

1 **Can multi-scale thermal infrared imaging help validate and monitor water stress in**  
2 **alluvial forests?**

3  
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25 **Abstract.**

26 Alluvial forests are fragile and sensitive to drought induced by climate change and  
27 exacerbated by altered flow regimes. Our ability to detect and map their sensitivity to drought  
28 is therefore crucial to evaluate the effects of climate change and adjust management practices.  
29 In such a context, we explore the potential of multi-scale thermal infrared imagery (TIR) to  
30 diagnose the sensitivity of alluvial forests to drought events. In summer 2022, we sampled  
31 leaves and phloem on *Populus nigra* trees from two sites with contrasted hydrological  
32 connectivity along the Ain River (France) in order to investigate the seasonality of water  
33 stress and act as ground truth for airborne TIR images. To map forest sensitivity to drought,  
34 we then used a set of TIR data from four existing airborne campaigns and Landsat archives  
35 over a larger spatial and temporal extent. Field data showed that stress conditions were  
36 reached for both sites during summer but were higher in the site with lower groundwater  
37 connectivity, which was also the case for individual tree crown temperature. At the forest plot  
38 scale, canopy temperature was linked to forest connectivity for two of four airborne TIR  
39 campaigns, with higher values in the more degraded reaches. The data from the Landsat  
40 archives at the landscape scale was used to locate the areas of the riparian forest impacted by  
41 a historical drought event, and monitor their recovery. TIR data showed promising results to  
42 help detect and map tree water stress in riparian environments. However, stress is not detected  
43 in all TIR campaigns, demonstrating that in-field ecophysiological measurements are  
44 complementary to validate observations and one-shot acquisitions are not enough to diagnose  
45 stress. More integrative indicators of drought stress are needed at a seasonal scale, one-shot  
46 acquisitions on a given day can inform potential heat disturbance effects but do not really give  
47 information on the cumulative effects of heat pulses over the whole vegetative season (ramp-  
48 disturbance effect). Landsat data was useful to identify trends but may be less representative

49 of stress due to coarse spatial resolution and potential confounding factors related to changes  
50 in successional stages (tree height and density...) at larger temporal scales.

51

52 **Key words.**

53 Riparian Vegetation, Water Stress, Thermal Infrared Remote Sensing, Multi-method  
54 Approach, Anthropogenic Alterations

55

56 **1. Introduction.**

57 Alluvial forests are unique environments characterized by complex feedback processes with  
58 their river system that provide a set of ecosystem services (Naiman et al., 1988; Riis et al.,  
59 2020). Alluvial forests are biodiversity hotspots (Naiman et al., 1993), improve water quality  
60 (Dosskey et al., 2010; Tabacchi et al., 2000), help stabilize channel banks (Simon and  
61 Collison, 2002) and play a key role in ensuring the sustainability of good hydromorphological  
62 conditions in river systems (González del Tánago et al., 2021).

63 Human development in the 20<sup>th</sup> century has however caused increased pressure on riparian  
64 vegetation (Bravard et al., 1997; Breton et al., 2023; Comiti et al., 2011; Poff et al., 2007).

65 Activities such as gravel mining, damming and engineering works have induced tree stress  
66 and increased mortality (Scott et al., 2000), and led to a reduction of forest renewal by  
67 increasing bank resistance or reducing channel mobility in other cases (Décamps et al., 1988;  
68 Dépret et al., 2023). Key riparian species are also sensitive to climate change (O’Briain,  
69 2019). Higher temperatures and variations in precipitations and flow regimes brought about  
70 by climate change can impact tree water use and water availability (Rivaes et al., 2013; Stella  
71 et al., 2013b). Groundwater decline due to drought or anthropogenic alterations can induce

72 stress and higher mortality in riparian woodlands, and lead to long-term shifts in species  
73 composition (Breton et al., 2023; Janssen et al., 2020; Kibler et al., 2021; Rohde et al., 2021).  
74 Therefore, it is important to monitor the response of riparian vegetation to events affecting  
75 water availability such as channel incision due to damming or mining and droughts due to  
76 climate change. Such a monitoring effort can then help target conservation and restoration  
77 actions to mitigate negative impacts of global change on riparian forests and assess their  
78 efficiency.

79 Field surveys are traditionally used to assess the water status of individual trees. For example,  
80 tree rings can give insights into events happening within the lifespan of an individual and  
81 radial growth anomalies can occur during dryer years (Dufour and Piégay, 2008; Singer et al.,  
82 2013; Stella et al., 2013a). Other less destructive methods such as sampling tree leaves or  
83 phloem can also inform about tree water status. The measure of leaf water potential (LWP)  
84 can indicate on-going water stress of the tree at the time of sampling (Brodribb and Holbrook,  
85 2003; Scholander et al., 1965). Carbon isotope analysis of the phloem content can help  
86 determine how efficient a tree is at using water to produce biomass (intrinsic Water Use  
87 Efficiency, iWUE), which responds to water deficits such as drought-related stress (Klein et  
88 al., 2016; Seibt et al., 2008). One of the main limitations of such approaches however is that  
89 they are time-intensive and cannot easily be conducted at a large scale within a short time  
90 window.

91 At the scale of communities or river reaches, remote-sensing has been used to assess forest  
92 status with imagery in the visible and near infrared spectrum (Kibler et al., 2021), or in the  
93 thermal infrared (TIR) spectrum (Mayes et al., 2020). In these cases, TIR data is used since  
94 differences in canopy temperatures are a good indicator of differences in evapotranspiration.  
95 Even though remote sensing applications are increasingly common to study riparian

96 vegetation, the use of TIR sensors has been mostly limited to the use of satellite-based sensors  
 97 in arid or semi-arid climates where drought events are common today [Table 1].

98 **Table 1. - Examples of recent thermal infrared studies focusing on riparian vegetation.**

Study	Sensor	Vector	Objective	Climate	Time Series
(Neale et al., 2011)	FLIR SC640	Aircraft	Estimating evapo-transpiration	Mediterranean (California)	
(Gokool et al., 2017)	MODIS + Landsat	Satellite	Retrieving daily evapo-transpiration from satellite data	Semi-Arid (South Africa)	X
(Fairfax and Small, 2018)	Landsat	Satellite	Assessing the impact of beaver damming on evapo-transpiration	Arid (Nevada)	X
(Lurtz et al., 2020)	Landsat	Satellite	Relationship between evapo-transpiration and modeled water table depths	Semi-Arid (Colorado)	X
(Cieřkowski et al., 2020)	Landsat	Satellite	Comparing thermal and optical indexes to detect stress	Temperate transitional (Poland)	X
(Mayes et al., 2020)	Landsat	Satellite	Understanding water use at landscape scale	Semi-Arid (Mexico)	X
(Sankey et al., 2021)	SenseFly thermoMAP	UAV	Detecting genetic traits differences	Semi-Arid (Central Arizona)	

100 One of the advantages of using data from satellites is that they can cover large spatial scales at  
101 a temporal resolution that cannot be matched by airborne surveys (*e.g.* 8-day repeat coverage  
102 for Landsat data and near daily coverage for MODIS data). This enables monitoring the  
103 impact of specific climate events by investigating the consequences of drought or changes in  
104 groundwater accessibility on water use by riparian trees (Lurtz et al., 2020; Mayes et al.,  
105 2020). A key disadvantage of satellite data however, is that spatial resolution is coarse for  
106 spectral bands in the TIR region, for instance with 100 meters at-sensor resolution for Landsat  
107 8 and 9 which is then resampled afterwards to fit the 30 meters resolution of Landsat  
108 products.

109 Finer spatial resolution is available when mounting TIR sensors on airborne vectors such as  
110 an airplane or a UAV and can also be used to assess the health status of the vegetation or  
111 evapo-transpiration (Neale et al., 2011). A recent study showed that it was possible to detect  
112 genetic traits differences between cottonwood trees from different populations planted in the  
113 same plantation (Sankey et al., 2021). This suggests that airborne TIR imagery can also  
114 provide critical insights into riparian vegetation, but studies using such vectors are rare.

115 Additionally, for aircraft acquisitions that can provide the combination of spatial extent and  
116 resolution which is the most interesting to stakeholders, acquiring repeated observations over  
117 an entire phenological seasons is usually difficult. While airborne TIR images are not often  
118 used to assess vegetation health, acquisitions over river corridors are already frequent and  
119 have led to multiple publications focusing on the surface temperature of rivers (see for  
120 instance a guide aimed at practitioners (Dugdale, 2016)). Indeed, knowledge of river  
121 temperature helps target areas of ecological interest for species living in the river and helps  
122 conservation efforts. Thermal refuges are of increasingly high importance for species in

120 warming waters (Dugdale et al., 2016; Wilbur et al., 2020), and monitoring the thermal  
121 functioning of rivers provides insights to assess the impact of restoration projects (Marteau et  
122 al., 2022a). Data acquired for such studies also covers riparian vegetation but usually only  
123 focuses on the shading effects provided by trees and on their ability to lower water  
124 temperature during summer (Dugdale et al., 2018; Marteau et al., 2022b; Wawrzyniak et al.,  
125 2017; Whitley et al., 2006).

126 Therefore, thermal infrared data is available from either satellite or airborne sensors with  
127 promising potential to provide feedback about riparian vegetation health but this potential has  
128 yet to be assessed and confronted to in-field eco-physiological measurements of water stress  
129 in the riparian environment.

130 In this context, this study aims at exploring the potential of multi-scale thermal infrared data  
131 to diagnose the sensitivity of riparian forests to drought events.

132 To address those aims, we (1) tested the validity of TIR information as a good indicator of  
133 water stress by coupling field evidence from ecophysiological surveys conducted at an  
134 individual scale during summer 2022 with a simultaneous TIR acquisition during the peak of  
135 the drought period. This enabled (2) the comparison of 4 existing historical airborne TIR  
136 acquisitions to study the response sensitivity of the forest to a set of potential drought events  
137 over a 50 km reach and then (3) the replication of a similar temporal analysis using TIR  
138 satellite data in order to identify lasting changes at the corridor scale over a longer time scale  
139 (*e.g.*, 34 years).

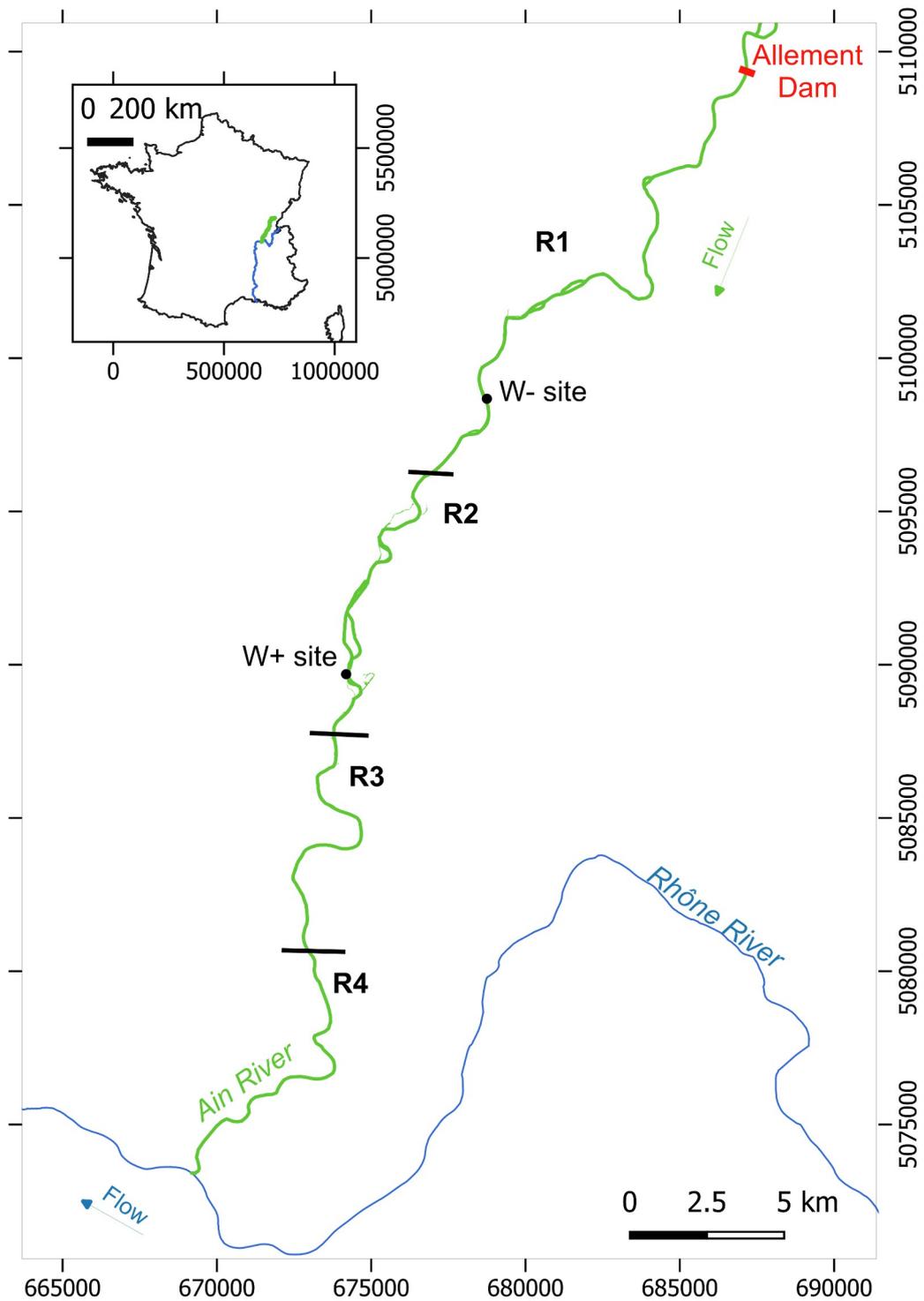
140 Our chosen study site was the riparian corridor of the Ain River because (1) it is large enough  
141 to be approached using Landsat imagery, (2) airborne TIR images have historically been  
142 acquired over the corridor to study river temperature, and (3) the forest is dominated by the

143 phreatophyte *Populus nigra* populations established on sites with contrasting connectivity to  
144 the river.

