian vegetation was also different during those two periods: the surfaces
486 recruited spiked in summer 2019 and were far lower than the surfaces destroyed by the following

487 hydrological period (*i.e.*, 10 000 m²), while recruitment was low during summer 2020 and, as a result,
488 surfaces recruited this year were lower than the surfaces destroyed by the following hydrological
489 period (*i.e.*, 2500 m²). Our results suggest that vegetation that was able to establish and resist the
490 floods from the 2019–2020 period was also able to resist the floods from the 2020–2021 period. An
491 increasing resistance to flooding due to the rapid establishment of roots and the need for morphogenic
492 change to uproot the plant for erosion has been shown in the literature for the perennial willows and
493 poplars (Bywater-Reyes et al., 2015; Karrenberg et al., 2002; Politti et al., 2018), which contribute to
494 the dynamics of vegetation patches in our study reach.

495

496 5.3. – Implications for river management and restoration.

497 González et al. (2015) considered active hydro-geomorphic actions such as water releases and bar
498 levelling as parts of a larger panel of tools to restore riparian vegetation in rivers. The same measures
499 can be also used to prevent vegetation encroachment, with studies citing both vegetation clearing
500 (Janssen et al., 2023) and water releases (Loire et al., 2021; Rivaes et al., 2015) as potential tools to
501 reset the system and remove established vegetation. In this study, we were able to assess the effects
502 of vegetation clearing and of hydrological drivers on the dynamics of pioneer vegetation over a six
503 years period.

504 On the Isère River, mechanical interventions on vegetated bars had a lasting impact on vegetation
505 encroachment and forced a reset of the system. Before the rejuvenation operations, the vegetation
506 cover in the reach was very high – reaching \approx 70% of the total channel (Serlet et al., 2018). In 2023,
507 vegetation cover occupied less than 30% of the surface of all bars that had been rejuvenated since the
508 first clearing operations took place in 2017 which corresponds to \approx 9% of the total channel. Targeted
509 interventions on the more vegetated bars in 2021 and in 2022 had a local but visible effect on the
510 repartition of vegetation at the reach scale but were still sensitive to recruitment and establishment
511 during summer 2023. The nature of the vegetation was also different from before the rejuvenation

512 works as it was comprised of vegetation at the pioneer stage rather than the post-pioneer and mature
513 upland forests that were described before (Corenblit et al., 2020).

514 Morphogenic flows in the Isère River currently target bar submersion and erosive processes using a
515 range of discharges between $200 \text{ m}^3 \cdot \text{s}^{-1}$ and $250 \text{ m}^3 \cdot \text{s}^{-1}$ (Melun, 2024). Our results show that attempting
516 to destroy established vegetation by erosion and uprooting would require the use of higher peak
517 discharges (e.g., $\geq 350 \text{ m}^3 \cdot \text{s}^{-1}$) and would need to target the winter or spring following high vegetation
518 recruitment, before it can establish further. While these values are not realistic for managing
519 vegetation in the reach as attaining these discharges cannot be guaranteed, our results also show that
520 promoting bar submersion can be effective to prevent the recruitment of vegetation. In particular, we
521 identified a threshold between $125 \text{ m}^3 \cdot \text{s}^{-1}$ and $135 \text{ m}^3 \cdot \text{s}^{-1}$ that was inversely correlated to the area of
522 recruited patches each year. These values are lower than the current flow level targets for
523 morphogenic releases in the reach and could be tested to sustain submersion over a longer time period
524 during the seed dispersal period. Therefore, while previous studies suggested that timing flow releases
525 with the peak of the dispersal period was an active restoration method that could be implemented to
526 promote the recruitment of Salicaceae (Mahoney and Rood, 1998; Rood et al., 2003), we suggest that
527 flow releases following seed dispersal and germination can help prevent the development of
528 vegetation in rivers where management needs are different.

529

530 **6. Conclusion.**

531 Bi-yearly colour aerial images acquired over a 6 km reach enabled the retrieval of both riparian
532 vegetation and information on the surface grain size of gravel bars. The spatio-temporal dynamics of
533 riparian vegetation were explored, showing a trend towards vegetation encroachment at the reach
534 scale. The inter-annual variability of changes in vegetation cover was high and driven by patch
535 recruitment and lateral expansion while vegetation destruction was mainly the result of targeted

536 rejuvenation actions. Encroachment was quick following rejuvenations actions and reached 20% of bar
537 surfaces in a two-year period.

