

1 Abiotic constraints and recreational rock 2 climbing shape cliff vegetation in Freyr, Belgium

3 Authors

4 Sarane Coen^{1,2,3} (<https://orcid.org/0009-0009-1317-1582>)

5 Georgia R. Harrison⁴ (<https://orcid.org/0000-0003-0544-4471>)

6 Amre van den Maagdenberg³

7 Christophe Metsu^{5,6} (<https://orcid.org/0000-0002-5153-9351>)

8 Laura M. Boggess⁷ (<https://orcid.org/0000-0003-4959-8464>)

9 Francisco Velasquez-Espin⁸ (<https://orcid.org/0009-0000-7606-2505>)

10 Martí March-Salas^{9,10} (<https://orcid.org/0000-0001-5347-4056>)

11 Kobe Burdack¹¹

12 Elias P. Goossens¹² (<https://orcid.org/0000-0002-4994-2029>)

13 Koenraad Van Meerbeek^{*5,6} (<https://orcid.org/0000-0002-9260-3815>)

14 Jonas J. Lembrechts^{*2,3} (<https://orcid.org/0000-0002-1933-0750>)

15 *Shared last

16 The author's institutional affiliations

17 ¹Institute of Botany, University of Liège, Belgium

18 ²Ecology & Biodiversity (E&B), Utrecht University, The Netherlands

19 ³Research Group of Plants and Ecosystems (PLECO), Department of Biology, University of Antwerp,
20 Belgium

21 ⁴Department of Animal and Range Sciences, Montana State University, Bozeman, MT USA.

22 ⁵Department of Earth and Environmental Sciences, KU Leuven, Leuven, Belgium

23 ⁶KU Leuven Plant Institute, KU Leuven, Leuven, Belgium

24 ⁷Ecology and Conservation Biology, Mars Hill University, North Carolina, USA

25 ⁸Department of Biology and Biochemistry, University of Houston, 4302 University Drive, Houston,
26 Texas 77004, USA

27 ⁹Area of Biodiversity and Conservation, Department of Biology and Geology, Physics and Inorganic
28 Chemistry, Universidad Rey Juan Carlos-ESCET, Móstoles, Madrid, Spain

29 ¹⁰Instituto de Investigación en Cambio Global (IICG-URJC), Universidad Rey Juan Carlos, Móstoles,
30 Madrid, Spain

31 ¹¹Faculty of Engineering Technology, KU Leuven, Leuven, Belgium

32 ¹²Department of Biology, Research group WILD, VUB, Brussels, Belgium.

33 **Correspondence**

34 Sarane Coen

35 Institute of Botany, University of Liège, Belgium

36 Ecology & Biodiversity (E&B), Utrecht University, The Netherlands

37 Research Group of Plants and Ecosystems (PLECO), Department of Biology, University of Antwerp,
38 Belgium

39 Email: coen.sarane@gmail.com

40 **Funding information**

41 J.J.L. has received funding from the European Union's Horizon Europe research and innovation
42 programme under the Marie Skłodowska-Curie grant agreement No. 101207721.

43 S.C. received support from the French Community of Belgium in the form of funding for a FRIA
44 scholarship No. 40038334.

45 **Acknowledgements**

46 We thank Michiel Nollet, Lucas De Ridder-Rombaut, Sebastiaan Verbeke, Nick Vandermeeren, and
47 Robbe Roskams for their assistance with fieldwork on the cliff faces. We are grateful to David Léduc,
48 site manager of Freyr, as well as to Klim- en Bergsportfederatie, Club Alpin Belge, and the Belgian
49 Rebolting Team for their collaboration and practical support. We also thank the climbers who
50 completed the survey that informed our estimates of climbing intensity.

51 **Author contributions**

52 S.C., K.V.M., and J.J.L. conceived the research idea. S.C. handled data curation, project administration,
53 visualisation, and wrote the original draft. Formal analyses were conducted by S.C., G.R.H., K.V.M., and
54 J.J.L.; S.C., L.M.B., K.B., M.M.S., F.V.E., K.V.M., and J.J.L. developed the methodological protocols. Data
55 were collected by S.C., K.B., E.P.G., A.M., C.M., K.V.M., and J.J.L. The project was supervised by J.J.L.
56 and K.V.M. All authors contributed to validation, review, and editing of the manuscript.

57

58 ABSTRACT

59 **Aim:** Cliff ecosystems support diverse vascular plant communities due to high abiotic heterogeneity
60 and their historical role as climatic refugia. However, cliffs are increasingly exposed to disturbances
61 from recreational rock climbing. The ecological effects of climbing likely depend on abiotic cliff
62 characteristics—such as slope, aspect, and microtopography—but these context-dependent
63 relationships have not been consistently evaluated. We investigated how climbing intensity and abiotic
64 characteristics jointly influence plant diversity, community composition, and community-level
65 functional tendencies in a local case study on limestone cliff faces.

66 **Location:** Freyr limestone cliffs, Belgium

67 **Methods:** Vegetation was surveyed in the Freyr climbing site in Belgium using a standardised protocol
68 across 248 1-m² subplots distributed along 13 vertical transects spanning unclimbed, low-intensity,
69 and high-intensity climbing routes. We quantified species richness, vegetation cover, Shannon
70 diversity, community composition, and community-weighted means (CWMs) of Grime's CSR strategies
71 (competitive, stress-tolerant, and ruderal) to characterise community-level functional tendencies. We
72 examined the effects of climbing intensity, and of abiotic factors such as slope, aspect, and
73 microtopography using generalised linear mixed models, multivariate ordination and an indicator
74 species analysis.

75 **Results:** A total of 103 vascular plant species were recorded. Abiotic factors influenced vegetation
76 patterns more than climbing intensity. Species richness and vegetation cover were highest at moderate
77 levels of microtopography, lower at steep slopes, and cover was reduced on south-facing cliffs. The
78 effect of climbing intensity depended on microtopography: species richness peaked at low climbing
79 intensity on cliffs with pronounced microtopography, consistent with the intermediate disturbance
80 hypothesis. Functional and species composition shifted with climbing: competitive and stress-tolerant
81 species were more abundant on unclimbed cliffs whereas ruderal species were more abundant on
82 climbed cliffs. Unclimbed cliffs were also associated with threatened (red-listed) indicator species.

83 **Conclusions:** At Freyr, cliff vegetation was mainly structured by abiotic conditions, with climbing acting
84 as a context-dependent disturbance that influenced community composition and functional
85 tendencies rather than overall cover or richness. Cliffs with high microtopography under moderate
86 climbing disturbance supported the highest diversity, while unclimbed cliffs harboured more stress-
87 tolerant and red-listed species, indicating a potential loss of vulnerable vegetation under increasing
88 climbing pressure. These site-specific findings underscore the need for multi-site studies and detailed

89 analyses of abiotic factors to better understand and manage the ecological impacts of recreational
90 climbing.