145

## 146 **2. Study Site.**

147 The Ain River is one of the main tributaries of the upper Rhône River in France. During the  
148 20<sup>th</sup> century, a chain of dams was built on this large, meandering gravel-bed river for  
149 hydroelectric purposes, causing a sediment starvation that propagates downstream by 500  
150 meters per year on average and results in a channel incision of 1 to 2 meters (Rollet, 2007;  
151 Rollet et al., 2014). We focus on a 50-km long reach that starts downstream of this chain of  
152 dams and ends at the confluence with the Rhône River, which is referred to as the lower Ain  
153 River valley [Figure 1]. The upstream section of the lower Ain River (R1) is characterized by  
154 sediment starvation and channel incision induced by the upstream dams. Downstream from  
155 this incised reach, elevation of the riverbed is stable and active channel meandering occurs  
156 and rejuvenates the riparian forest (R2). Further downstream is another reach affected by  
157 sediment starvation because of the combined effects of sediment trapping in R2 and limited  
158 local production due to lateral constraints by morainic deposits (R3). Finally, the reach of the  
159 river that leads to its confluence with the Rhône River has been historically affected by  
160 regressive incision from the Rhône River but now produces enough sediment due to channel  
161 meandering to lead to aggradation near the confluence in recent years (R4).



162 **Figure 1. - Location of the lower Ain River and of the different study reaches. The**  
 163 **coordinate system is EPSG 2154.**

164 Traditional on-field forestry surveys conducted in 2008 and 2017 by the French National  
 165 Forestry Office (ONF) (Dumas, 2017; Dumas and Perrin, 2006) showed that the dominant

166 species in the riparian forest is *Populus nigra*. These studies also suggested that sediment  
167 starvation and channel incision have an impact on the health of the riparian forest along the  
168 Ain River, with a relative loss of native pioneer species (*Populus nigra*, *Salix alba*, *Salix*  
169 *eleagnos*) in favour of exotic (*Fallopia japonica*) or post-pioneer (*Fraxinus excelsior*) species.  
170 They also mention high mortality of poplar trees in the sector near the confluence that they  
171 attribute to drought events in the early 2000s.  
172 This degraded health status of the riparian forest is visible as a downward trend in NDVI from  
173 Landsat images (Lejot et al., 2011) as well as from a study looking at the long-term impacts of  
174 channel incision by coupling the data from the ONF field surveys with LiDAR and  
175 hyperspectral datasets (Godfroy et al., 2023). This later study highlighted that in reach R1, the  
176 relative elevation of the riparian forest from the base flow waterline is higher than in reach R2  
177 which results in an increased distance to the groundwater table and a dryer environment. The  
178 Ain River is therefore an interesting site to address our aims because the known changes in  
179 forest composition, structure, and reflectance in the visible and near-infrared spectrum based  
180 on forest stationary conditions suggests that water stress occurs during the growing period and  
181 that some poplar trees are known to have responded to a dryer environment.

182

### 183 **3. Materials & Methods.**

#### 184 3.1. Forestry data used for field validation.

185 Fieldwork was conducted approximately every two weeks between May 5<sup>th</sup> and September  
186 27<sup>th</sup> 2022 (Table A). It focused on two sampling sites with differing geomorphic conditions : a  
187 site on the R2 reach where poplar trees were assumed to be well-connected to the river system  
188 (W+), and a site on the R1 reach where poplars were assumed to be disconnected from the

189 river and to have more limited access to groundwater (W-). Site selection was based on  
190 previous work on the riparian forest by Dufour (2005) and Godfroy et al. (2023).

191 For each site, ten *Populus nigra* tree were selected, their diameter at breast height was  
192 measured, and they were sampled during multiple campaigns at midday, near the solar zenith.

193

### 194 3.1.2. Leaf water potential.

195 For each campaign, two green leaf shoots were collected for each of the trees and enclosed  
196 in aluminium sheets to maintain dark conditions until measurements in the lab. Then, shoots  
197 were kept in a cooling box for conservation before measuring leaf water potential (LWP)  
198 using a pressure chamber once back at the laboratory (Scholander et al., 1965). The leaf petiol  
199 was cut with a sharp blade and inserted in the pressure chamber. The mean pressure of the two  
200 selected leaves per tree was used as, a single replicate.

201

### 202 3.1.2. Phloem collection and iWUE calculation.

203 Phloem of the trees was sampled every other field campaigns and used to estimate the  
204 intrinsic water use efficiency (iWUE) of each tree according to the protocol published by  
205 Gerle et al. (2023), based on relative ratios in carbon isotopes in the phloem content (Vernay  
206 et al., 2020).

207 Briefly, phloem was collected with a bark-corer (9mm) at breast height in the trunk of each  
208 tree and immersed in 1.5 mL exudation solution (15 mM polyphosphate buffer: sodium  
209 hexametaphosphate, Sigma, München, Germany). Phloem was then removed from the tube  
210 before freezing the solution. The samples were then freeze-dried and rehydrated with  
211 deionised water in a tin capsule. After drying in a oven, the  $^{13}\text{C}/^{12}\text{C}$  phloem ratio was analyzed  
212 with a spectrometer and expressed in ‰ relative to Vienna Pee Dee Belemnite (VPDB).

213 Calculations to obtain iWUE are detailed in Gerle et al. (2023) and Vernay et al. (2020).

214

215 3.2. Remote-sensing data.

216 *3.2.1. Airborne TIR images.*

217 Images in the TIR spectrum (7.5–14  $\mu\text{m}$ ) were acquired in summer over four campaigns since  
218 2010 [Table 2] with sensors mounted on an ultralight aircraft (2010 and 2022) or a helicopter  
219 (2011 and 2014). Three different sensors were used: a Thermo Tracer TH7800 for the first  
220 flight, a VarioCAM hr head for the second flight and a VarioCAM hr research 600 for the  
221 third and fourth flights. These cameras can detect temperature differences of around 0.1°C  
222 within an image. Days of flight were chosen based on the weather and during low-flow  
223 conditions.

224 **Table 2 – Airborne TIR campaigns used in this study.**

<b>Date</b>	<b>Time</b>	<b>Spatial resolution (m)</b>	<b>Mean daily discharge (<math>\text{m}^3\cdot\text{s}^{-1}</math>)</b>	<b>Daily maximum temperature (<math>^{\circ}\text{C}</math>)</b>
July 30 <sup>th</sup> 2010	15:00 to 16:00	1.50	14.3	25.6
June 28 <sup>th</sup> 2011	17:45 to 18:30	0.70	13.4	34.0
July 3 <sup>rd</sup> 2014	18:00 to 19:00	0.60	14.1	30.2
July 19 <sup>th</sup> 2022	12:20 to 13:30	0.35	16.2	37.1

225

226 The images from the first three campaigns are historical TIR data available on the Ain River  
227 that originally aimed at studying river temperature to identify the inter-annual variability of  
228 cold-water patches. Therefore, the processing of these images is documented by the authors of  
229 the initial study (Wawrzyniak et al., 2016). The last airborne campaign was coordinated with

230 the field campaign of this study (July 19<sup>th</sup>) and the flight took place during the same time-  
231 frame.

232

### 233 *3.2.2. Satellite TIR images.*

234 Landsat satellites were used as a source of coarse historical TIR data starting from 1990 by  
235 using level 2 images from Landsat 5, 8 and 9. Images from Landsat 7 were not considered for  
236 the analysis due to the Scan Line Corrector failure that occurred in 2003. At most one image  
237 was selected per year from the available data based on a rule of precipitation during the  
238 summer period (a six-day window with no precipitation). This rule was defined according to  
239 results from preliminary analysis of airborne TIR data that showed the difficulty of assessing  
240 vegetation water stress from TIR imagery with wet antecedent conditions. This ensured that  
241 selected images were from the driest possible period for every summer.

242

### 243 3.3. Additional information for geo-referencing, selecting and characterizing the study area.

244 Historical aerial photographs produced by the French Institut National de l'Information  
245 Géographique et Forestière (IGN) were used to provide information about land cover for dates  
246 close to the TIR campaigns. They were originally used by Wawrzyniak et al. (2016) to help  
247 manually geo-reference the TIR images. Aerial color images were also acquired during the  
248 2022 campaign by mounting a camera (Nikon Z6, 35 mm lens) on the aircraft during the TIR  
249 acquisition and were similarly used to assess land cover in 2022 and to help with the geo-  
250 referencing process.

251 Topo-bathymetric LiDAR data was acquired during August 2015 and covers the upstream half  
252 of the study reach ( $\approx 20$  km), initially to study river bathymetry (Lague and Feldmann, 2020).  
253 This dataset was then used to characterize the stationary conditions of the riparian forest in

254 Godfroy et al. (2023). It is used in this study to help assess the changes in canopy temperature  
255 based on forest stationary conditions.

256 The data was acquired with an Optech Titan sensor flown on an airplane, and resulted in the  
257 acquisition of a point cloud with a final density of around 18.6 points per meter square for  
258 each of the two lasers of the sensor (Green for bathymetry and NIR for above-ground) and  
259 with a mean vertical accuracy of  $\approx 10$  cm.

260 Vegetation surveys were conducted by ONF in 2007 and 2017 at the request of local  
261 stakeholders. It led to the survey of *ca.* 1200 forest plots in the study reach with the goal of  
262 providing extensive information on species distribution and health in the riparian forest of the  
263 Ain River and, as such, one plot was assessed per hectare of forest. These vegetation plots  
264 were used to provide an extensive analysis grid common to the forest surveys and to previous  
265 studies (Godfroy et al., 2023) that covers the lower Ain River corridor. That grid is used for  
266 selecting forest patches within which Landsat TIR information is analyzed.

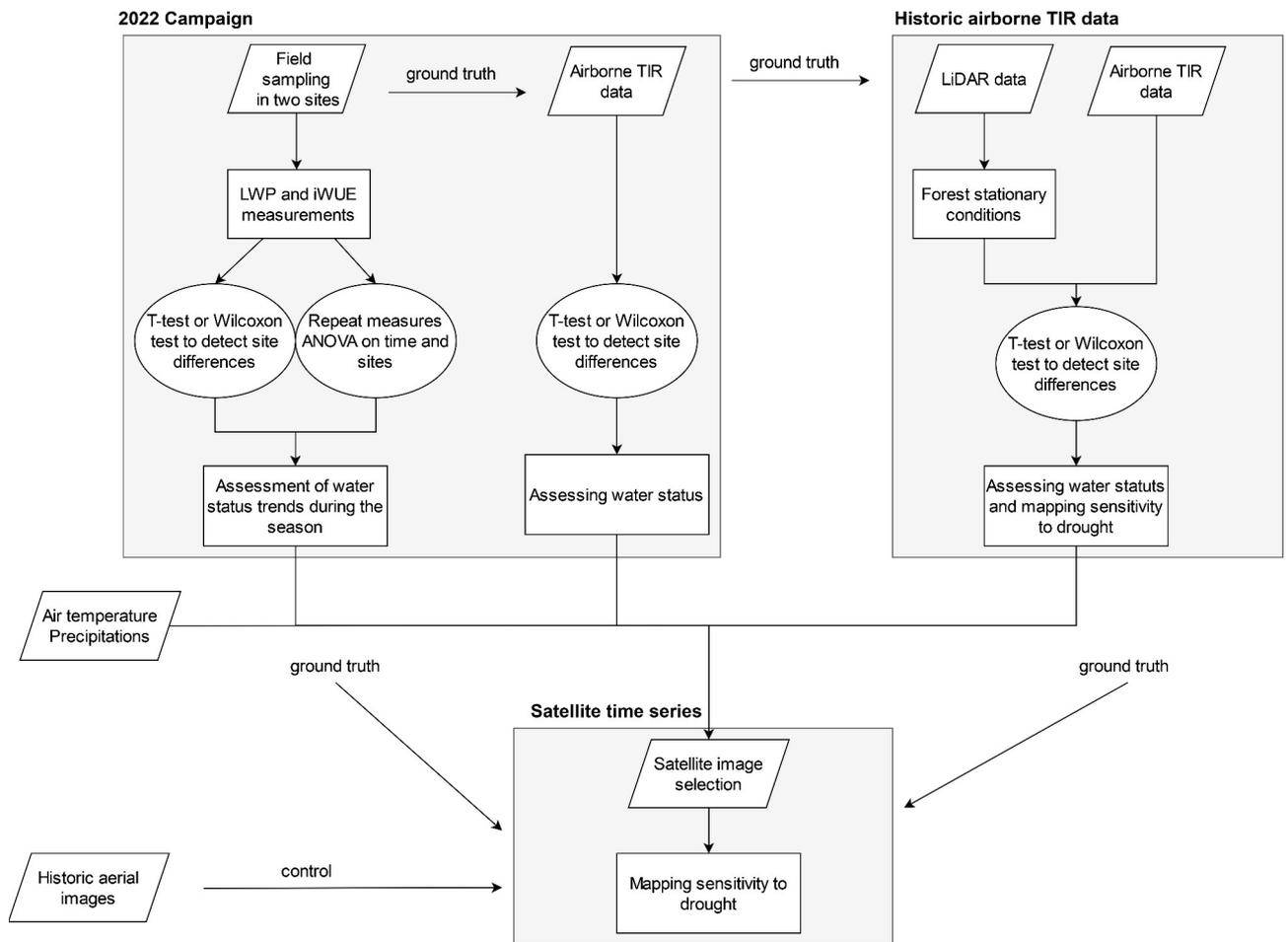
267 Information about daily temperature and precipitations near the study site was accessed from  
268 the open access archives of the Ambérieu-en-Bugey meteorological station available from the  
269 Global Surface Summary of the Day (GSSD) provided by the National Centers for  
270 Environmental Information (cf. <https://www.ncei.noaa.gov/metadata/geoportal/>). Information  
271 on the flow level of the Ain River was measured at the hydrological station located at Pont  
272 d'Ain and provided by the French Ministry of Ecological Transition (cf.  
273 <https://www.hydro.eaufrance.fr/>).

274

### 275 3.4. General workflow.

276 Data analysis first focused on the campaign conducted during summer 2022 to investigate  
277 how the water status of *Populus nigra* differed between the two study sites [Figure 2]. Water

278 status was assessed by monitoring changes in leaf water potential and intrinsic water use  
 279 efficiency from field samples. Differences in LWP or iWUE between poplars and between  
 280 sites were then assessed by running Student's test or Wilcoxon Mann-Whitney test depending  
 281 on data distribution for each date. Differences between each week were assessed by using a  
 282 pairwise t-test. Repeated measures ANOVA were then conducted to test the effects of  
 283 environment (W+ and W-) and time on LWP and iWUE values (environment, time and their  
 284 interaction were used as explanatory variables in the ANOVAs). Meteorological data from the  
 285 start of the campaign to its end were then plotted to recontextualize the observed trends in  
 286 water stress with the trends in air temperature and precipitations of summer 2022.



287  
 288 **Figure 2 – General workflow of our study.**

289 TIR data from 2022 was then analyzed by first looking at differences in tree crown  
290 temperature between poplars from the two sites. In order to retrieve tree crown temperature,  
291 the shape of each tree crown was first delineated manually by combining GPS positions of  
292 each tree with the aerial images synchronous to the TIR acquisition and canopy height  
293 extracted from the LiDAR dataset. Median temperature was then extracted for each tree  
294 crown from TIR data. Differences between the poplars on the two sites were then assessed  
295 using the same tests as for LWP and iWUE.