538 Vegetation recruitment was found to be mainly driven by the hydrological conditions during the seed
539 dispersal and germination period. Flows between $125 \text{ m}^3 \cdot \text{s}^{-1}$ and $135 \text{ m}^3 \cdot \text{s}^{-1}$ were found to be correlated
540 with lower recruitment and are hypothesized to contribute to the mortality of young seedlings through
541 the submersion of the bars they generate. Under favourable hydrological conditions, vegetation
542 established on areas previously dominated by fine sediments or water, suggesting a preference for
543 areas of low elevation where fine sediments deposit and humidity is high. On the other hand, floods
544 were shown to have little impacts on vegetation in the reach: hydrological periods with high excess
545 flow power only led to vegetation destruction when vegetation had less than one year to establish.
546 This suggests that riparian vegetation in the Isère River establishes very quickly once seedlings survive
547 summer floods.

548 Our results have strong implications on the use of river flow in vegetation management. While current
549 practices suggest that eco-morphogenic flows could be used to limit vegetation encroachment in rivers
550 by destroying established vegetation, we suggest that sustaining summer flows could be used to limit
551 vegetation establishment.

552

553

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559 l’aménagement et du logement Auvergne-Rhône-Alpes, Direction départementale des territoires de

560 Savoie, Electricité de France, Syndicat Mixte de l'Isère et de l'Arc en Combe de Savoie and Assemblée
561 du Pays Tarentaise Vanoise.

562 **Author contributions.**

563 Julien Godfroy, Laurent Borgniet, Hervé Piégay, Gabriel Melun and Philippe Janssen conceived the
564 ideas and designed methodology. Thibault Boissy collected the data. Julien Godfroy analysed the
565 data. Julien Godfroy led the writing of the manuscript. All authors contributed critically to the draft
566 and gave final approval for publication.

567 **Data availability statement.**

568 Raw data used in this study is the property of the stakeholders of the Isère River and is not made
569 publicly available as it contains sensitive information outside the active channel. Post-processed data
570 supporting the analysis, including masked and normalized aerial images over the active channel,
571 classification results and the vegetation layers for each date, are available at
572 <https://doi.org/10.57745/USUGYX>.

573 **Conflict of interest statement.**

574 The authors have no conflict of interest to declare.

575 **Statement on inclusion.**

576 Our study was conducted by scientists and stakeholders that were based in the country where the
577 study was carried out. Both researchers and stakeholders at the national (Office Français de la
578 Biodiversité) and local (Syndicat Mixte de l'Isère et de l'Arc en Combe de Savoie) level worked on the
579 design of the study, and the data used was collected by the local stakeholders. They were associated
580 to the writing of the paper and made critical contributions to the drafts. The results of this study were
581 also conveyed more broadly to the other stakeholders involved in the integrative management of the
582 Isère River basin (Plan de pérennisation de l'Isère en Combe de Savoie).

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808

809 **Supplementary Materials.**

810 **Appendix T1.** – Characteristics of the datasets used in this study.

Date	Spatial resolution	Daily mean discharge	Days between flights
2018-06-27	5 cm	53 m ³ .s ⁻¹	115
2018-10-19	5 cm	15 m ³ .s ⁻¹	
2019-04-23	5 cm	59 m ³ .s ⁻¹	108
2019-08-09	5 cm	13 m ³ .s ⁻¹	
2020-06-16	5 cm	76 m ³ .s ⁻¹	107
2020-10-01	5 cm	43 m ³ .s ⁻¹	
2021-07-02	5 cm	54 m ³ .s ⁻¹	105
2021-10-15	5 cm	39 m ³ .s ⁻¹	
2022-05-16	5 cm	78 m ³ .s ⁻¹	144
2022-10-07	5 cm	17 m ³ .s ⁻¹	
2023-05-30	5 cm	112 m ³ .s ⁻¹	133
2023-10-10	5 cm	33 m ³ .s ⁻¹	

811

812 **Appendix T2.** – Confusion matrix of the classification on the validation data.

		Predicted class of the validation patches				
		Fine	Coarse	Shadow	Vegetation	Water
Actual class of the validation patches	Fine	5743	109	19	1	148
	Coarse	151	5788	0	1	40
	Shadow	26	0	5941	3	30
	Vegetation	3	2	1	5991	3
	Water	159	67	36	3	5735

813

814 **Appendix T3.** – Proportion of each surface class that became vegetation after the vegetative period.

815 Both vegetation patches ($\geq 2\text{m}^2$) and isolated (i.e., sparse) vegetation units ($< 2\text{ m}^2$) are shown.