91 KEYWORDS

92 Cliff ecology, rock climbing, disturbance, environmental filtering, Grime CSR, functional traits,
93 Intermediate disturbance hypothesis, microclimate

94 INTRODUCTION

95 Rock cliffs host high vascular plant diversity, including rare and endemic species adapted to harsh,
96 vertical environments ([Datar and Watve 2018](#); [Ellenberg 2009](#); [García et al. 2020](#); [Kuntz and Larson](#)
97 [2006](#); [Larson et al. 2005](#); [March-Salas et al. 2023](#); [Rusterholz et al. 2011](#); [Harrison et al. 2024](#);
98 [Fitzsimons and Michael 2017](#)). The steep and fragmented nature of cliffs creates isolated habitats
99 where pressure from competitive species is reduced ([Larson et al. 2005](#)). As a result, cliffs serve as one
100 of the last holdouts for stress-tolerant taxa ([Ellenberg 2009](#)). However, spatial isolation also limits
101 dispersal and connectivity. Under accelerating climate change and increasing anthropogenic pressure
102 on biodiversity ([Hooper et al. 2012](#); [Calvin et al. 2023](#); [IPBES 2019](#); [De Pryck 2021](#)), this restricted
103 migration may increase local extinction risk ([March-Salas et al. 2025](#)). Cliff ecosystems are therefore
104 both rich in biodiversity and vulnerable to global change. Yet, they remain poorly studied and deserve
105 greater ecological attention ([Capó et al. 2023](#); [Cartwright 2019](#); [March-Salas et al. 2023](#)).

106 Cliff biodiversity arises from limited historical disturbance, and fine-scale environmental
107 heterogeneity. Variations in slope, aspect, and microtopography create gradients in solar radiation,
108 moisture, and nutrient availability, resulting in a patchwork of microclimates that can buffer or amplify
109 environmental extremes ([Larson et al. 2005](#); [Bartlett et al. 1990](#); [Bramer et al. 2018](#)). Cliff surfaces
110 exhibit more complex radiation dynamics than horizontal or gently sloped surfaces ([Geiger et al. 2012](#);
111 [Larson et al. 2005](#)). South-facing slopes in the Northern Hemisphere generally receive more solar
112 radiation and are warmer and drier, while north-facing slopes are cooler and more humid ([Suggitt et](#)
113 [al. 2011](#)). These microclimatic gradients are expected to strongly influence plant establishment,
114 development and persistence; however, in situ measurements on vertical cliff faces remain limited,
115 leaving key mechanisms misunderstood and calling for many more case studies. As a result, cliffs can
116 function simultaneously as refugia for cold-adapted species, facilitating persistence during climatic
117 change, and as stepping stones for warm-adapted taxa extending their ranges ([De Frenne et al. 2013](#);
118 [Dobrowski 2011](#); [Speziale and Ezcurra 2015](#); [Keppel et al. 2015](#)).

119 Slope and microtopography strongly shape plant communities too ([Boggess et al. 2021](#); [Holzschuh](#)
120 [2016](#)). Steep cliffs are often well-drained and nutrient-poor, while cliffs with shallower slopes tend to
121 trap more debris and moisture, creating sites suitable for plant establishment and growth ([Bartlett et](#)
122 [al. 1990](#); [Clark and Hessel 2015](#); [Harrison et al. 2022](#); [Kuntz and Larson 2006](#); [Larson et al. 2005](#)).
123 Microtopography—such as crevices, pockets, fractures, and surface irregularities—provides microsites
124 for soil and water retention, creating places for propagule catchment and plant growth ([Nuzzo 1996](#);
125 [Larson et al. 2005](#)). Increased microtopography often enhances species richness. Thus, the particular
126 abiotic conditions of cliffs act as a strong mediator of the presence and structure of cliff vegetation
127 ([Kuntz and Larson 2006](#); [Harrison et al. 2022](#)).

128 Due to their steepness and inaccessibility, cliffs have historically experienced lower levels of grazing,
129 trampling, and development compared to surrounding landscapes.. However, the rising popularity of
130 outdoor recreation, such as rock climbing ([Aubel and Lefèvre 2022](#)), has resulted in increased
131 disturbance regimes in these fragile ecosystems. For instance, the Flemish climbing federation (Klim-
132 en Bergsportfederatie, KBF) recorded a 28% increase in its members between 2017 and 2024 (KBF,
133 personal communication, 2024). Rock climbing can cause both mechanical and chemical disturbances,
134 through vegetation and loose rock removal upon establishment, trampling and the use of climbing
135 chalk ([Morales-Armijo et al. 2024](#)) ; Léduc, personal communication, 2024).

136 Research on the impact of rock climbing on vascular plants has revealed mixed results ([Boggess et al.](#)
137 [2021](#); [Holzschuh 2016](#)). Most studies have documented negative effects of climbing, including reduced
138 species richness and abundance ([Camp and Knight 1998](#); [McMillan and Larson 2002](#); [Müller et al. 2004](#);
139 [Lorite et al. 2017](#); [Schmera et al. 2018](#)), as well as shifts in community composition ([Morales-Armijo et](#)
140 [al. 2024](#)). These compositional changes can involve a decline in specialist rock-dwelling species ([Müller](#)
141 [et al. 2004](#); [Lorite et al. 2017](#); [Camp and Knight 1998](#)). In contrast, March-Salas et al. ([2018](#)) reported
142 a shift primarily affecting generalist species, with reduced richness and cover of non-specialists in
143 climbed areas. Nevertheless, several studies detected weak or no direct effect of climbing once
144 variation in abiotic cliff characteristics was accounted for ([Clark and Hessel 2015](#); [Boggess et al. 2017](#);
145 [Harrison et al. 2022](#); [Nuzzo 1996](#); [Kuntz and Larson 2006](#)).

146 Together, these findings indicate that the ecological effects of rock climbing on plants can be context
147 dependent, varying with baseline vegetation composition, abiotic cliff characteristics, and climbing
148 intensity ([Holzschuh 2016](#); [Boggess et al. 2021](#)). In addition, climbers tend to preferentially select cliff
149 faces that are naturally less vegetated, further complicating causal inference ([Boggess et al. 2017](#);
150 [Farris 1998](#); [Kuntz and Larson 2006](#); [Müller et al. 2004](#)). Consequently, expanding research to
151 additional sites and new locations is essential to disentangle disturbance effects from abiotic filtering.

152 Previous studies have been clustered in North America, with European research mostly limited to
153 Switzerland and Spain ([Boggess et al. 2021](#)).

154 To enable more robust and long-term reproducibility and global comparability ([Boggess et al. 2021](#)),
155 the international MIREN Rocks network, a global research community of rock climbing ecologists,
156 developed a standardised sampling protocol (Appendix A.1). This protocol provides a consistent
157 framework for quantifying vegetation, abiotic conditions, and microclimatic variation on vertical rock
158 faces. Here, we present the first application of this protocol in Belgium, at the limestone cliffs of Freyr.
159 Beyond extending the geographical coverage of existing research, this study applies a multi-faceted
160 approach to estimate climbing intensity (Appendix A.2), and integrates in situ microclimatic
161 measurements on vertical cliff faces.

162 Beyond species richness and cover, we focus on community composition and functional traits to
163 capture subtle responses. Functional strategies are interpreted through Grime's ([1977](#)) CSR
164 framework, which classifies species as competitive (C), stress-tolerant (S), or ruderal (R), with most
165 species combining strategies to cope with local stress (e.g., limited resources) and disturbance (e.g.,
166 biomass removal). Harsh cliff conditions favour stress-tolerant strategies, yet disturbance may shift
167 communities toward ruderal species, potentially altering functional composition ([Rusterholz et al.](#)
168 [2011](#)).