296 Tree diameter from the field survey and tree height from the LiDAR data were tested to  
297 predict LWP, iWUE and TIR as potential variables to explain differences between individuals  
298 on a given site for all dates during summer 2022 for which data was available.

299 Data analysis then focused on using existing airborne TIR imagery to detect water stress and  
300 map the sensitivity of the riparian forest to drought. The relationship between canopy  
301 temperature and stationary conditions at the level of forest plots was investigated for all four  
302 campaigns.

303 In order to retrieve information on canopy temperature for each forestry plots, the closest  
304 aerial images available to each campaign were used to screen forest plots and mask areas  
305 which were not vegetated. The D90 of canopy temperatures was then extracted in each plot in  
306 order to minimize effects from shadowing.

307 Stationary conditions were assessed by detrending LiDAR-derived DEM using the Fluvial  
308 Corridor Toolbox (Roux et al., 2015). This detrended DEM was obtained by subtracting the  
309 elevation of the water level of the river from the elevation of the terrestrial floodplain level,  
310 resulting in values of elevation relative to the water level under low flow conditions ( $Q = 16$   
311  $\text{m}^3.\text{s}^{-1}$ ). The mean value from the detrended DEM was then extracted for all forest plots in the  
312 coverage of the LiDAR data and used to create two classes of stationary conditions based on

313 previous literature on the Ain River (Dufour and Piégay, 2008; Godfroy et al., 2023), with a  
314 "good connectivity" class for plots lower than 2.5 meters above low flow and a  
315 "disconnected" class for plots 2.5 meters above low flow and higher.

316 The same methodology as for the analysis of the 2022 data was then applied by running  
317 Student's test or Wilcoxon Mann-Whitney test depending on data distribution for each  
318 campaign in order to detect differences in canopy temperature due to stationary conditions.

319 The campaign from 2022 was used as a control of the expected response in temperature  
320 canopy under known stress conditions, and hydrological and meteorological data for the week  
321 before the campaign were plotted to contextualize the observations.

322 Maps of riparian forest sensitivity to drought were then produced using relevant campaigns.

323 In order for maps to be comparable despite differences in atmospheric conditions at the time  
324 of survey, canopy temperature values were discretized using quartile statistics.

325 Similar maps were then created using Landsat data to highlight lasting changes since 1990.

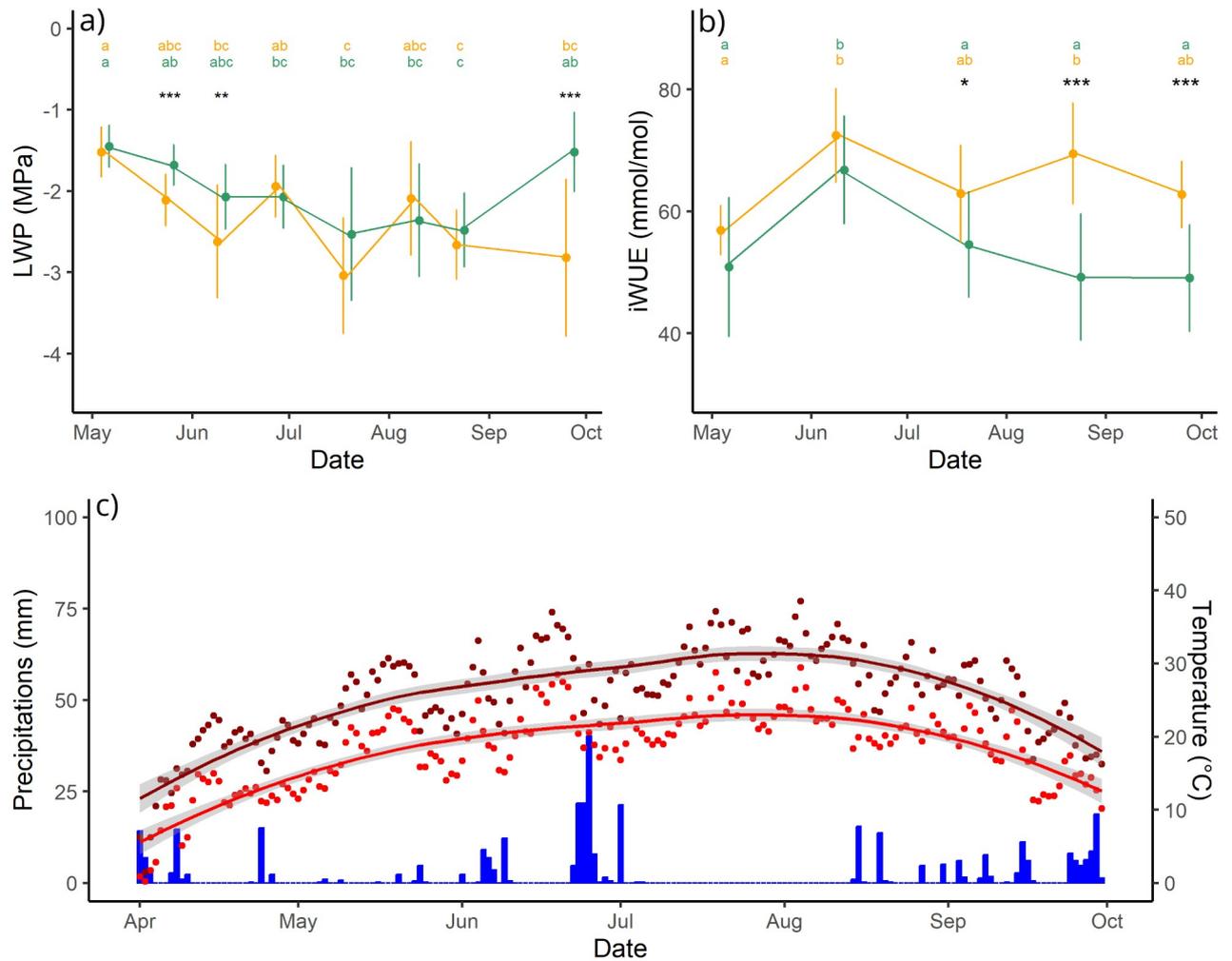
326 Image selection is described in the Satellite TIR images section of the manuscript. At-sensor  
327 resolution (100–120 meters) was higher than the spatial footprint of the forest plots (20 meters  
328 radius, with 100 meters between plot centers) so mean temperature was extracted for each plot  
329 and a map was produced to help assess and discuss the impact of the original footprint of the  
330 sensor on the results by using available orthophotos.

331

## 332 **4. Results.**

### 333 4.1. Field validation of tree water stress and TIR response.

334 Field surveys show that the water status of poplar trees varies during the summer period. In  
335 particular, both sites reached stress conditions during summer since LWP values were  
336 generally lower than -1.75 MPa from June to September [Figure 3a].



337

338 **Figure 3. - (a) Leaf water potential (LWP) and (b) intrinsic water use efficiency (iWUE)**

339 **on the W+ site (green) and W- site (orange), and (c) precipitations (blue) and daily mean**

340 **(red) and daily maximum (dark red) temperatures for the study period.** Stars indicate

341 differences in LWP or iWUE between the two sites (Wilcoxon or Student tests depending on data distribution, \*

342 =  $p < 0.05$ , \*\* =  $p < 0.001$ , \*\*\* =  $p < 0.001$ ) and letters indicate the differences between dates from pairwise t

343 tests.

344 LWP values at the connected site (W+, in green) decreased until reaching a plateau near -2.5

345 MPa starting from the middle of June before increasing back to -1.5 MPa at the end of

346 September. For the disconnected site (W-, in orange), LWP decreased faster than for the

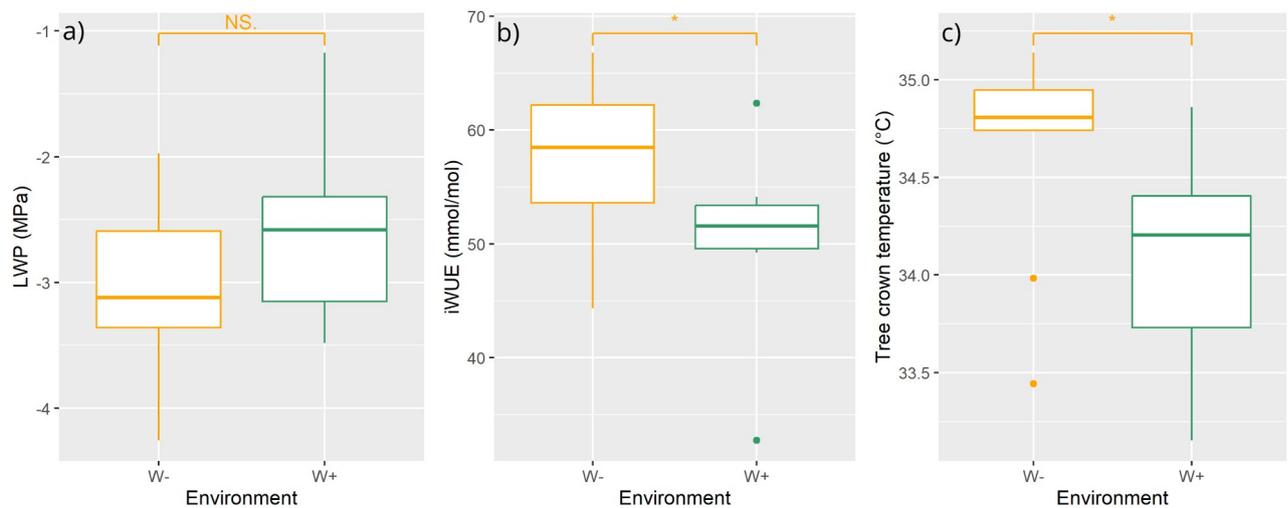
347 connected site, reaching values near -3 MPa on average in July. A more contrasted recovery is

348 observed for these poplars since LWP values at the end of September were still around -3  
349 MPa on average, but variability between individuals increased which suggests partial  
350 recovery. In addition, LWP values were more variable for poplars on the disconnected site  
351 with short recovery periods (June 28th and August 9th) where mean LWP values were closer  
352 to that of the connected site. Differences between sites were most significant at the beginning  
353 of the stress period when LWP values increased with each subsequent campaign (May 25<sup>th</sup>  
354 and June 10<sup>th</sup>) and at the end of September (September 26<sup>th</sup>) after which the connected site  
355 fully benefits from the recovery period.

356 Although iWUE measurements were sparser and notably not conducted on June 28<sup>th</sup> or  
357 August 9<sup>th</sup> (dates for which LWP values fluctuated for the disconnected site), they show a  
358 trend similar to LWP measurements [Figure 3b]. An increase in iWUE values was observed  
359 for both sites during summer, from 40–45 mmol/mol and reaching values near 50 mmol/mol  
360 and 60 mmol/mol on average for the connected and disconnected sites respectively. A  
361 recovery period is identified at the end of summer, with decreasing iWUE values starting  
362 August 23<sup>rd</sup> for the connected site and at the end of September for the disconnected site.  
363 While iWUE values were on average higher for poplars in the disconnected site, differences  
364 between the two sites were significant during summer (July 19<sup>th</sup> and August 23<sup>rd</sup>) and at the  
365 end of September.

366 The ANOVA tests performed on repeated measures indicated a significant effect of the  
367 environment (W+ vs. W-,  $p = 0.017$ ,  $F = 8.583$ ), of the week of the sampling ( $p < 10^{-3}$ ,  $F =$   
368  $10.034$ ) and of the environment  $\times$  week interaction ( $p < 10^{-3}$ ,  $F = 4.413$ ) on LWP. For iWUE,  
369 only an effect of the week ( $p < 10^{-3}$ ,  $F = 15.625$ ) and of the environment  $\times$  week interaction ( $p$   
370  $< 10^{-3}$ ,  $F = 3.412$ ) were significant, but not the effect of the environment alone ( $p = 0.099$ ,  $F =$   
371  $4.091$ ).

372 Those two indicators converge towards an increasing water stress in *Populus nigra* during  
 373 summer and a recovery in late August or during September, which follows temperature and  
 374 rainfall trends for summer 2022 [Figure 3c].  
 375 Tree sampling and TIR data were acquired simultaneously on July 19<sup>th</sup>, which means all three  
 376 indicators of water stress can be compared for the disconnected (W-, in orange) and connected  
 377 (W+, in green) sites for the same date. On this date, LWP values were lower in the  
 378 disconnected site than in the connected site (Figure 4a), and the opposite was true for iWUE  
 379 values (Figure 4b). However these differences were only significant for iWUE ( $p < 0.05$ ).  
 380 This validates a higher stress status in the disconnected site. Canopy temperature of sampled  
 381 trees was higher in the disconnected site ( $p < 0.05$ ) which also suggests higher water stress for  
 382 poplars in the disconnected site (Figure 4c).