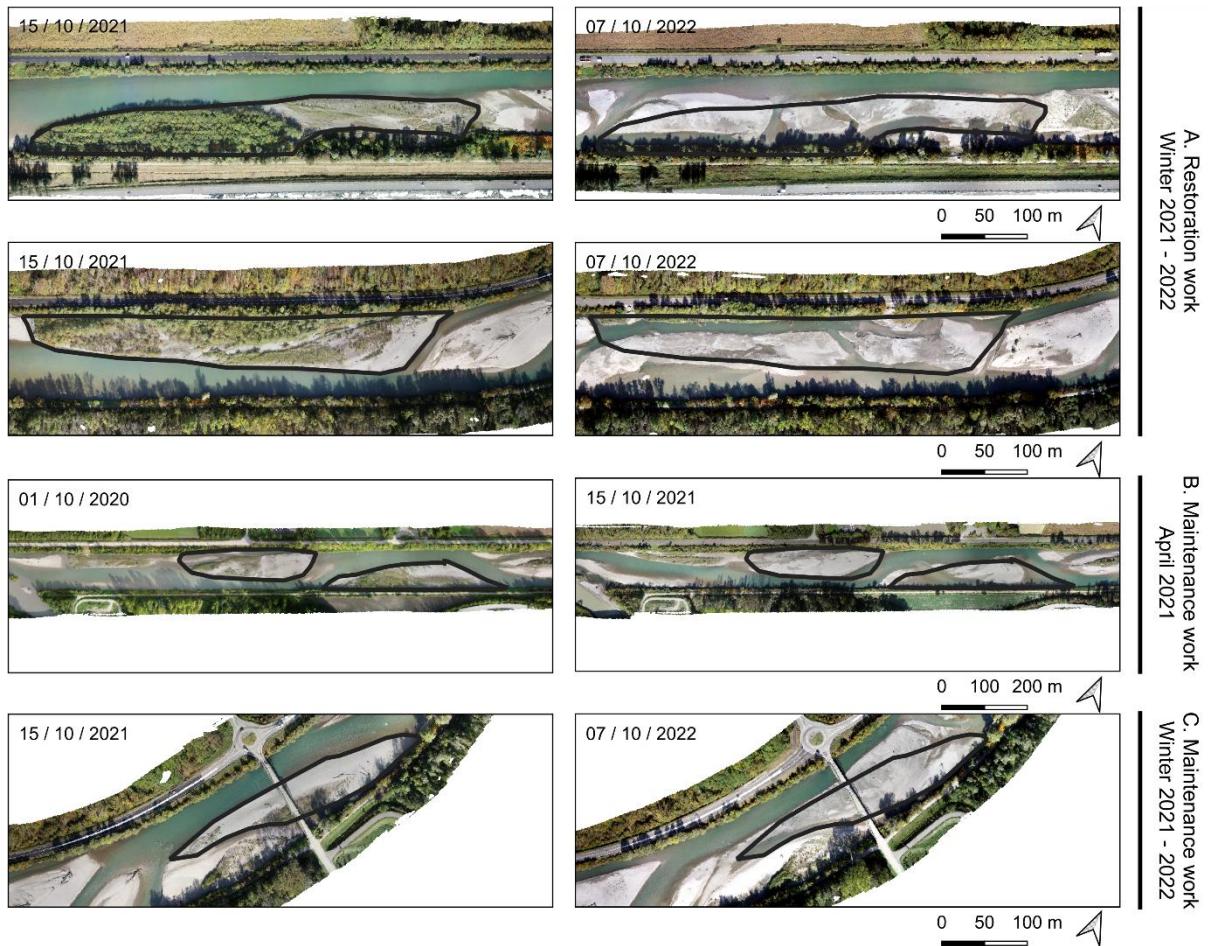
	Fine		Coarse		Water	
	<i>Patches</i>	<i>Sparse</i>	<i>Patches</i>	<i>Sparse</i>	<i>Patches</i>	<i>Sparse</i>
2018	0%	0%	0%	0%	0%	1%
2019	24%	3%	4%	2%	7%	2%
2020	17%	2%	2%	1%	4%	2%
2021	7%	2%	1%	0%	1%	2%
2022	15%	3%	2%	1%	1%	1%
2023	38%	3%	5%	2%	14%	5%

816

817

818 **Appendix F1.** – Before–after view of the bars that underwent (A) vegetation clearing work during
819 winter 2021–2022, (B) maintenance work in April 2021 and (C) maintenance work during winter
820 2021–2022.