169 Specifically, we examine how species richness, vegetation cover, functional traits (CSR strategies), and
170 community composition vary along gradients of climbing intensity, slope, aspect, and microtopography
171 in Freyr, to clarify how disturbance interacts with abiotic filtering to shape cliff-face vegetation.

172 METHODS

173 Study site

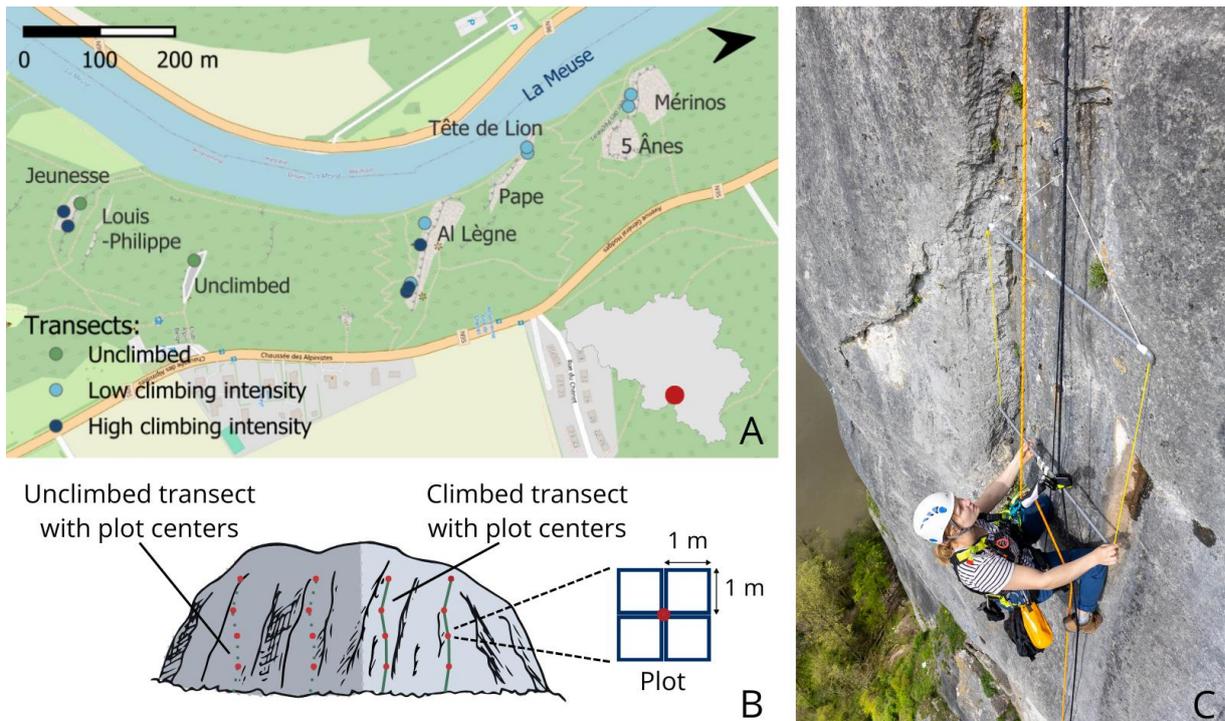
174 The study was conducted at the Freyr climbing site in the Meuse Valley, Belgium (50.223697, 4.895386)
175 (fig. 1A). This Natura 2000 protected area is characterised by limestone cliffs and species-rich
176 calcareous grasslands of high conservation value ([Newton et al. 2012](#)). Freyr has been Belgium's largest
177 climbing crag in terms of both area and number of routes since the 1930s and supports year-round
178 climbing. ([Bott 2022](#)). The Club Alpin Belge (CAB) manages site access and maintenance.

179 Data acquisition

180 Vegetation data were collected following the *MIREN Rocks* survey protocol (Appendix A.1). Fieldwork
181 was conducted in June and July 2024, during peak flowering. Vegetation surveys followed a hierarchical
182 structure (site > zone > transect > plot > subplot). The site corresponded to the Freyr climbing area (fig.

183 1A), within which eight cliff zones (i.e. standalone cliffs) were surveyed. Thirteen vertical transects
184 were established across these zones (fig. 1A), encompassing a total of 18 climbing routes. A single
185 transect could include sections of multiple routes, as routes in Freyr can deviate from the vertical axis
186 or consist of multiple routes above each other. Survey transect length varied between 18 and 76 m,
187 depending on cliff height. Along each transect, between three and nine plots were placed, with longer
188 transects having more plots, resulting in a total of 62 plots. Each plot covered 4 m² and consisted of
189 four 1 m² subplots (fig. 1B-C), yielding 248 subplots in total. Subplots were the primary observational
190 units for vegetation data collection and analysis, with plots, transects, and zones treated as nested
191 random effects.

192 At the plot level, we measured abiotic cliff characteristics, including height above the base of the cliff
193 and aspect, which was determined using a compass. At the subplot level, slope was measured with an
194 inclinometer, and microtopography was quantified using a surface heterogeneity index ranging from
195 one (smooth rock surface) to ten (high surface heterogeneity characterised by abundant fissures,
196 cracks, crevices, and/or pockets) (Appendix A.1 fig.7; [Boggess et al. 2021](#); [Boggess et al. 2017](#)). All
197 vascular plant species present in each subplot were identified, and both total vegetation cover and
198 species-specific cover were visually estimated as percentages of subplot area. Overall, 71.8% of taxa
199 were identified to species level, 16.5% to genus level only, and 11.6% remained unidentified. Taxa
200 identified only to genus were treated as distinct operational taxonomic units in all diversity and
201 community analyses.



203

204 **Figure 1:** A) Map (OpenStreetMap) of the Freyr study area, showing the different cliff zones (grey). Each dot represents a
 205 transect; colours indicate the dominant climbing intensity level of the transect. Inset: location of Freyr in Belgium B)
 206 Sampling design for cliff faces with climbed and unclimbed zones. The vegetation was surveyed along vertical transects,
 207 with each plot composed of four 1 m² subplots. C) Picture of the vegetation survey on the cliff of Al Lègne, performed while
 208 rappelling down, using a 1 m² frame attached to a rope as subplot.

209 Climbing intensity

210 The MIREN Rocks protocol recommends monitoring vegetation in paired climbed and unclimbed
 211 transects. This was, however, impractical for our study because unclimbed zones were scarce or unsafe
 212 to access. Instead, we initially assessed climbing intensity of the climbed transects with the Climbing
 213 Use Index (CUI) based on guidebook star ratings and walking approach time (Clark and Hessler 2015),
 214 and PCA-based metrics derived from online ascent logs and guidebook information. However, all of
 215 these approaches proved unsuitable for Freyr. The CUI produced only a narrow range of values and
 216 relied on subjective route ratings that reflect appreciation rather than actual use in our study area.
 217 PCA-based scores did not match local climbers' experience (see Appendix A.2. for details). Given these
 218 limitations, climbing intensity was estimated using a survey of eight experienced local climbers, who
 219 ranked the sampled routes according to perceived use.