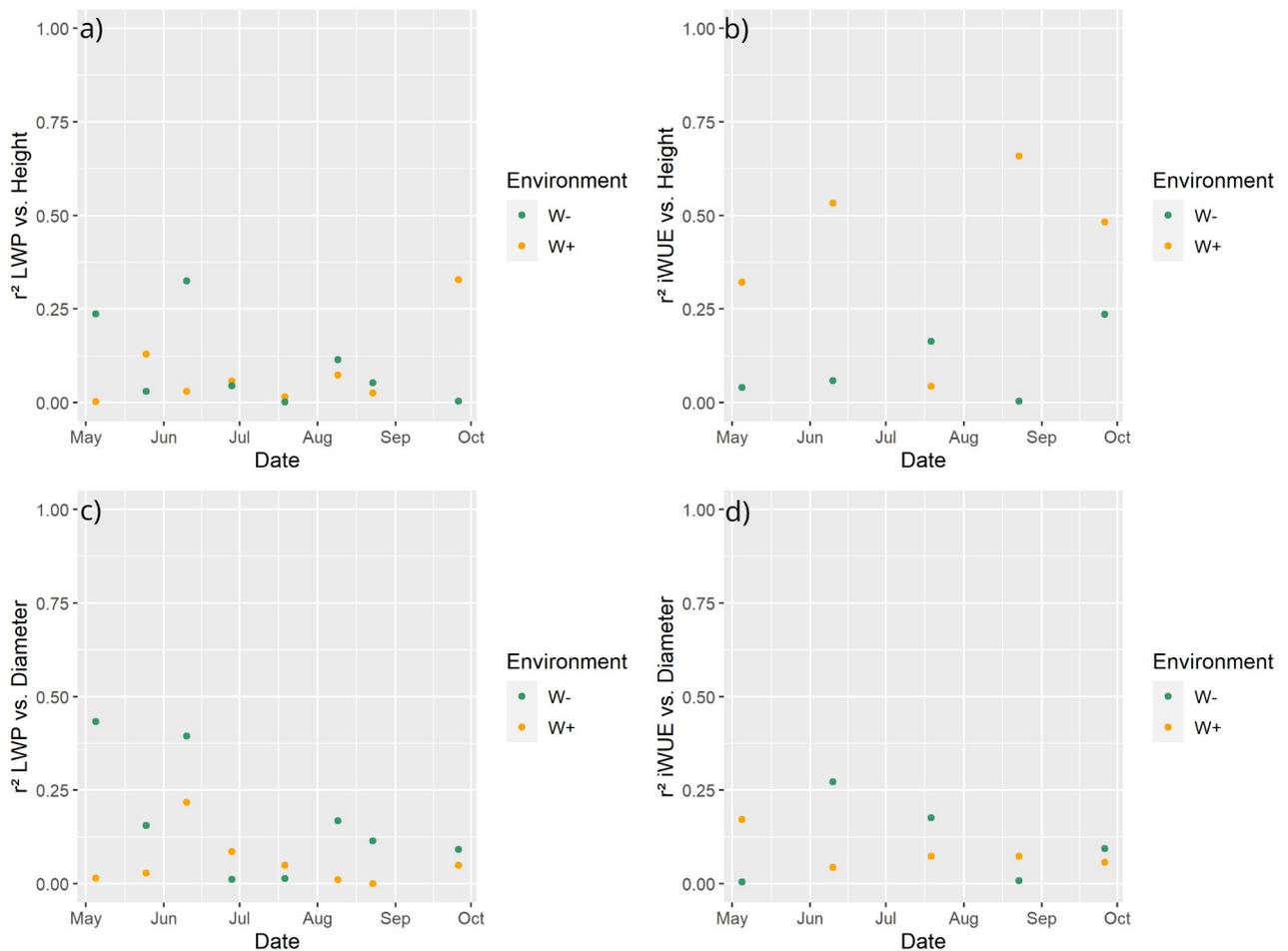
383

384 **Figure 4. - (a) Leaf water potential, (b) intrinsic water use efficiency and (c) tree crown**  
 385 **temperature depending on the study site on July 19<sup>th</sup> 2022.** Stars indicate differences between the  
 386 sites (Wilcoxon or Student tests depending on data distribution, \* =  $p < 0.05$ , \*\* =  $p < 0.001$ , \*\*\* =  $p < 0.001$ ).

387

388 The relationship between tree physiognomy and the three indicators of water stress was also  
 389 investigated (Figure 5). While  $r^2$  values were low for most of the dates, intra-seasonal

390 variability was also recorded. Higher  $r^2$  values were reached for LWP at the beginning of  
 391 summer for trees in the disconnected site (from 0.25 to 0.50) and for iWUE at the beginning  
 392 and the end of the summer for trees in the connected site (from 0.25 to 0.70). The relationship  
 393 between canopy temperature and tree height or tree diameter also reached high  $r^2$  values for  
 394 the trees at the disconnected site (0.55 and 0.60 respectively). Overall, when a high  $r^2$  value  
 395 was recorded, bigger trees (higher diameter or height) were less stressed (higher LWP, lower  
 396 iWUE and lower canopy temperatures) than smaller ones. Tree diameter was better correlated  
 397 than tree height for the disconnected site while the opposite was true for the connected site.

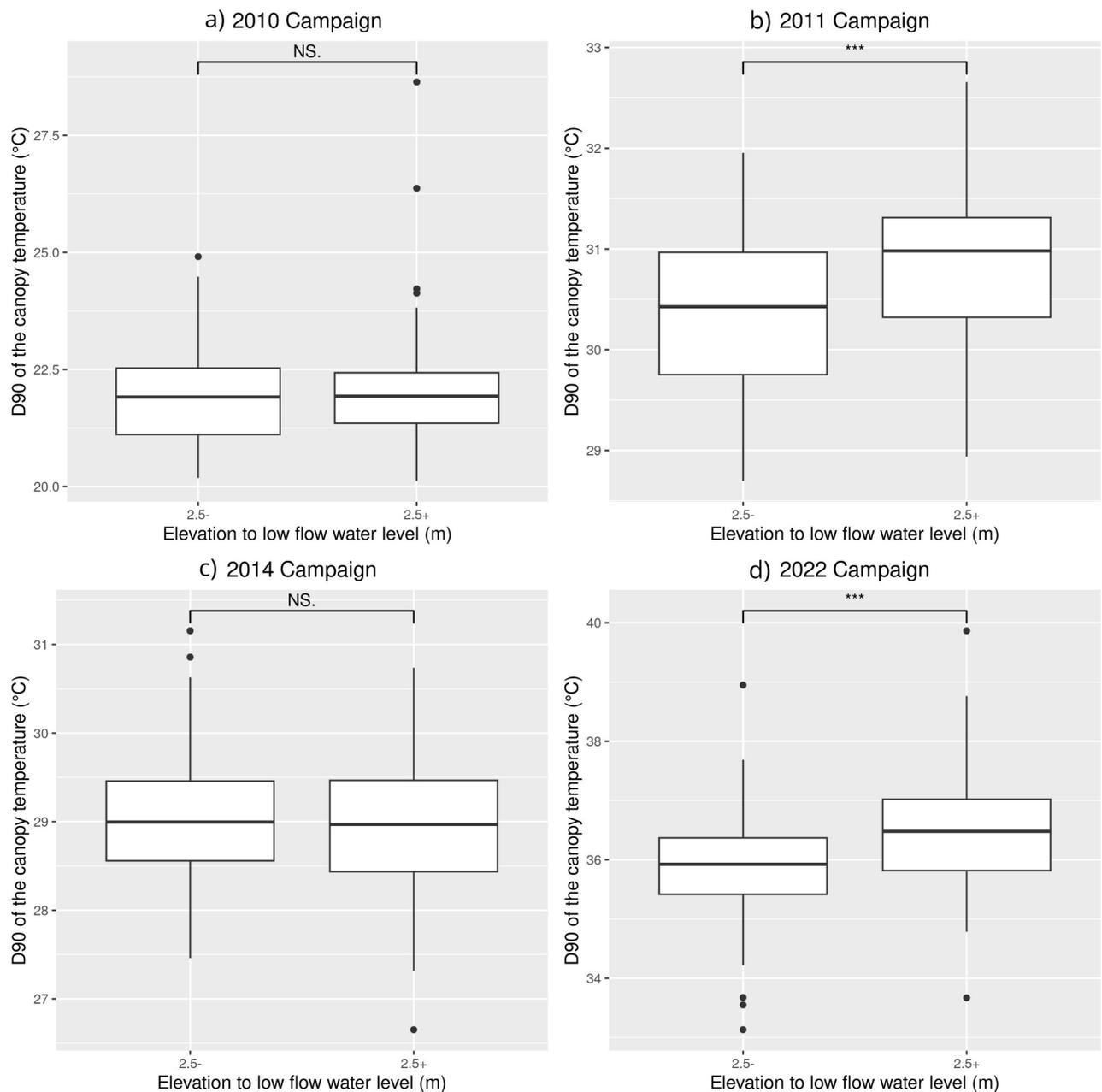


398  
 399 **Figure 5. -  $r^2$  of the linear relationship between (a) LWP and tree height, (b) iWUE and**  
 400 **tree height, (c) LWP and tree diameter and (d) iWUE and tree diameter for each**  
 401 **sampling date and site.**

402

403 4.2. Inter-annual differences in airborne TIR response to summer conditions.

404 In the absence of *in-situ* assessment of the water status of the riparian forest for the historical  
405 airborne surveys, only the thermal response of riparian vegetation to differences in  
406 connectivity can be assessed. Using a breakpoint value of 2.5 meters above the low flow  
407 water level as the main variable to consider shifts in connectivity, the results from an analysis  
408 at the plot level in 2022 were similar to those of the analysis at the level of individual tree  
409 crowns [Figure 6].



410

411 **Figure 6. Canopy temperature at the plot level depending on plot elevation to low flow**  
 412 **water level for all four airborne TIR campaigns: a) 2010, b) 2011, c) 2014 and d) 2022.**

413 Stars indicate the differences between plots (results from Wilcoxon or Student tests depending on data  
 414 distribution \* =  $p < 0.05$ , \*\* =  $p < 0.001$ , \*\*\* =  $p < 0.001$ ).

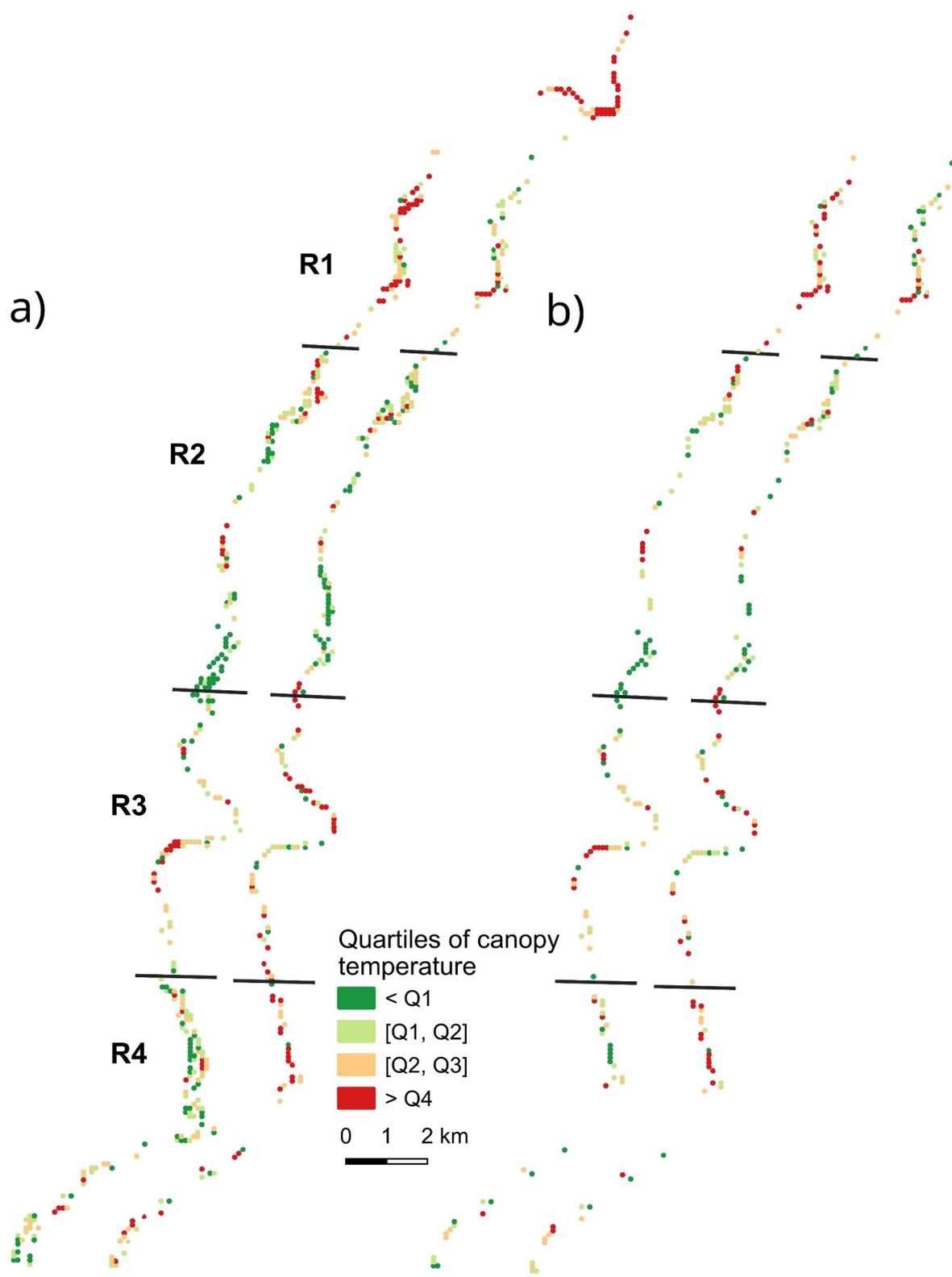
415 Indeed, in 2022 [Figure 6d] canopy temperature were higher for plots with higher elevation to  
 416 the water table ( $p < 0.001$ ) which is similar to the higher crown temperatures for trees in the  
 417 disconnected site. While this was also true for the 2011 [Figure 6b] campaign ( $p$ -value <

418 0.001), there were no differences in canopy temperature between the two classes of stationary  
419 conditions (connected for plots where elevation to low flow is < 2.5 meters and disconnected  
420 for plots where it is > 2.5 meters) for 2010 [Figure 6a] and 2014 [Figure 6c]. These results  
421 show temporal variability in the thermal response of riparian vegetation to its stationary  
422 conditions.

423

#### 424 4.3. Mapping of airborne TIR responses to summer droughts: 2011 vs. 2022.

425 The statistical distribution of canopy temperatures was mapped for 2011 and 2022 using  
426 quartiles [Figure 7a], and a confrontation of the two maps (keeping only forest plots for which  
427 thermal information was available in both 2011 and 2022) is shown [Figure 7b].