821



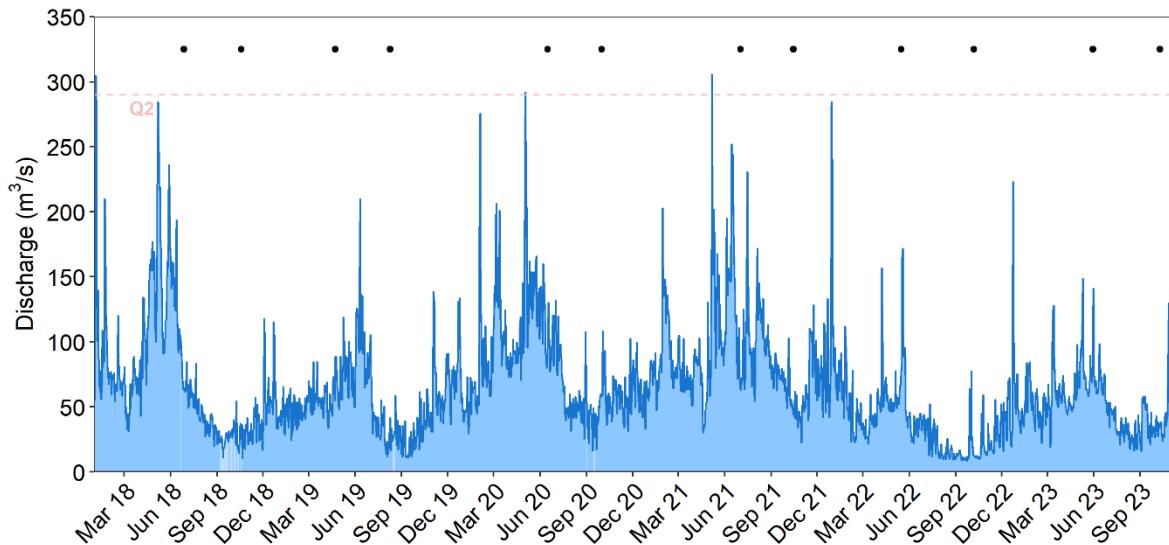
822

823

824

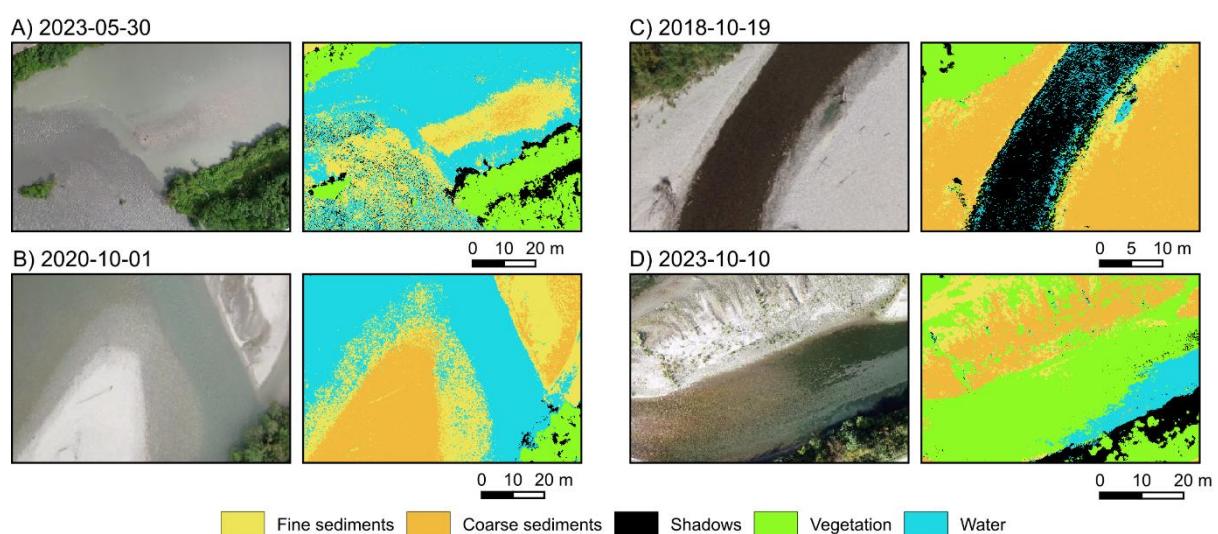
825

826 **Appendix F2.** – Daily mean discharge series merging the data from the Grignon station (from 2022)
827 and the reconstructed series from the Conflans and Venthon stations (before 2022). The black dots are
828 the date of acquisition for each aerial image and areas where the curve is interpolated are shown in
829 light blue.