220 Agreement among individual rankings was tested using Kendall's coefficient of concordance (W), which
 221 indicated moderate alignment ($W=0.473$, $p<0.001$). To avoid overconfidence with the ranking, climbing
 222 intensity was classified at the plot level into three categories based on average ranks: unclimbed ($n=28$
 223 subplots), low ($n=116$), and high ($n=104$); plot-level classification allowed climbing intensity to vary

224 within transects composed of different climbing routes. The proportion of climbers whose ranking
225 placed each route in the same category was calculated to evaluate how well the classification reflected
226 climbers' perceptions. Most routes showed high agreement (11 of 18 routes >85%), while only two
227 routes showed <50% consensus, indicating that surveys of experienced local climbers provided a
228 reliable estimate of climbing intensity at Freyr.

229 Data analysis

230 Alpha diversity metrics

231 Diversity metrics were quantified at subplot level as follows: species richness was calculated as the
232 number of vascular plant species, total vegetation cover was estimated as proportion of each subplot
233 covered by vascular plants, and Shannon diversity index, calculated using the vegan package ([Oksanen
234 et al. 2024](#)), integrating species richness and relative abundances to account for evenness, from
235 species-specific cover values to account for both richness and evenness. Subplots without vegetation
236 were excluded from this calculation, resulting in n = 150 subplots. Simpson index was calculated as
237 well and displayed in Appendix A.5.

238 Species richness, Shannon diversity and total vegetation cover were analysed using generalised linear
239 mixed models (GLMMs; glmmTMB package ([Brooks et al. 2017](#))). Species richness was modelled with
240 a negative binomial GLMM to account for overdispersion, total vegetation cover with a beta regression
241 GLMM and Shannon diversity with a GLMM using a Tweedie distribution with zero-inflation to
242 accommodate positive continuous values with many zeros.

243 Fixed effects included climbing intensity (factor with three levels: unclimbed, low, high), aspect (factor
244 with two levels: north- vs. south-facing), slope (continuous, standardised to mean = 0 and SD = 1), and
245 microtopography (discrete, standardised). Based on visual inspection, a quadratic effect of
246 microtopography was included for species richness and total vegetation cover. Interactions between
247 climbing and microtopography were included for all response variables to account for the potential
248 differences in how microtopography modifies the effect of climbing on vegetation structure and
249 diversity.

250 To account for the nested sampling design (subplot < plot < transect < zone), all models included a
251 hierarchical random-effects structure. If convergence or singularity issues arose, the random-effects
252 structure was simplified. Specifically, species richness models included plots nested in transects nested
253 within zones; total vegetation cover models included plots nested within transects; and Shannon
254 diversity included plots only.

255 Functional diversity (CSR strategies)
256 Plant strategies were quantified using Grime's CSR framework (C = competitive, S = stress-tolerant, R
257 = ruderal) based on the KS index, which assigns each species proportional C, S, and R values (summing
258 to one) derived from functional traits related to growth, persistence, and disturbance tolerance
259 [\(Ivanova and Zolotova 2023\)](#). When species-level values were unavailable, genus-level means derived
260 from the KS dataset were used, but only when within-genus variation in CSR values (standard
261 deviation) was below 1. For each subplot, relative abundance in the community-weighted mean of
262 each strategy was calculated by multiplying each species' CSR values by its proportional cover,
263 summing these values across species, and dividing by the total cover of all species with available CSR
264 data. These CWM values reflect the dominant functional strategies within each plant community.
265 Subplots with less than 70% representation of species with known CSR values were excluded, resulting
266 in 136 subplots, nested within 51 plots, 13 transects, and 8 zones.

267 Separate Tweedie GLMMs were fitted for each strategy (C, S, R) to test for effects of climbing intensity,
268 slope, aspect, and microtopography. Quadratic terms were excluded as they were not expected, and
269 interaction terms were not included to avoid overfitting. Random-effects structures were adapted to
270 each strategy to ensure model convergence: stress-tolerant CWMs included plots nested in transects
271 and zones, competitive CWMs excluded zones, and ruderal CWMs included plots only.

272 Differences in occurrence between strategies were evaluated using a linear model with CWM as the
273 response variable and strategy (C, S, R) as a fixed effect, and (subplot < plot < transect < zone) as
274 random effects with the lme4 package [\(Bates et al. 2015\)](#). Pairwise comparisons between the
275 strategies were performed using the emmeans package [\(Lenth and Piaskowski 2017\)](#) and Tukey-
276 adjusted p-values.

277 Model diagnostics for both alpha diversity and CSR models, including residual distribution,
278 overdispersion, and multicollinearity, were assessed using DHARMA [\(Hartig 2024\)](#) and performance
279 [\(Lüdecke et al. 2021\)](#), which was also used to calculate marginal and conditional R² for all models.
280 Figures and model predictions were generated using ggplot2 [\(Wickham 2009\)](#) and ggeffects [\(Lüdecke](#)
281 [2018\)](#).

282 Community composition

283 We tested whether species composition differed among climbing intensities and abiotic cliff
284 characteristics using multivariate analyses. Analyses were restricted to subplots containing at least
285 three species because extremely low species counts prevented reliable multivariate analysis due to
286 near-zero stress in the NMDS. Community dissimilarities were calculated using Bray–Curtis distances.
287 Patterns in community composition were visualised with non-metric multidimensional scaling (NMDS;

288 k=3) using the ‘metaMDS’ function from the vegan package ([Oksanen et al. 2024](#)). Effects of climbing
289 intensity (categorical; unclimbed, low, high), aspect (categorical; North–South), slope (continuous),
290 and microtopography (continuous) were tested using permutational multivariate analysis of variance
291 (PERMANOVA) implemented with function ‘adonis2’ (999 permutations, marginal tests). This
292 approach evaluates whether multivariate community centroids differ among factor levels or along
293 environmental gradients while accounting for other predictors. Homogeneity of multivariate
294 dispersion was assessed for categorical predictors (climbing intensity and aspect) using ‘betadisper’
295 function followed by permutation-based ANOVA to test for differences in within-group dispersion.
296 NMDS ordinations were visualised using ggplot2.

297 Indicator species were identified using multilevel pattern analysis (function ‘multipat’, 999
298 permutations) from the indicpecies package ([De Cáceres and Legendre 2009](#)). This uses permutations
299 to test the strength of association between each species and level of climbing intensity, without
300 including the additional environmental covariates. In contrast to the multivariate analyses, the
301 indicator species analysis was conducted on the full species matrix.

302 Microclimate

303 Microclimatic variation was assessed using fifteen TOMST thermologgers ([Wild et al. 2019](#)). They
304 measured the air temperature 8 cm in front of the cliff face, and were equipped with the factory-
305 provided single shield. They were placed at different heights on north- and south-facing cliffs for
306 approximately one year. Two sets of three sensors were installed along a vertical gradient on one
307 north-facing and one south-facing cliff; the remaining sensors were distributed across additional cliffs.
308 In total, six sensors were located on north-facing cliffs and nine on south-facing cliffs. Daily mean,
309 minimum, and maximum temperatures were calculated and analysed with linear mixed-effects models
310 to test for seasonal differences between aspects. See Appendix A.3 for full methodological details. The
311 microclimate data were insufficient to be included as an explanatory variable in analyses.