428

429 **Figure 7. - (a) Spatial distribution of the different quartiles of canopy temperature in**  
430 **2011 and 2022 for all ONF plots, and (b) a 2011-2022 comparison using only plots for**  
431 **which data was available both in 2011 and 2022.**

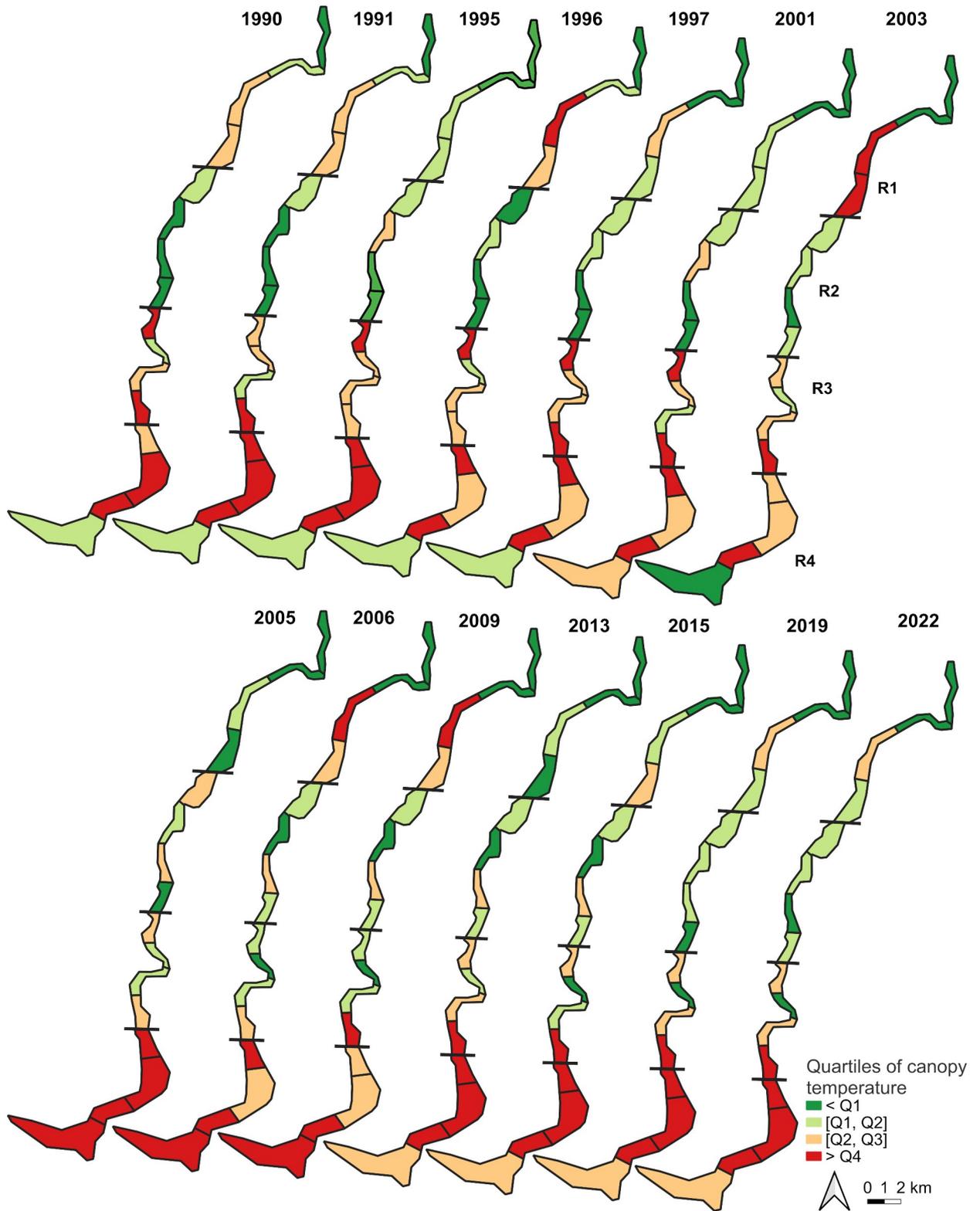
432 Both acquisitions covered most of the riparian corridor near the main channel of the Ain River  
433 and poorly covered larger areas of riparian forest that were created by river meandering. The  
434 higher canopy temperatures were concentrated in the R1 and R3 reaches, while the lower  
435 canopy temperatures are located in the R2 and R4 reaches. As R1 was the most  
436 geomorphologically-degraded reach suffering from historic channel incision and R2 was the  
437 most geomorphologically-stable reach the spatial distribution of canopy temperature suggests  
438 that the health of the riparian forest and its sensitivity to drought reflect 20th century changes  
439 in geomorphology.

440 The distribution of canopy temperatures in 2022 was different from the one in 2011 with  
441 lower quartiles of canopy temperature located in the R2 reach and higher quartiles being  
442 located in the R3 and R4 reaches. Changes mainly occurred in the R1 and R4 reaches.

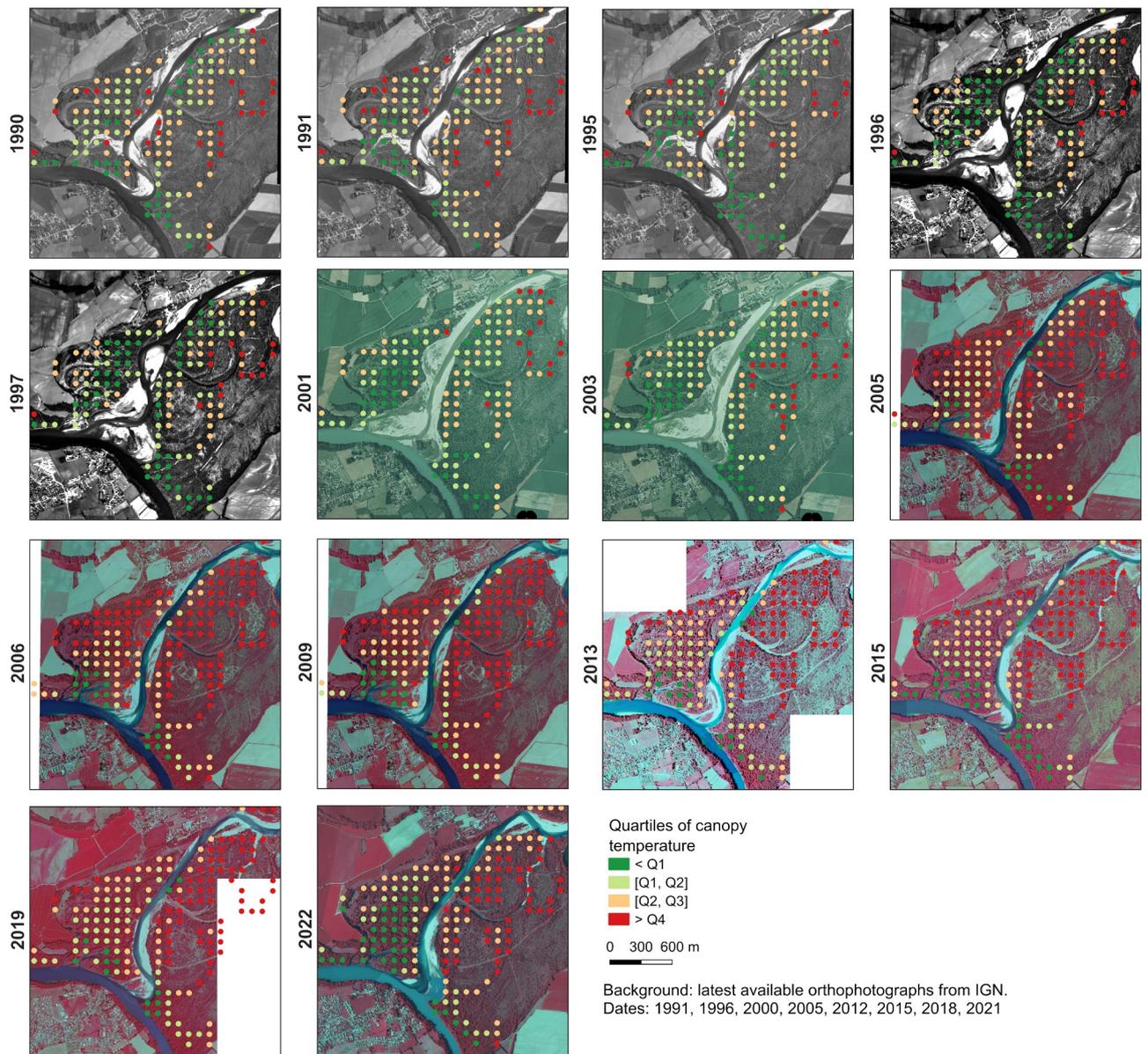
443

#### 444 4.4. Mapping large-scale riparian thermal changes using Landsat archives.

445 Based on the results from field validation, selection of TIR Landsat data was constrained by  
446 precipitation (no rain during a six-day period preceding observations from 2011 and 2022)  
447 and by cloud cover (visually inspecting each image). With one image selected per year at a  
448 maximum, this resulted in only 14 images selected for the 1990–2022 period. The resulting  
449 maps also compared the distribution of temperature quartiles for each year but used all  
450 available forest plots due to the higher spatial extent of satellite data [Figure 8].



452 **Figure 8. Spatial distribution of canopy temperature quartiles at the corridor scale from**  
453 **Landsat data between 1990 and 2022.** Each reach was subdivided in four sectors of similar river length  
454 and the information for each forest plot was summarized for each sector to facilitate reading.  
455 The spatial distribution of canopy temperature quartiles was very similar for each year  
456 satellite data was available. Plots showing higher temperature were mainly distributed in the  
457 R1, R3 and R4 reaches, with most of the plots in the R2 reach being consistently among the  
458 coolest plots. Lower temperatures also feature plots near the confluence with the Rhône River,  
459 in the middle section of the R3 reach since the start of the new century, and plots near the  
460 main channel or in the upstream-most section of reach R1.  
461 Although the resolution of satellite thermal data is coarse, strong and meaningful change is  
462 still highlighted in the area near the confluence with the Rhône River that is of interest to  
463 understand forest dynamics in regards to meteorological and geomorphic constraints [Figure  
464 9].



465

466 **Figure 9. - Evolution of the sector at the confluence with the Rhône River since 1990.**

467 On the aerial images, a simplification of channel morphology towards a lower active channel  
 468 width is visible between 1995 and 2001 and illustrates the geomorphic changes that occurred  
 469 at the end of the century in this section of the lower Ain River valley. In the case of the  
 470 riparian forest, canopy temperature progressively increased relative to the rest of the lower  
 471 Ain River valley between 2000 and 2005 leading to a shift from lower quartiles of

472 temperature to higher quartiles of temperature. This shift is more pronounced on the left side  
473 of the river as the right side appears to have recovered by 2022.

474

## 475 **5. Discussion.**

### 476 5.1. Summer water stress conditions of poplar trees depend on water accessibility.

477 Our results suggest that the water stress conditions of riparian poplars depend on water  
478 accessibility. Differences in the health status of trees were observed for all indicators (tree  
479 crown temperature, LWP, iWUE), showing the contrasted response of *Populus nigra* to  
480 drought based on its stationary conditions. The trees from the dryer site were consistently  
481 showing higher water stress on one or more of the indicators used in this study, compared to  
482 trees from the wetter site.

483 When considering a stress threshold at -1.75 MPa value for the leaf water potential, as  
484 reported for *Populus nigra* stomatal closure in previous studies along the Drac and Isère rivers  
485 (Lambs et al., 2006), stress conditions were reached for poplar individuals in summer 2022  
486 regardless of their stationary conditions. While trees in more connected reaches of the river  
487 were displaying lower stress according to the indicators (higher LWP, lower iWUE and lower  
488 canopy temperature), they still experienced water scarcity. Therefore, the threat of drought-  
489 induced changes in riparian forests can even affect river reaches with good geomorphic  
490 conditions, and be exacerbated in river reaches where anthropogenic activity has already  
491 historically altered water accessibility.

492 Such a change following a drought event was recorded in this study. Long-term change  
493 analysis based on Landsat archives showed a shift in the riparian forest health status near the  
494 confluence with the Rhône River. The timing of this change corresponds to the hottest and  
495 driest summer recorded in France: the 2003 drought (Black et al., 2004; Luterbacher et al.,

496 2004). The fact that satellite data was able to detect such a change despite its coarse pixel  
497 resolution suggests that the shift in forest structure is significant, with a more open canopy,  
498 and a temperature measurement at sensor mixing the signals from the canopy, the understory  
499 and potentially even the forest ground. This would be consistent with the field surveys by  
500 ONF noticing a higher number of dead poplars in this sector and that they attributed to that  
501 summer 2003 drought. Similar vegetation dieback has been recorded using remote sensing  
502 data in other river systems following drought events (e.g. Kibler et al., 2021).

503 Tree response to stress however may vary. While tree mortality and the decline of the poplar  
504 forest is one possible response to stress, it is not the only one. Water scarcity can also impact  
505 tree growth and lead to changes in productivity of riparian trees (Lambs et al., 2006; Monclus  
506 et al., 2006; Smith et al., 1991) and therefore impact both the ability of trees to act as carbon  
507 sinks and their structure (e.g. diameter, height, leaf size). Specifically, for *Populus nigra*,  
508 plasticity has been shown across a range of climatic conditions in Europe, with individuals  
509 from certain locations (e.g. Spain) being more adapted to drier environments by having slower  
510 growth with smaller leaves and a faster stomatal closure (Viger et al., 2016).

511 In our study, the relationships between tree height or diameter and several indicators were  
512 tested but were only significant for a few dates. Trees on the disconnected site with a larger  
513 diameter had higher LWP at the beginning of the stress period and lower canopy temperatures  
514 during the TIR acquisition. On the other hand, trees from the connected site with a higher tree  
515 height had a better recovery at the end of the summer. Interestingly, the allometric relationship  
516 between tree height and tree diameter was distinct for the two sites (see Figure B). This  
517 highlights how water scarcity, albeit due to channel incision (Godfroy et al., 2023; Rollet et  
518 al., 2014), has already affected tree physiognomy in our study reach and may explain the  
519 observed differences in tree response along the height – diameter gradient between sites.

520 Indeed, smaller trees are similar but differences in height-to-diameter ratios are observed  
521 when trees grow larger.

522 In addition, not all historic airborne TIR data were collected under stressful conditions, as  
523 only two campaigns showed changes in canopy temperature based on the vertical connectivity  
524 of forest plot. Lower temperatures and higher precipitations before the two other campaigns  
525 (Figure C) may not have led to stress conditions. Although our number of observations was  
526 too low to be significant, the meteorological conditions in a three-day time period before the  
527 campaign were strongly correlated to TIR-derived stress indicators (Figure D). This is also  
528 supported by the intra-seasonal variability of the stress signal as measured by the LWP of the  
529 poplar trees that showed a partial recovery for dates sampled following rainfall events.

530 This shows that identifying the role played by water accessibility in the response of riparian  
531 forests to drought is difficult because it related to both river-related dynamics (such as access  
532 to groundwater following channel incision or depending on flows during summer) and  
533 meteorological dynamics (such as precipitations temporarily replenishing water content). In  
534 addition, while we studied *Populus nigra* as it is a phreatophyte species, other species in the  
535 riparian forest such as *Fraxinus excelsior* do not have the same root systems and are limited to  
536 the vadose zone for accessing water resources (Dufour, 2005).

537

## 538 5.2. The benefits of a multi-tool approach to understand tree response to water stress.

539 Our results also shed light on the tools available to assess and monitor water stress in riparian  
540 forests. One of our main findings in regards with the study design is that all our indicators  
541 were sensitive to acquisition timing and did not always detect differences in stress intensity  
542 based on forest stationary conditions, making it difficult to assess the stress status of the forest  
543 or individual trees.