830

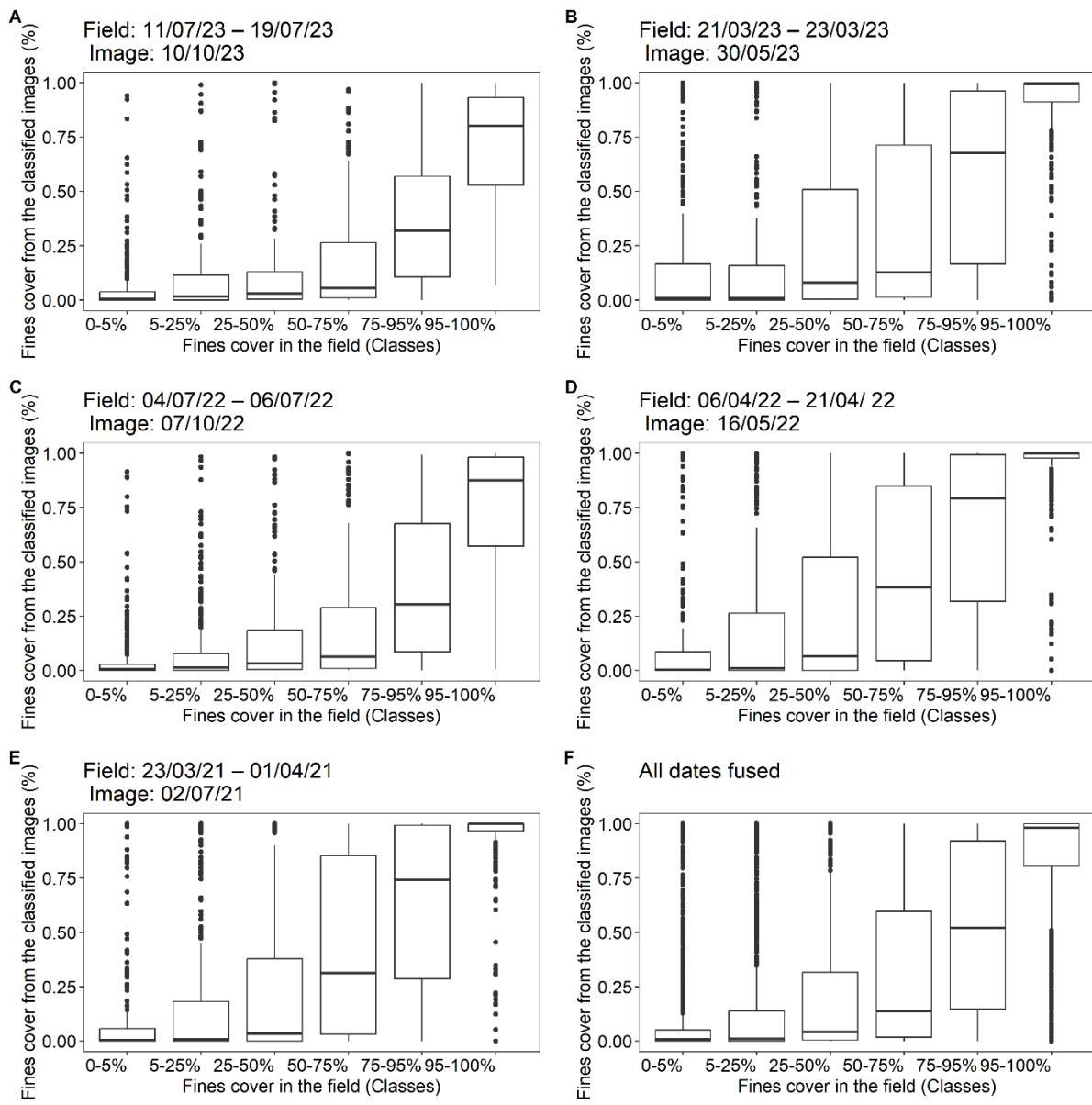
831 **Appendix F3.** – Panels highlighting the limitations of the trained classifier: (A) high turbidity, (B) low
832 and transparent water table where submerged sediments are visible, (C) low water table on darker
833 substrate and (D) biofilm cover.



834

835

836 **Appendix F4.** – Comparison between the relative cover of fine sediments observed in field and
 837 predicted on the images.



838

839