312 All analyses were conducted in R version 4.3.1 (R core team, 2023).

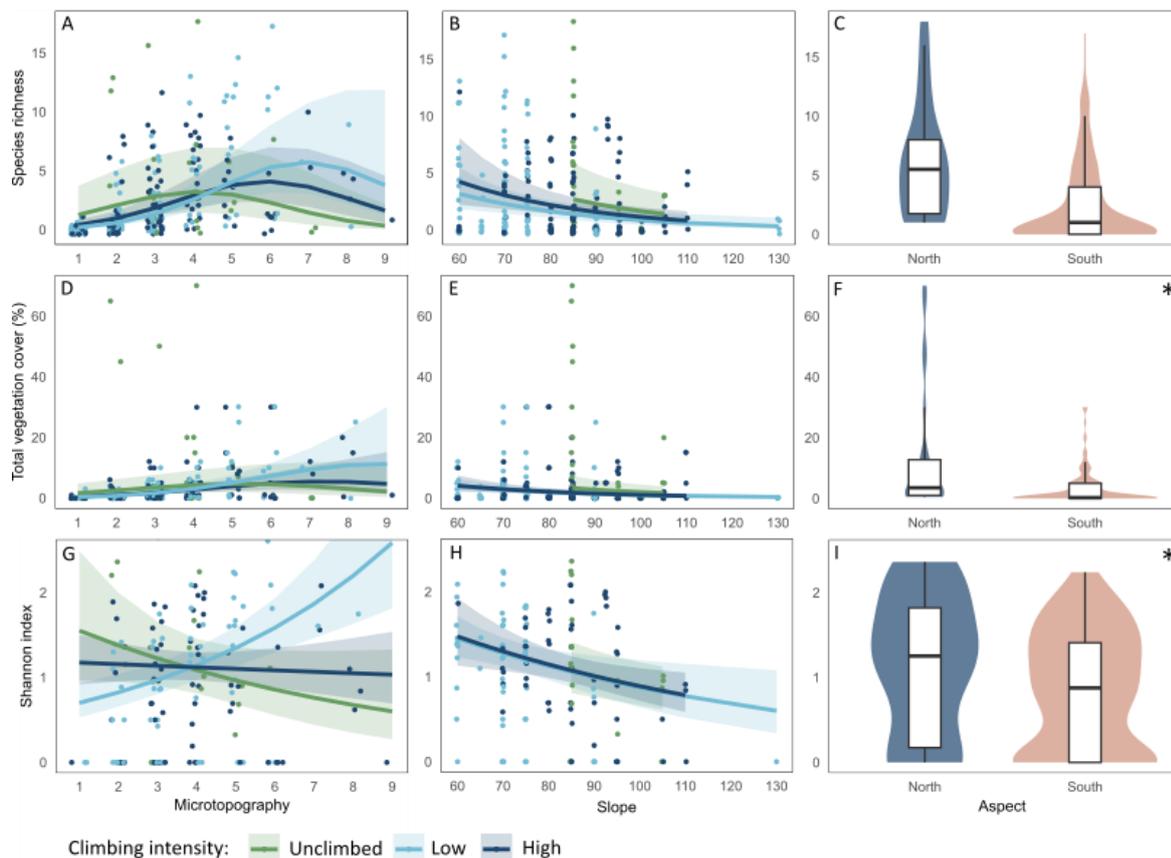
313 RESULTS

314 Alpha diversity

315 Across all plots, we recorded 103 vascular plant species. Species richness, total vegetation cover, and
316 Shannon diversity showed significant interactions between microtopography and climbing intensity
317 (tab. 1). Species richness had a negative quadratic relationship with microtopography, and response
318 varied by climbing intensity. Species richness was higher under low than under high climbing intensity

319 across the microtopographic gradient, whereas unclimbed plots supported lower richness than highly
 320 climbed plots (fig. 2A). Because only a small number of plots combined high microtopographic
 321 complexity with low climbing intensity, we repeated the analyses excluding these plots; the observed
 322 interaction patterns remained qualitatively similar (Appendix A.4). Vegetation cover showed a similar
 323 relationship, where cover was higher with greater microtopography, but the interaction terms with
 324 climbing intensity were smaller in magnitude than those observed for species richness (fig 2D).
 325 Shannon diversity was higher only under low climbing intensity compared to high climbing intensity
 326 (fig. 2G).

327 Besides microtopography, there were differences in vegetation across other abiotic variables
 328 regardless of climbing intensity. Slope had a negative effect on all vegetation metrics: steeper cliffs
 329 consistently supported lower species richness, reduced vegetation cover, and lower Shannon diversity
 330 (fig 2B,E,H). Aspect influenced vegetation cover (fig 2F) and diversity (fig. 2G) but not species richness
 331 (fig. 2C), with south-facing cliffs exhibiting lower cover and reduced Shannon diversity compared to
 332 north-facing cliffs. Patterns for the Simpson index closely matched those observed for the Shannon
 333 index (Appendix A.5.).



334
 335 **Figure 2: : Effects of climbing intensity and abiotic cliff characteristics on alpha diversity metrics. Species richness (A–C),**
 336 **total vegetation cover (D–F), and Shannon diversity index (G–I) are shown in relation to microtopography (A, D, G) and**

337 slope (B, E, H). The number of subplots for species richness and vegetation cover is n=248, for the Shannon index n=150.
 338 Points represent observed values in subplots and are coloured by climbing intensity. Lines show fitted relationships from
 339 generalized linear mixed models (GLMMs), with shaded areas indicating 95% confidence intervals. Panels C, F, and I display
 340 violin plots with embedded boxplots summarizing the distribution (median and interquartile range) of each response
 341 variable by cliff aspect, with significant differences among aspect indicated by *(p<0.05).

342

343 Table 1: Fixed-effect estimates from separate generalised linear mixed models (GLMMs). Cells show the estimate, its p-
 344 value (in parentheses). Significant predictor effects are in bold. Marginal and conditional R² values from the GLMMs
 345 indicate the proportion of variance explained by fixed effects and by both fixed and random effects, respectively.
 346 Continuous predictors were not standardised prior to analysis; estimates are presented on the original measurement scale.

Predictor	Species richness	Total vegetation cover	Shannon index
Intercept	1.440 (p=0.007)	-2.718 (p<0.001)	0.439 (p=0.002)
Microtopography (linear)	0.880 (p<0.001)	0.853 (p<0.001)	-0.027 (p=0.66)
Microtopography (quadratic)	-0.276 (p<0.001)	-0.172 (p=0.003)	-
Slope	-0.458 (p=0.002)	-0.478 (p<0.001)	-0.175 (p=0.01)
Aspect (North/South)	-0.730 (p=0.18)	-1.196 (p=0.01)	-0.321 (p=0.02)
Low climbing intensity	-0.290 (p=0.30)	-0.050 (p=0.86)	-0.126 (p=0.30)
Unclimbed intensity	0.372 (p=0.34)	0.640 (p=0.13)	0.054 (p=0.73)
Microtopography : low climbing	0.331 (p=0.04)	0.291 (p=0.10)	0.305 (p<0.001)
Microtopography : unclimbed	-0.573 (p=0.03)	-0.428 (p=0.010)	-0.175 (p=0.20)
R ² _{Marginal}	0.542	0.369	0.337
R ² _{Conditional}	0.797	0.579	0.575

347

348 CSR strategies

349 The relative abundance (community-weighted means) of plant functional strategies (CSR) was strongly
 350 uneven. Stress-tolerant plants dominated cliff vegetation, while competitive species were less
 351 abundant and ruderal species were least abundant (Appendix 7). Post hoc Tukey tests confirmed that
 352 the relative abundance of all three strategy types differed significantly from each other (Tukey-
 353 adjusted p < 0.001) although the explanatory power of this was low (R² = 0.13).