544 In-field sampling was used successfully to detect the occurrence of water stress from May to  
545 September and both LWP and iWUE showed an expected increase in stress intensity during  
546 the summer. On the other hand, while both indicators detected differences between  
547 individuals from the two sites, the dates for which results were the most significant did not  
548 match except during the recovery period at the end of September.

549 Similarly, using airborne TIR data did not yield relevant results for all of the dates data was  
550 available. Due to a lack of historic *in-situ* assessment of tree water status, it is not possible to  
551 know if stress conditions would have been reported using another indicator. However, these  
552 results show that the ideal acquisition window for studying vegetation health using thermal  
553 data is not the same as the ideal acquisition window to study river temperature (*i.e.* the aim of  
554 the initial historical study): data to study interactions between the river and groundwater  
555 (Wawrzyniak et al., 2016) does not always provide accurate information on vegetation status.  
556 However, it also means that having such wider implications in mind when planning data  
557 collection can help optimise efforts and resources in making data relevant beyond its original  
558 use.

559 Considering ground truth, a careful study design based on antecedent hydro-climatic  
560 conditions and a deep knowledge of geomorphic conditions in a river corridor seem crucial to  
561 studies seeking to employ airborne TIR imagery to assess water stress.

562 However, some level of variability is to be expected since different indicators of water stress  
563 do not correspond to the same biophysical processes and might not share the same response  
564 time to environmental variables (Volaire, 2018), and water stress progresses from the upper  
565 shoots in the tree crown to the tree trunk and the roots (Lambs et al., 2006).

566 For example, stomatal closure, which has been linked to changes in transpiration and an  
567 increase in leaf temperature (Hsiao and Acevedo, 1974; Jones, 1999), occurs slowly over a

568 range of LWP depending on the rate of stress (Jones and Rawson, 1979). In *Populus nigra*,  
569 higher rates of stomatal closure were related to higher iWUE (Viger et al., 2016). After  
570 rewatering, the reopening of stomata lags behind recovery in LWP and the duration of this lag  
571 is dependent on the degree of water stress and the species considered (Liang and Zhang,  
572 1999). This is consistent with our observations of a delay between stress measured through  
573 LWP and the increase in iWUE.

574 Overall, multi-tool monitoring would allow to better understand the stress signal of the  
575 riparian forests in their natural conditions, and the delays between the responses that can be  
576 measured at the canopy level with remote sensing and the events leading to stress conditions.  
577 Additionally, using optical indexes or LiDAR data to retrieve parameters such as canopy  
578 greenness and structure would also allow us to better understand under which conditions the  
579 stress experienced by riparian trees leads to responses such as leaves yellowing, biomass loss,  
580 and reduced growth.

581

### 582 5.3. Good practices in designing campaigns to monitor water stress in riparian ecosystems.

583 Based on this information and our results, designing campaigns to assess or monitor water  
584 stress in riparian environment at large scales should couple TIR acquisitions with  
585 complementary data to cross-validate the observations such as *in-situ* measurements of water  
586 stress at target sites and optical remote-sensing indexes. Acquisitions would need to be  
587 planned depending on when the stress is expected to occur, the lag in response for the  
588 indicators used and their sensitivity to other environmental variables. Under these conditions,  
589 multi-date surveys are recommended to obtain multiple observations for a given summer. Our  
590 results also show that the acquisition of TIR data should target the peak of the stress period

591 and avoid dates with recent precipitations that could replenish water resources and affect air  
592 humidity.

593 Another way to acquire TIR data would then be to rely on drones or satellites as other  
594 airborne vectors such as helicopters or airplanes are less flexible in regards to the time of  
595 acquisition and are more costly which limits both the ability to acquire data multiple times per  
596 season and to target the most optimal window for TIR sensing of vegetation health. However,  
597 the use of satellite TIR data is limited by the poor spatial resolution of current TIR sensor  
598 which impacts their applicability to an early detection of water stress as they may be more  
599 sensitive to more delayed responses to stress conditions such as changes in canopy structure  
600 due to vegetation dieback and shifts in communities. Drone acquisitions suffer from lower  
601 spatial extent but the technology is emerging for the study of river temperature (Dugdale et  
602 al., 2022; Redana et al., 2024) after initial lowcost sensors proved prone to temperature  
603 drifting (Dugdale et al., 2019). They would however provide the high temporal resolution  
604 needed to better study how canopy temperature can be used as an indicator of water stress in  
605 complex and heterogeneous riparian environments by coupling drone acquisitions with *in-situ*  
606 surveys like those conducted in this study.

607 Multi-scale acquisition could help answer some of these issues by providing higher resolution  
608 data as ground truth which is increasingly common when processing satellite data  
609 (Carbonneau et al., 2020) or by using repeated drone surveys to help identify when more  
610 costly airborne acquisitions over a larger spatial extent should be conducted.

611

#### 612 5.4. Production of spatially-explicit knowledge for stakeholders and river management.

613 The results in this study builds upon previous research by validating the hypothesis that the  
614 riparian forest of the Ain River experiences water stress in the summer. Previous studies

615 focused on an upstream–downstream gradient (Lejot et al., 2011) or on stationary conditions  
616 (Dufour, 2005; Godfroy et al., 2023; Rollet et al., 2014) to highlight differences in the health  
617 status of the riparian forest but did not directly approach water stress issues. Here, we were  
618 able to confirm the presence of water stress during the summer using ecophysiological *in-situ*  
619 measurement. We then discovered that water stress conditions impacted *Populus nigra*  
620 individuals in areas where we expected an unstressful water availability, shedding light on the  
621 fact that even better-connected sectors of the riparian forest were experiencing water stress  
622 during summer 2022.

623 By applying an approach previously known to have an impact on the growth of *Fraxinus*  
624 *excelsior* (Dufour, 2005) and on the distribution of species in the riparian forest and its  
625 structure (Godfroy et al., 2023), focused on stationary conditions, we also detected a direct  
626 effect of stationary conditions on the stress status of *Populus nigra*. Drier stationary  
627 conditions due to channel incision led to higher levels of water stress during summer, and  
628 surveyed poplar trees experienced stress conditions earlier than those in reaches in better  
629 geomorphic conditions. These results support previous studies on other river systems that  
630 linked channel incision with an increase in mortality due to stress (Scott et al., 2000).

631 While *in-situ* assessments of the water status focused on *Populus nigra* because it was both  
632 the dominant species in the Ain River valley and of interest to local stakeholders, other  
633 riparian species might respond differently to droughts as previously described in section 5.1.  
634 This could limit results reproducibility and interpretability, yet in our study the relationship  
635 between forest stationary conditions and canopy temperature from airborne sensors held true  
636 at both the level of individual tree crowns and forest plots.

637 Overall, our results show that acquiring and using TIR data at various spatial scales can help  
638 monitor the health status of riparian forests and their response to anthropic pressures to  
639 generate actionable information for river managers.

640 In our case, we were able to detect a higher sensitivity of the riparian forest in Reach R4 to  
641 the 2003 drought which may partly be explained by lower water access compared to Reach  
642 R2 and by more pioneer populations compared to reaches R1 and R3. Since protecting  
643 pioneer environments is one of the priorities of local stakeholders, our results indicate that  
644 current gravel reinjection actions that focus on reaches R1 and R2 might not be sufficient to  
645 protect riparian environments closer to the confluence with the Rhône River from drought. In  
646 order to reduce the impact of climate change and warming weather on riparian forest, gravel  
647 reinjections actions closer to this sector could be considered to limit channel straightening and  
648 incision and improve groundwater access for pioneer trees. In a social context where the  
649 construction of a dam on the Rhône River upstream the Ain–Rhône confluence is being  
650 discussed, careful consideration needs to be given to the already fragilized riparian forest in  
651 this sector as this could exacerbate water stress issues and contribute to a shift towards dryer  
652 communities to the detriment of pioneer species such as *Populus nigra*.

653

#### 654 **Conclusion.**

655 Water stress conditions were detected using *in-situ* measurements and showed increased water  
656 stress on black poplars located in dryer terrestrial river margins. Airborne thermal infrared  
657 remote sensing was able to detect this difference in water status due to an increase in canopy  
658 temperature and enabled mapping of the riparian forest response to drought events on a 50 km  
659 reach of the Ain River. Differences in the spatial distribution of forest sensitivity over a 10-  
660 year period highlighted forest degradation in one of the reaches and provided feedback on

661 current restoration efforts. The use of satellite thermal infrared data was investigated but due  
662 to coarse spatial resolution and the limited spatial extent of riparian corridors, observations  
663 were more likely related to larger scale canopy openness and proximity to other land covers  
664 rather than water status, therefore highlighting the need for further studies coupling satellite  
665 data with ground truth acquired at higher resolution. Nonetheless, lasting degradations were  
666 detected near the confluence of the Ain River with the Rhône River which can be attributed to  
667 the combined effects of the 2003 drought event and groundwater levels lowering caused by  
668 channel incision. However, the variation of the stress signal during summer and the fact that  
669 not all surveys showed a temperature effect on tree canopy related to drought show the  
670 importance of multi-date and multi-source surveys when attempting to assess or monitor  
671 water stress in complex heterogeneous environments such as riparian forests. Despite these  
672 requirements, airborne TIR mapping appears as a promising tool to monitor the response of  
673 river systems to climate change thanks to its ability to inform about the water status of the  
674 riparian forest and to additionally provide crucial information about other components of river  
675 systems such as water surface temperature., but it needs to be repeated over the seasons and  
676 years to detect long term effects of such potential ramp disturbance (e.g. cumulative effects of  
677 heat pulses that are supposed to be more frequent and more intense).

678

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689

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695

696 **CRedit Statement.**

697 **J. Godfroy:** Conceptualization, Methodology, Formal Analysis, Investigation, Data Curation,  
698 Writing - Original Draft, Writing – Review & Editing., **P. Malherbe:** Formal Analysis, Data  
699 Curation, Writing – Review & Editing., **F. Gerle:** Formal Analysis, Data Curation, Writing –  
700 Review & Editing., **B. Marteau:** Investigation, Methodology, Data Curation, Writing –  
701 Review & Editing., **P. Lochin:** Investigation, Data Curation, Writing – Review & Editing., **S.**  
702 **Puijalon:** Writing – Review & Editing, Supervision., **J. Lejot:** Writing – Review & Editing,  
703 Supervision., **A. Vernay:** Conceptualization, Formal Analysis, Investigation, Data Curation,  
704 Writing – Review & Editing, Supervision, Funding acquisition., **H. Piégay:**  
705 Conceptualization, Methodology, Writing – Review & Editing, Supervision, Funding  
706 acquisition.

707 **Appendixes.**

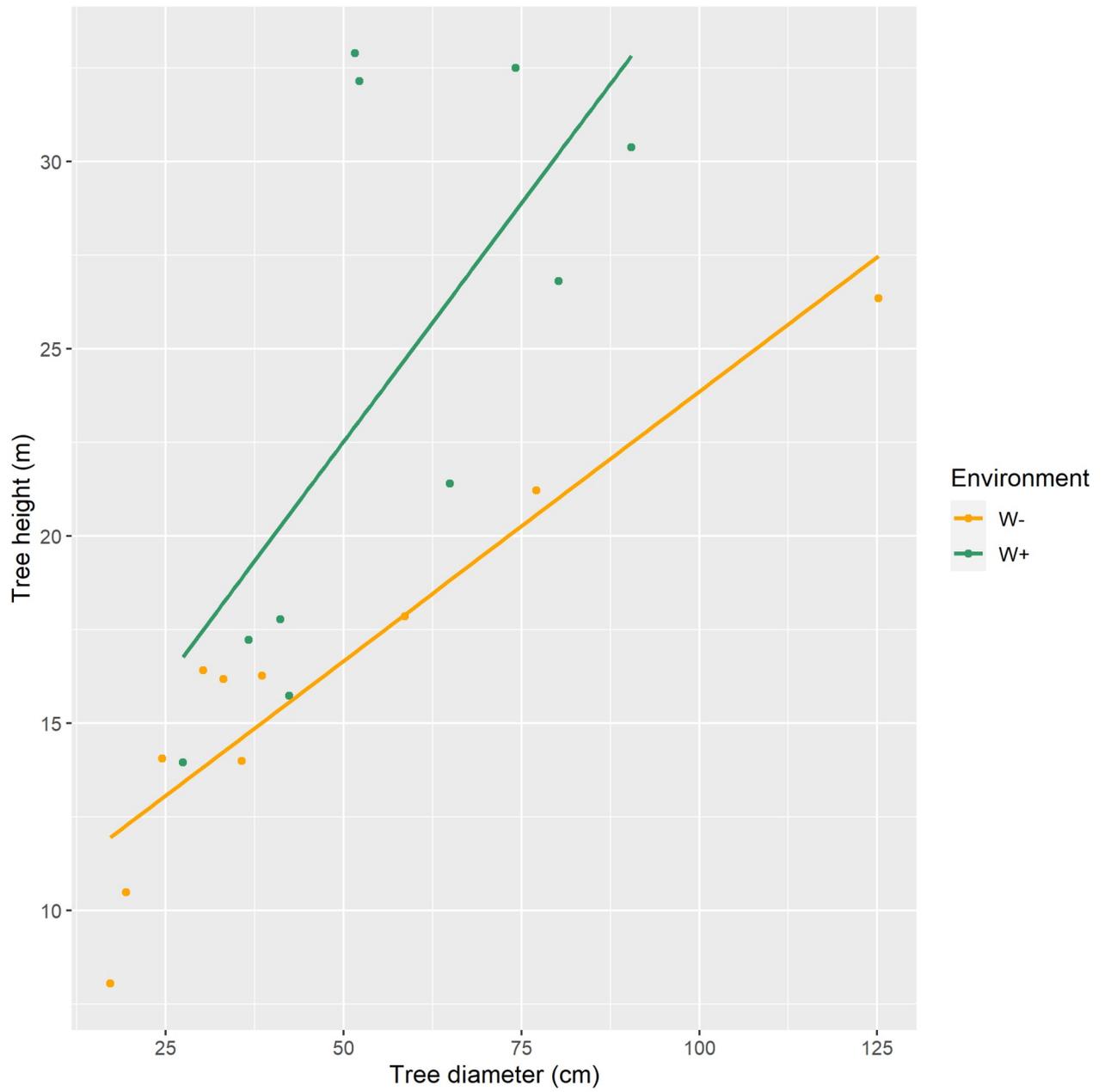
708 Table A. Dates for which leaf and phloem sampling was conducted during 2022.