354 Responses of community-weighted means to climbing intensity and abiotic conditions differed among
 355 strategies (tab. 2, fig. 3). Competitive plants were more abundant on unclimbed subplots than in highly
 356 intensity climbed plots and were more present on south-facing than on north-facing cliffs. Their
 357 relative abundance was higher with increasing microtopographic heterogeneity and lower with
 358 steeper slope (fig 3A-C). Stress-tolerant plants showed a similar but weaker response to climbing
 359 intensity. Their relative abundance was higher in unclimbed than in highly climbed plots, with only a

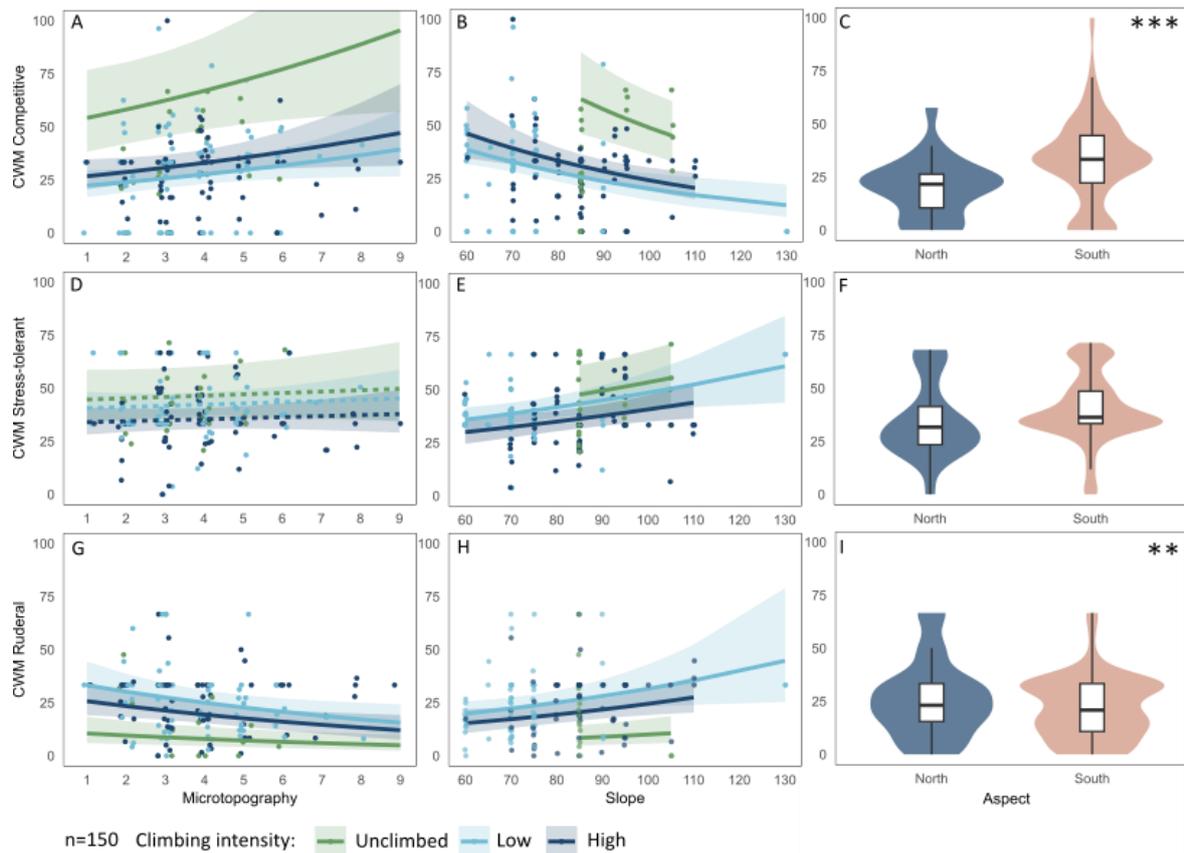
360 marginal, non-significant positive effect under low climbing intensity. In contrast to competitive
 361 species, stress-tolerant plants increased with slope, while microtopography and aspect had no
 362 detectable effects (fig. 3D-F). Ruderals showed the opposite pattern with climbing. Their relative
 363 abundance was lower in unclimbed than in highly climbed plots. Ruderal plants were more abundant
 364 on steeper cliffs, declined with increasing microtopographic heterogeneity, and were less common on
 365 south-facing cliffs (fig.G-I).

366 *Table 2: Drivers of Community-weighted means (CWM) of CSR plant strategies. Fixed-effect estimates from separate*
 367 *generalised linear mixed models (GLMMs) with zero-inflated Tweedie distribution for each strategy-specific CWM. Cells*
 368 *show the estimate, its p-value (in parentheses). Significant values are bolted. Marginal and conditional R² values from*
 369 *the GLMMs indicate the proportion of variance explained by fixed effects and by both fixed and random effects,*
 370 *respectively.*

Predictor	Competitive (C)	Stress-tolerant (S)	Ruderal (R)
Intercept	2.578 (p < 0.001)	3.343 (p < 0.001)	3.731 (p < 0.001)
Microtopography	0.121 (p=0.05)	0.023 (p = 0.55)	-0.161 (p = 0.01)
Slope	-0.225 (<0.001)	0.105 (p = 0.008)	0.161 (p = 0.02)
Aspect (north/south)	0.850 (p <0.001)	0.219 (p = 0.11)	-0.681 (p=0.002)
Low climbing intensity	-0.180 (p=0.16)	0.177 (p = 0.07)	0.255 (p = 0.12)
Unclimbed intensity	0.707 (p <0.001)	0.273 (p = 0.04)	-0.894 (p=0.002)
R ² _{Marginal}	0.226	0.129	0.208
R ² _{Conditional}	0.226	0.189	0.245

371

372



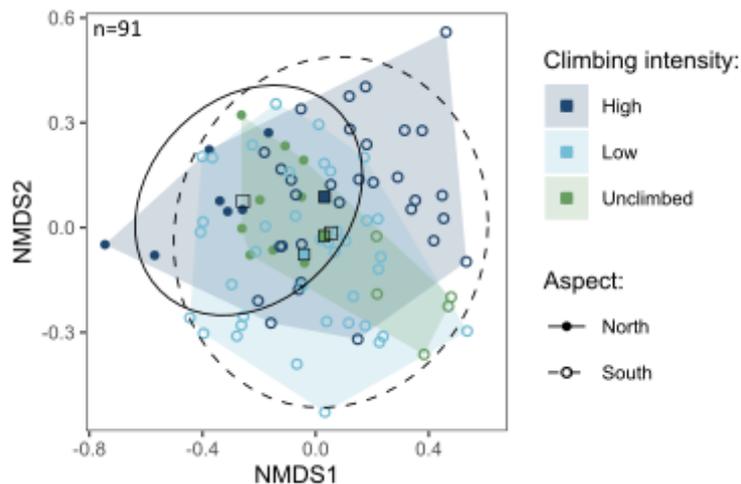
373
 374 **Figure 3: Effects of climbing intensity and abiotic cliff characteristics on the community-weighted mean (CWM) of Grime**
 375 **plant strategies. Competitive (A–C), Stress-tolerant (D–F), and ruderal (G–I) strategies are shown in relation to**
 376 **microtopography (A, D, G) and slope (B, E, H). Points represent observed values and are coloured by climbing intensity.**
 377 **Lines show fitted relationships from generalized linear mixed models (GLMMs), with shaded areas indicating 95%**
 378 **confidence intervals. Panels C, F, and I display violin plots with embedded boxplots summarizing the distribution (median**
 379 **and interquartile range) of each response variable by cliff aspect, with significant differences among aspect indicated by**
 380 ****($p < 0.01$) and ***($p < 0.001$).**

381 Community composition

382 Community composition differed among climbing intensities and abiotic cliff characteristics. Climbing
 383 intensity ($R^2 = 0.06$, $p = 0.001$), North–South aspect ($R^2 = 0.05$, $p = 0.001$), slope ($R^2 = 0.03$, $p = 0.001$),
 384 and microtopography ($R^2 = 0.02$, $p = 0.001$) each explained a significant portion of variation in
 385 community composition (PERMANOVA tests, fig 4). Together, these attributes explained 15.5% of the
 386 total variation. Multivariate dispersion differed between North- and South-facing aspects ($p < 0.001$)
 387 but not among climbing categories ($p = 0.15$). This means that aspect effects reflect both shifts in
 388 centroid and variability, whereas climbing intensity effects are limited to centroid shifts (Beta
 389 dispersion analysis, Figure 4).

390 A total of 19 indicator species of climbing-intensity categories were identified (detailed results table in
 391 Appendix A.6). Only one species was indicative of low climbing intensity and two species were

392 associated with high climbing intensity, whereas, in comparison, sixteen species were associated with
393 unclimbed plots, three of which are threatened and listed on the local red list (*Scabiosa columbaria* L.,
394 *Hieracium murorum* Jord., *Festuca pallens* Host.). In total, six red-listed species were observed across
395 all plots, including *Lactuca perennis* L., *Melampyrum arvense* L., and *Thymus praecox* Opiz.



396

397 **Figure 4: NMDS ordination (3D, stress = 0.18) of species composition across subplots with different climbing intensities and**
398 **cliff aspects. Points represent subplots with at least three species, colored by climbing intensity and filled by aspect.**
399 **Colored convex hulls illustrate the range of sites for each climbing category, while ellipses show 95% confidence intervals**
400 **for North- and South-facing aspects. Squares indicate group centroids; coloured for climbing intensity, empty for aspect.**

401 Microclimate

402 The cliff aspect strongly influenced microclimatic temperatures. South-facing cliffs were generally
403 warmer than north-facing cliffs, but the magnitude of this difference varied seasonally. For daily mean
404 temperatures, differences by aspect were most pronounced and highly significant in summer, moderately significant in spring and fall, and small and non-significant in winter. Daily maximum
405 temperatures differed significantly in all seasons, and daily minimum temperatures were significantly
406 higher on south-facing cliffs in summer (Appendix A.2).
407

408 DISCUSSION

409 We surveyed vascular plant communities on Freyr's limestone cliffs in Belgium to examine how
410 recreational rock climbing influences cliff vegetation across a range of abiotic cliff characteristics.
411 Climbing intensity alone did not show strong main effects on alpha diversity, but was associated with
412 shifts in functional structure and species composition and the effects depended on microtopographic
413 heterogeneity. Together, these results suggested that the ecological effects of rock climbing can be
414 subtle, expressed primarily through species turnover rather than reductions in vegetation cover, and
415 strongly shaped by abiotic constraints.

416 Effects of rock climbing on vegetation diversity, functional structure, and species 417 composition

418 Rock climbing did not lead to uniform reductions in richness, cover, or diversity; instead, it primarily
419 altered community composition and functional structure. The observed changes in diversity patterns
420 due to climbing intensity were underpinned by shifts in the relative dominance of plant functional
421 strategies and species composition.

422 Species composition on unclimbed cliffs likely reflected a combination of historical disturbance
423 regimes and strong microhabitat filtering. Unclimbed cliffs had both a higher relative abundance of
424 competitors (C-strategy) and stress-tolerant (S-strategy) species in comparison with low and high level
425 intensity climbed cliffs. Within this low-disturbance context, competitive species may dominate locally,
426 leading to reduced diversity through competitive exclusion, while stress-tolerant species persist in
427 harsher micro-sites (e.g., steeper slope, less microtopography). Competitive species associated with
428 unclimbed areas included *Sesleria caerulea* (L.) Ard. and *Silene nutans* L.. These areas also hosted
429 stress-tolerant taxa such as *Umbilicus rupestris* (Salisb.) Dandy, *Carex humilis* Leyss., *Erysimum cheiri*
430 (L.) Crantz, *Sedum album* L., and threatened (red-listed) species, including *Scabiosa columbaria* L.,
431 *Hieracium murorum* L., and *Festuca pallens* L., which were not significantly associated as indicator
432 species with climbed areas. This finding indicates a potential loss of vulnerable vegetation due to
433 climbing. Although not identical, these results reflect a pattern partly in line with Müller et al. (2004),
434 who found that specialised rock species—often stress-tolerant—occurred less frequently on climbed
435 cliffs than on unclimbed cliffs in Switzerland.

436 Low-intensity climbing areas hosted a mixture of species with competitive, stress-tolerant, or ruderal
437 strategies. Moderate disturbance reduced local competitive dominance and resulted in a peak in
438 species richness, while stress-tolerant species remained present, but several that were associated with
439 unclimbed cliffs in the indicator species analysis no longer showed this association. This indicates that,
440 even where richness is highest, low-intensity climbing can potentially reduce community uniqueness.

441 Highly climbed cliffs supported disturbance-driven communities, where frequent biomass removal
442 created open space for ruderal and opportunistic species. Indicator species associated with high
443 climbing intensity, such as *Taraxacum sp.*, demonstrated that these communities favor disturbance-
444 adapted taxa. This is similar to literature, potentially at the expense of more specialised taxa, which
445 are replaced (Radujković et al. 2025). In contrast, March-Salas et al. (2018) found that fewer generalist
446 species occurred on climbed cliffs, while specialised and moderately specialized rock species were not
447 significantly affected by climbing intensity.

448 Only one and two indicator species were associated with low- and high-intensity climbing, respectively,
449 compared to 16 species associated with unclimbed cliffs. This pattern suggests that climbing
450 disturbance may reduce the complexity and uniqueness of cliff communities. ([Radujković et al. 2025](#)).
451 However, the three climbing intensity levels were unevenly represented (unclimbed = 28; low = 116;
452 high = 104). The lower number of unclimbed plots may have reduced within-group variation, increasing
453 the likelihood of detecting significant indicator species. However, overall species richness did not differ
454 among climbing levels. Overall, stress-tolerant species dominated in cliff ecosystems due to harsh
455 conditions ([Larson et al. 2005](#)). These shifts in functional strategies explained the observed patterns in
456 alpha-diversity, showing that climbing primarily affects species composition rather than vegetation
457 cover.