	<b>Campaign Date</b>							
	May 5 <sup>th</sup>	May 25 <sup>th</sup>	June 10 <sup>th</sup>	June 28 <sup>th</sup>	July 19 <sup>th</sup>	Aug. 9 <sup>th</sup>	Aug. 23 <sup>rd</sup>	Sept. 27 <sup>th</sup>
<b>Leaf</b>	X	X	X	X	X	X	X	X
<b>Phloem</b>	X		X		X		X	X

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710

711 Figure B. Relationship between tree height and tree diameter on the W+ site ( $r^2 = 0.47$ ,  $p =$   
712  $0.0285$ ) and the W- site ( $r^2 = 0.85$ ,  $p = 0.0001$ ).

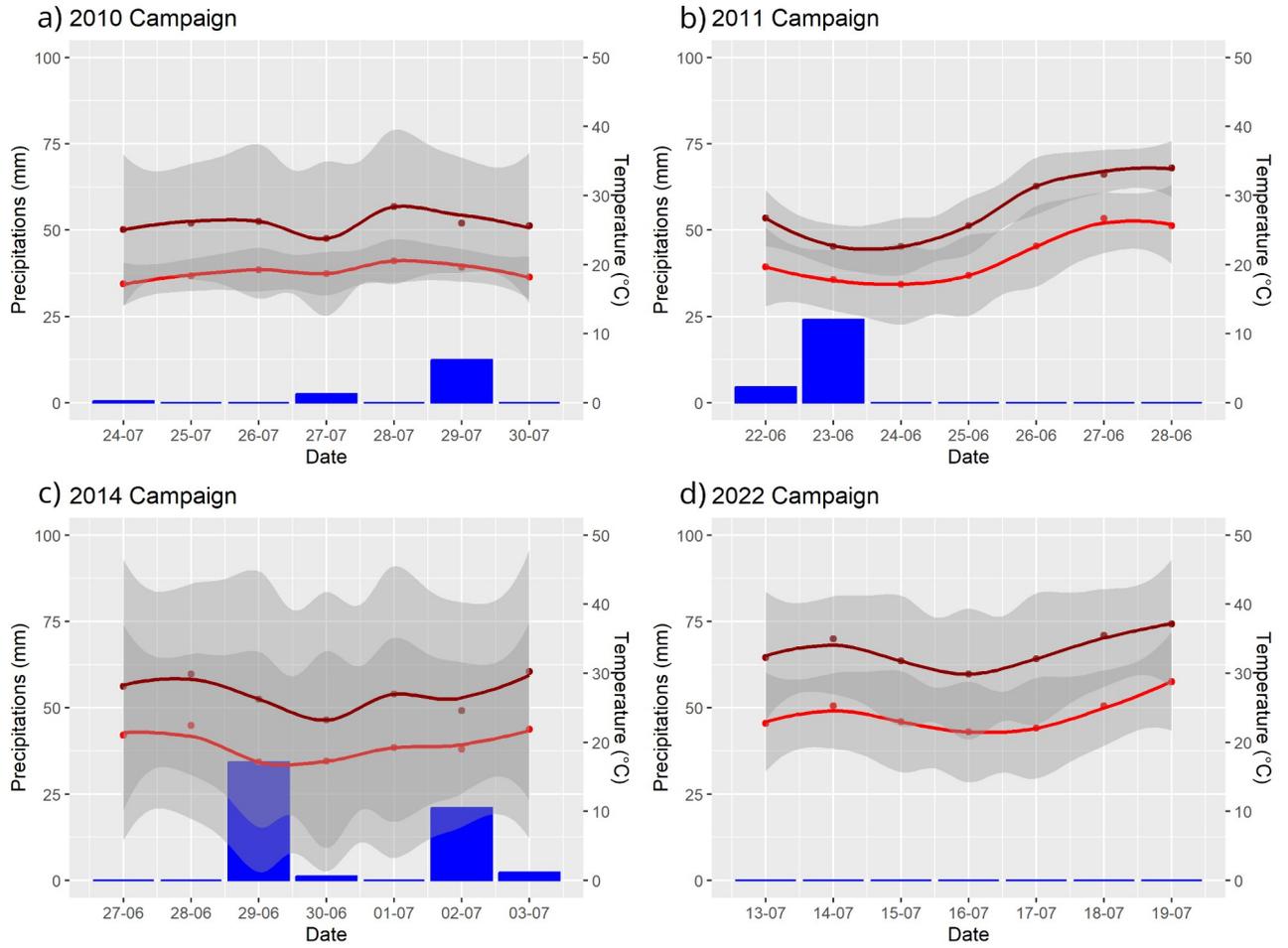


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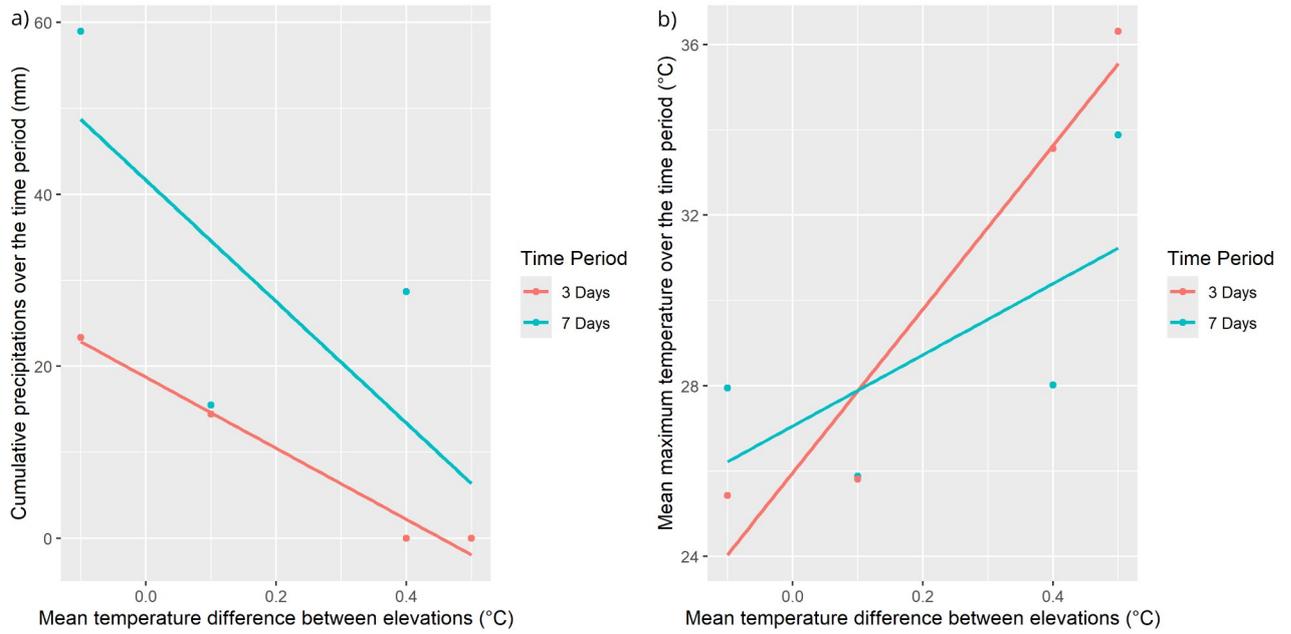
716 Figure C. Precipitation (blue), daily mean (red) and daily maximum (dark red) temperatures  
717 for a 7-day period preceding each airborne TIR campaign: a) 2010, b) 2011, c) 2014 and d)  
718 2022.



719

720

721 Figure D. Relationship between a) cumulative precipitations and b) mean maximum  
722 temperature for a 3-day or 7-day time period and the observed difference in mean  
723 temperatures from TIR data between the two elevation classes: connected (< 2.5 m from base  
724 flow) and disconnected (> 2.5 m from base flow).



725

726 **Bibliography.**

- Black, E., Blackburn, M., Harrison, G., Hoskins, B., Methven, J., 2004. Factors contributing to the summer 2003 European heatwave. *Weather* 59, 217–223. <https://doi.org/10.1256/wea.74.04>
- Bravard, J.-P., Amoros, C., Pautou, G., Bornette, G., Bournaud, M., Creuzé des Châtelliers, M., Gibert, J., Peiry, J.-L., Perrin, J.-F., Tachet, H., 1997. River incision in south-east France: morphological phenomena and ecological effects. *Regul. Rivers Res. Manag.* 13, 75–90. [https://doi.org/10.1002/\(SICI\)1099-1646\(199701\)13:1<75::AID-RRR444>3.0.CO;2-6](https://doi.org/10.1002/(SICI)1099-1646(199701)13:1<75::AID-RRR444>3.0.CO;2-6)
- Breton, V., Girel, J., Janssen, P., 2023. Long-term changes in the riparian vegetation of a large, highly anthropized river: Towards less hygrophilous and more competitive communities. *Ecol. Indic.* 155, 111015. <https://doi.org/10.1016/j.ecolind.2023.111015>
- Brodribb, T.J., Holbrook, N.M., 2003. Stomatal Closure during Leaf Dehydration, Correlation with Other Leaf Physiological Traits. *Plant Physiol.* 132, 2166–2173. <https://doi.org/10.1104/pp.103.023879>
- Carbonneau, P.E., Belletti, B., Micotti, M., Lastoria, B., Casaioli, M., Mariani, S., Marchetti, G., Bizzi, S., 2020. UAV-based training for fully fuzzy classification of Sentinel-2 fluvial scenes. *Earth Surf. Process. Landf.* 45, 3120–3140. <https://doi.org/10.1002/esp.4955>
- Ciezkowski, W., Kleniewska, M., Chormański, J., 2020. Thermal and Optical Indices for Wetland Habitats, are They Showing the Same Thing? *IEEE J. Sel. Top. Appl. Earth Obs. Remote Sens.* 13, 3951–3957. <https://doi.org/10.1109/JSTARS.2020.3008864>
- Comiti, F., Da Canal, M., Surian, N., Mao, L., Picco, L., Lenzi, M.A., 2011. Channel adjustments and vegetation cover dynamics in a large gravel bed river over the last 200years. *Geomorphology* 125, 147–159. <https://doi.org/10.1016/j.geomorph.2010.09.011>

- Décamps, H., Fortuné, M., Gazelle, F., Pautou, G., 1988. Historical influence of man on the riparian dynamics of a fluvial landscape. *Landsc. Ecol.* 1, 163–173. <https://doi.org/10.1007/BF00162742>
- Dépret, T., Thommeret, N., Piégay, H., Gautier, E., 2023. Can lateral mobility be restored along a highly domesticated low-energy gravel-bed river? *J. Environ. Manage.* 325, 116485. <https://doi.org/10.1016/j.jenvman.2022.116485>
- Dosskey, M.G., Vidon, P., Gurwick, N.P., Allan, C.J., Duval, T.P., Lowrance, R., 2010. The Role of Riparian Vegetation in Protecting and Improving Chemical Water Quality in Streams1. *JAWRA J. Am. Water Resour. Assoc.* 46, 261–277. <https://doi.org/10.1111/j.1752-1688.2010.00419.x>
- Dufour, S., 2005. Contrôles naturels anthropiques de la structure et de la dynamique des forêts riveraines (PhD Thesis). Université Jean Moulin Lyon III, Lyon.
- Dufour, S., Piégay, H., 2008. Geomorphological Controls of *Fraxinus Excelsior* Growth and Regeneration in Floodplain Forests. *Ecology* 89, 205–215. <https://doi.org/10.1890/06-1768.1>
- Dugdale, S.J., 2016. A practitioner’s guide to thermal infrared remote sensing of rivers and streams: recent advances, precautions and considerations. *WIREs Water* 3, 251–268. <https://doi.org/10.1002/wat2.1135>
- Dugdale, S.J., Franssen, J., Corey, E., Bergeron, N.E., Lapointe, M., Cunjak, R.A., 2016. Main stem movement of Atlantic salmon parr in response to high river temperature. *Ecol. Freshw. Fish* 25, 429–445. <https://doi.org/10.1111/eff.12224>
- Dugdale, S.J., Kelleher, C.A., Malcolm, I.A., Caldwell, S., Hannah, D.M., 2019. Assessing the potential of drone-based thermal infrared imagery for quantifying river temperature heterogeneity. *Hydrol. Process.* 33, 1152–1163. <https://doi.org/10.1002/hyp.13395>

- Dugdale, S.J., Klaus, J., Hannah, D.M., 2022. Looking to the Skies: Realising the Combined Potential of Drones and Thermal Infrared Imagery to Advance Hydrological Process Understanding in Headwaters. *Water Resour. Res.* 58, e2021WR031168. <https://doi.org/10.1029/2021WR031168>
- Dugdale, S.J., Malcolm, I.A., Kantola, K., Hannah, D.M., 2018. Stream temperature under contrasting riparian forest cover: Understanding thermal dynamics and heat exchange processes. *Sci. Total Environ.* 610–611, 1375–1389. <https://doi.org/10.1016/j.scitotenv.2017.08.198>
- Dumas, S., 2017. Inventaire des boisements forestiers de la Basse Vallée de l’Ain. Office National des Forêts.
- Dumas, S., Perrin, V., 2006. Le suivi de la forêt alluviale de la Basse Vallée de l’Ain : Inventaire de niveau II de 2006. Office National des Forêts.
- Fairfax, E., Small, E.E., 2018. Using remote sensing to assess the impact of beaver damming on riparian evapotranspiration in an arid landscape. *Ecohydrology* 11, e1993. <https://doi.org/10.1002/eco.1993>
- Gerle, F., Malherbe, P., Boisselet, C., Lafleuriel, D., Godfroy, J., Lochin, P., Marteau, B., Piégay, H., Puijalon, S., Vernay, A., 2023. Intrinsic water use efficiency estimate: an isotopic method.
- Godfroy, J., Lejot, J., Demarchi, L., Bizzi, S., Michel, K., Piégay, H., 2023. Combining Hyperspectral, LiDAR, and Forestry Data to Characterize Riparian Forests along Age and Hydrological Gradients. *Remote Sens.* 15, 17. <https://doi.org/10.3390/rs15010017>
- Gokool, S., Jarmain, C., Riddell, E., Swemmer, A., Lerm, R., Chetty, K.T., 2017. Quantifying riparian total evaporation along the Groot Letaba River: A comparison between infilled and

spatially downscaled satellite derived total evaporation estimates. *J. Arid Environ.* 147, 114–124. <https://doi.org/10.1016/j.jaridenv.2017.07.014>

González del Tánago, M., Martínez-Fernández, V., Aguiar, F.C., Bertoldi, W., Dufour, S., García de Jalón, D., Garófano-Gómez, V., Mandzukovski, D., Rodríguez-González, P.M., 2021. Improving river hydromorphological assessment through better integration of riparian vegetation: Scientific evidence and guidelines. *J. Environ. Manage.* 292, 112730. <https://doi.org/10.1016/j.jenvman.2021.112730>

Hsiao, T.C., Acevedo, E., 1974. Plant responses to water deficits, water-use efficiency, and drought resistance. *Agric. Meteorol., Plant modification for more efficient water use* 14, 59–84. [https://doi.org/10.1016/0002-1571\(74\)90011-9](https://doi.org/10.1016/0002-1571(74)90011-9)

Janssen, P., Stella, J.C., Piégay, H., Räßle, B., Pont, B., Faton, J.-M., Cornelissen, J.H.C., Evette, A., 2020. Divergence of riparian forest composition and functional traits from natural succession along a degraded river with multiple stressor legacies. *Sci. Total Environ.* 721, 137730. <https://doi.org/10.1016/j.scitotenv.2020.137730>

Jones, H.G., 1999. Use of infrared thermometry for estimation of stomatal conductance as a possible aid to irrigation scheduling. *Agric. For. Meteorol.* 95, 139–149. [https://doi.org/10.1016/S0168-1923\(99\)00030-1](https://doi.org/10.1016/S0168-1923(99)00030-1)

Jones, M.M., Rawson, H.M., 1979. Influence of Rate of Development of Leaf Water Deficits upon Photosynthesis, Leaf Conductance, Water Use Efficiency, and Osmotic Potential in Sorghum. *Physiol. Plant.* 45, 103–111. <https://doi.org/10.1111/j.1399-3054.1979.tb01672.x>

Kibler, C.L., Schmidt, E.C., Roberts, D.A., Stella, J.C., Kui, L., Lambert, A.M., Singer, M.B., 2021. A brown wave of riparian woodland mortality following groundwater declines during the 2012–2019 California drought. *Environ. Res. Lett.* 16, 084030. <https://doi.org/10.1088/1748-9326/ac1377>

- Klein, T., Rotenberg, E., Tatarinov, F., Yakir, D., 2016. Association between sap flow-derived and eddy covariance-derived measurements of forest canopy CO<sub>2</sub> uptake. *New Phytol.* 209, 436–446. <https://doi.org/10.1111/nph.13597>
- Lague, D., Feldmann, B., 2020. Topo-bathymetric airborne LiDAR for fluvial-geomorphology analysis, in: Paolo Tarolli, S.M.M. (Eds. ) (Ed.), *Remote Sensing of Geomorphology, Developments in Earth Surface Processes*. Elsevier, pp. 25–54. <https://doi.org/10.1016/B978-0-444-64177-9.00002-3>
- Lambs, L., Loubiat, M., Girel, J., Tissier, J., Peltier, J.-P., Marigo, G., 2006. Survival and acclimatation of *Populus nigra* to drier conditions after damming of an alpine river, southeast France. *Ann. For. Sci.* 63, 377–385. <https://doi.org/10.1051/forest:2006018>
- Lejot, J., Piégay, H., Hunter, P.D., Moulin, B., Gagnage, M., 2011. Utilisation de la télédétection pour la caractérisation des corridors fluviaux : exemples d’applications et enjeux actuels. *Géomorphologie Relief Process. Environ.* 17, 157–172. <https://doi.org/10.4000/geomorphologie.9362>
- Liang, J., Zhang, J., 1999. The relations of stomatal closure and reopening to xylem ABA concentration and leaf water potential during soil drying and rewatering. *Plant Growth Regul.* 29, 77–86. <https://doi.org/10.1023/A:1006207900619>
- Lurtz, M.R., Morrison, R.R., Gates, T.K., Senay, G.B., Bhaskar, A.S., Ketchum, D.G., 2020. Relationships between riparian evapotranspiration and groundwater depth along a semiarid irrigated river valley. *Hydrol. Process.* 34, 1714–1727. <https://doi.org/10.1002/hyp.13712>
- Luterbacher, J., Dietrich, D., Xoplaki, E., Grosjean, M., Wanner, H., 2004. European Seasonal and Annual Temperature Variability, Trends, and Extremes Since 1500. *Science* 303, 1499–1503. <https://doi.org/10.1126/science.1093877>

- Marteau, B., Michel, K., Piégay, H., 2022a. Can gravel augmentation restore thermal functions in gravel-bed rivers? A need to assess success within a trajectory-based before–after control–impact framework. *Hydrol. Process.* 36, e14480. <https://doi.org/10.1002/hyp.14480>
- Marteau, B., Piégay, H., Chandesris, A., Michel, K., Vaudor, L., 2022b. Riparian shading mitigates warming but cannot revert thermal alteration by impoundments in lowland rivers. *Earth Surf. Process. Landf.* n/a. <https://doi.org/10.1002/esp.5372>
- Mayes, M., Caylor, K.K., Singer, M.B., Stella, J.C., Roberts, D., Nagler, P., 2020. Climate sensitivity of water use by riparian woodlands at landscape scales. *Hydrol. Process.* 34, 4884–4903. <https://doi.org/10.1002/hyp.13942>
- Monclus, R., Dreyer, E., Villar, M., Delmotte, F.M., Delay, D., Petit, J.-M., Barbaroux, C., Le Thiec, D., Bréchet, C., Brignolas, F., 2006. Impact of drought on productivity and water use efficiency in 29 genotypes of *Populus deltoides*×*Populus nigra*. *New Phytol.* 169, 765–777. <https://doi.org/10.1111/j.1469-8137.2005.01630.x>
- Naiman, R., J., Decamps, H., Pastor, J., Johnston, C., 1988. The Potential Importance of Boundaries to Fluvial Ecosystems. *J. North Am. Benthol. Soc.* 7. <https://doi.org/10.2307/1467295>
- Naiman, R.J., Decamps, H., Pollock, M., 1993. The Role of Riparian Corridors in Maintaining Regional Biodiversity. *Ecol. Appl. Publ. Ecol. Soc. Am.* 3, 209–212. <https://doi.org/10.2307/1941822>
- Neale, C.M.U., Geli, H., Taghvaeian, S., Masih, A., Pack, R.T., Simms, R.D., Baker, M., Milliken, J.A., O’Meara, S., Witherall, A.J., 2011. Estimating evapotranspiration of riparian vegetation using high resolution multispectral, thermal infrared and lidar data, in: *Remote Sensing for Agriculture, Ecosystems, and Hydrology XIII*. SPIE, pp. 254–262. <https://doi.org/10.1117/12.903246>

O'Briain, R., 2019. Climate change and European rivers: An eco-hydromorphological perspective. *Ecohydrology* 12, e2099. <https://doi.org/10.1002/eco.2099>

Poff, N.L., Olden, J.D., Merritt, D.M., Pepin, D.M., 2007. Homogenization of regional river dynamics by dams and global biodiversity implications. *Proc. Natl. Acad. Sci.* 104, 5732–5737. <https://doi.org/10.1073/pnas.0609812104>

Redana, M., Lancaster, L.T., Chong, X.Y., Lip, Y.Y., Gibbins, C., 2024. An open-source method for producing reliable water temperature maps for ecological applications using non-radiometric sensors. *Remote Sens. Appl. Soc. Environ.* 34, 101184. <https://doi.org/10.1016/j.rsase.2024.101184>

Riis, T., Kelly-Quinn, M., Aguiar, F.C., Manolaki, P., Bruno, D., Bejarano, M.D., Clerici, N., Fernandes, M.R., Franco, J.C., Pettit, N., Portela, A.P., Tammeorg, O., Tammeorg, P., Rodríguez-González, P.M., Dufour, S., 2020. Global Overview of Ecosystem Services Provided by Riparian Vegetation. *BioScience* 70, 501–514. <https://doi.org/10.1093/biosci/biaa041>

Rivaes, R., Rodríguez-González, P.M., Albuquerque, A., Pinheiro, A.N., Egger, G., Ferreira, M.T., 2013. Riparian vegetation responses to altered flow regimes driven by climate change in Mediterranean rivers. *Ecohydrology* 6, 413–424. <https://doi.org/10.1002/eco.1287>

Rohde, M.M., Stella, J.C., Roberts, D.A., Singer, M.B., 2021. Groundwater dependence of riparian woodlands and the disrupting effect of anthropogenically altered streamflow. *Proc. Natl. Acad. Sci.* 118, e2026453118. <https://doi.org/10.1073/pnas.2026453118>

Rollet, A.J., 2007. Etude et gestion de la dynamique sédimentaire d'un tronçon fluvial à l'aval d'un barrage : le cas de la basse vallée de l'Ain (PhD Thesis). Université Jean Moulin Lyon 3.

Rollet, A.J., Piégay, H., Dufour, S., Bornette, G., Persat, H., 2014. Assessment of Consequences of Sediment Deficit on a Gravel River Bed Downstream of Dams in

Restoration Perspectives: Application of a Multicriteria, Hierarchical and Spatially Explicit Diagnosis. *River Res. Appl.* 30, 939–953. <https://doi.org/10.1002/rra.2689>

Roux, C., Alber, A., Bertrand, M., Vaudor, L., Piégay, H., 2015. “FluvialCorridor”: A new ArcGIS toolbox package for multiscale riverscape exploration. *Geomorphology, Geomorphology in the Geocomputing Landscape: GIS, DEMs, Spatial Analysis and statistics* 242, 29–37. <https://doi.org/10.1016/j.geomorph.2014.04.018>

Sankey, T., Hultine, K., Blasini, D., Koepke, D., Bransky, N., Grady, K., Cooper, H., Gehring, C., Allan, G., 2021. UAV thermal image detects genetic trait differences among populations and genotypes of Fremont cottonwood (*Populus fremontii*, Salicaceae). *Remote Sens. Ecol. Conserv.* 7, 245–258. <https://doi.org/10.1002/rse2.185>

Scholander, P.F., Bradstreet, E.D., Hemmingsen, E.A., Hammel, H.T., 1965. Sap Pressure in Vascular Plants. *Science* 148, 339–346. <https://doi.org/10.1126/science.148.3668.339>

Scott, M.L., Lines, G.C., Auble, G.T., 2000. Channel incision and patterns of cottonwood stress and mortality along the Mojave River, California. *J. Arid Environ.* 44, 399–414. <https://doi.org/10.1006/jare.1999.0614>

Seibt, U., Rajabi, A., Griffiths, H., Berry, J.A., 2008. Carbon isotopes and water use efficiency: sense and sensitivity. *Oecologia* 155, 441–454. <https://doi.org/10.1007/s00442-007-0932-7>

Simon, A., Collison, A.J.C., 2002. Quantifying the mechanical and hydrologic effects of riparian vegetation on streambank stability. *Earth Surf. Process. Landf.* 27, 527–546. <https://doi.org/10.1002/esp.325>

Singer, M.B., Stella, J.C., Dufour, S., Piégay, H., Wilson, R.J.S., Johnstone, L., 2013. Contrasting water-uptake and growth responses to drought in co-occurring riparian tree species. *Ecohydrology* 6, 402–412. <https://doi.org/10.1002/eco.1283>

- Smith, S.D., Wellington, A.B., Nachlinger, J.L., Fox, C.A., 1991. Functional Responses of Riparian Vegetation to Streamflow Diversion in the Eastern Sierra Nevada. *Ecol. Appl.* 1, 89–97. <https://doi.org/10.2307/1941850>
- Stella, J.C., Riddle, J., Piégay, H., Gagnage, M., Trémélo, M.-L., 2013a. Climate and local geomorphic interactions drive patterns of riparian forest decline along a Mediterranean Basin river. *Geomorphology, Process geomorphology and ecosystems: Disturbance regimes and interactions* 202, 101–114. <https://doi.org/10.1016/j.geomorph.2013.01.013>
- Stella, J.C., Rodríguez-González, P.M., Dufour, S., Bendix, J., 2013b. Riparian vegetation research in Mediterranean-climate regions: common patterns, ecological processes, and considerations for management. *Hydrobiologia* 719, 291–315. <https://doi.org/10.1007/s10750-012-1304-9>
- Tabacchi, E., Lambs, L., Guilloy, H., Planty-Tabacchi, A.-M., Muller, E., Décamps, H., 2000. Impacts of riparian vegetation on hydrological processes. *Hydrol. Process.* 14, 2959–2976. [https://doi.org/10.1002/1099-1085\(200011/12\)14:16/17<2959::AID-HYP129>3.0.CO;2-B](https://doi.org/10.1002/1099-1085(200011/12)14:16/17<2959::AID-HYP129>3.0.CO;2-B)
- Vernay, A., Tian, X., Chi, J., Linder, S., Mäkelä, A., Oren, R., Peichl, M., Stangl, Z.R., Torngern, P., Marshall, J.D., 2020. Estimating canopy gross primary production by combining phloem stable isotopes with canopy and mesophyll conductances. *Plant Cell Environ.* <https://doi.org/10.1111/pce.13835>
- Viger, M., Smith, H.K., Cohen, D., Dewoody, J., Trewin, H., Steenackers, M., Bastien, C., Taylor, G., 2016. Adaptive mechanisms and genomic plasticity for drought tolerance identified in European black poplar (*Populus nigra* L.). *Tree Physiol.* 36, 909–928. <https://doi.org/10.1093/treephys/tpw017>
- Volaire, F., 2018. A unified framework of plant adaptive strategies to drought: Crossing scales and disciplines. *Glob. Change Biol.* 24, 2929–2938. <https://doi.org/10.1111/gcb.14062>

- Wawrzyniak, V., Allemand, P., Bailly, S., Lejot, J., Piégay, H., 2017. Coupling LiDAR and thermal imagery to model the effects of riparian vegetation shade and groundwater inputs on summer river temperature. *Sci. Total Environ.* 592, 616–626. <https://doi.org/10.1016/j.scitotenv.2017.03.019>
- Wawrzyniak, V., Piégay, H., Allemand, P., Vaudor, L., Goma, R., Grandjean, P., 2016. Effects of geomorphology and groundwater level on the spatio-temporal variability of riverine cold water patches assessed using thermal infrared (TIR) remote sensing. *Remote Sens. Environ.* 175, 337–348. <https://doi.org/10.1016/j.rse.2015.12.050>
- Whitledge, G.W., Rabeni, C.F., Annis, G., Sowa, S.P., 2006. Riparian Shading and Groundwater Enhance Growth Potential for Smallmouth Bass in Ozark Streams. *Ecol. Appl.* 16, 1461–1473. [https://doi.org/10.1890/1051-0761\(2006\)016\[1461:RSAGEG\]2.0.CO;2](https://doi.org/10.1890/1051-0761(2006)016[1461:RSAGEG]2.0.CO;2)
- Wilbur, N.M., O’Sullivan, A.M., MacQuarrie, K.T.B., Linnansaari, T., Curry, R.A., 2020. Characterizing physical habitat preferences and thermal refuge occupancy of brook trout (*Salvelinus fontinalis*) and Atlantic salmon (*Salmo salar*) at high river temperatures. *River Res. Appl.* 36, 769–783. <https://doi.org/10.1002/rra.3570>