458 **Abiotic constraints**

459 Abiotic factors remained the dominant constraint on cliff vegetation, shaping patterns of richness,
460 cover, diversity, and functional composition.

461 Slope acted as a strong environmental filter. Steeper cliffs had lower vascular plant richness, cover,
462 and Shannon diversity, likely due to gravitational instability, reduced water retention, and limited soil
463 and nutrient accumulation. These effects were consistent across climbing intensities, suggesting that
464 slope constraints vegetation more strongly than climbing disturbance. This finding is similar to existing
465 studies ([Bartlett et al. 1990](#); [Clark and Hessler 2015](#); [Larson et al. 2005](#)). Ruderal and stress-tolerant
466 plants were more abundant on steeper slopes, while competitive plants declined, consistent with the
467 expectation that steep, harsh environments favour disturbance- and stress-adapted species ([Kraft et
468 al. 2015](#)), and competitive plants are disadvantaged in resource-poor, stressful environments ([Grime
469 1977](#)).

470 Aspect did not affect species richness, suggesting that both north- and south-facing cliffs offer
471 sufficient habitat heterogeneity for species persistence. This result is consistent with findings by
472 March-Salas et al. ([2018](#)) and Graham and Knight ([2004](#)), whereas Boggess ([2017](#)) reported the highest
473 vascular plant diversity on west-facing slopes. In contrast, vegetation cover was higher on north-facing
474 cliffs, likely reflecting cooler and more humid microclimatic conditions that promote greater plant
475 growth and biomass accumulation. NMDS results further indicated that north-facing communities
476 were compositionally less variable, suggesting stronger environmental filtering under cooler, moister
477 conditions. These findings align with Müller et al. (2004), who reported higher plant cover on shaded
478 cliffs. South-facing cliffs, by contrast, are warmer and drier, which constrains plant growth and likely
479 results in lower overall cover and more patchy vegetation, thereby reducing evenness and Shannon
480 diversity. On North faces, species with a ruderal plant strategy were more abundant, on South faces,

481 competitive strategies were more prevalent, though it is not clear what is driving this pattern.
482 Additional processes and environmental drivers likely contribute to this trend but are not captured in
483 our models of relative CSR strategy abundance, as indicated by their low R^2 values. Potential
484 interactions with climbing intensity, due to route selection, may also influence these patterns,
485 although limited data across aspects and climbing levels prevent further testing.

486 Microtopography, in our study area, was the main source of heterogeneity and availability of
487 microhabitats. Species richness and vegetation cover showed a hump-shaped response to
488 microtopography, with the highest values at intermediate levels of structural complexity. On smooth
489 cliff surfaces, plant establishment is constrained by the limited availability of suitable microhabitats,
490 whereas higher microtopographic complexity initially increases the number of microhabitats with
491 accumulation of soil, nutrients, moisture, and propagules ([Larson et al. 2005](#); [Kuntz and Larson 2006a](#);
492 [Kuntz and Larson 2006b](#)). Similar positive effects of microtopographic complexity on cliff vegetation
493 have been reported in numerous studies ([Clark and Hessel 2015](#); [Farris 1998](#); [March-Salas et al. 2018](#);
494 [Harrison et al. 2022](#); [Müller et al. 2004](#); [Nuzzo 1996](#)). As such, at lower microtopographic complexity,
495 the pattern is consistent with the habitat heterogeneity hypothesis, which predicts that environmental
496 complexity promotes species coexistence through niche differentiation, as species with differing
497 requirements can exploit varying conditions ([Graham and Duda 2011](#); [Stein et al. 2014](#)).

498 At high levels of microtopographic complexity, however, species richness declined again. Although the
499 number of plots representing the highest levels of microtopographic complexity was limited in this
500 study, one plausible explanation is increased competitive exclusion under locally less stressful
501 conditions, as indicated by the positive relationship between microtopography and the community-
502 weighted mean of competitive plant strategies. In addition, inspection of plot photographs revealed
503 that several high-complexity plots with low richness were characterised by pronounced blocky
504 overhangs or strong dominance by a single species, suggesting that suggesting that extremely complex
505 surfaces may become unsuitable for plant establishment when structural overhangs reduce substrate
506 stability or light availability, consistent with the strong negative effects of slope on vegetation observed
507 in this study.

508 Shannon diversity did not vary with microtopography, likely because this metric is influenced by
509 species evenness ([Shannon 1948](#)). Although greater microtopographic complexity can increase the
510 number of available microhabitats and species, dominance by a few competitive taxa may persist or
511 even increase, limiting changes in evenness.

512 **Climbing as context dependent disturbance mediated by habitat heterogeneity**
513 Species richness peaked at a low climbing intensity level, but only under conditions of high
514 microtopography, whereas both unclimbed and high-intensity climbed areas exhibited lower diversity.
515 This interaction indicates that the ecological effects of climbing depend strongly on underlying habitat
516 heterogeneity. These patterns are consistent with the Intermediate Disturbance Hypothesis ([Connell](#)
517 [1978](#); [Fox 1981](#)), but only under structurally heterogeneous conditions, suggesting that habitat
518 complexity mediates whether disturbance promotes coexistence. Together, our results indicate that
519 climbing does not impose a uniform pressure across cliffs, but instead interacts with abiotic constraints
520 that already structure these ecosystems.

521 **Future directions**

522 Although climbing and abiotic variables significantly influenced diversity and species composition,
523 most variation in community composition remained unexplained. Climbing disturbance and abiotic cliff
524 characteristics explained only ~15.5% of variation in community composition, with climbing intensity
525 explaining 6% specifically. This low explanatory power suggests that cliff plant communities may
526 additionally be shaped by factors not captured in this study, potentially finer-scale spatial processes
527 than we assessed.

528 Further research could consider detailed drivers, including substrate and soil analyses, as well as UAV-
529 derived digital elevation models combined with thermal imaging for microclimates to improve
530 understanding of microhabitat conditions ([Metsu et al. 2026](#)). Drone-based vegetation surveys could
531 also improve data collection in hazardous or difficult-to-access cliff areas ([Nyberg et al. 2024](#)). This
532 study focused on vascular plants, but cliff ecosystems also host mosses and lichens and include the cliff
533 edge and base, which may respond differently to climbing disturbance (Larson et al. 2005; Boggess et
534 al. 2021). Including these additional taxa and zones would provide a more comprehensive
535 understanding of climbing impacts at the ecosystem scale.

536 Multi-site studies across different climates, rock types, and management regimes, and long-term
537 monitoring will help to understand and assess the generality of the observed patterns ([Boggess et al.](#)
538 [2021](#)). Using standardised methodology and global networks such as MIREN Rocks could facilitate this
539 comparability among studies. This further research is essential to better understand and protect these
540 unique cliff ecosystems and to develop science-based strategies that balance outdoor recreation with
541 biodiversity conservation.

542 CONCLUSION

543 Our findings showed that the ecological impacts of climbing in Freyr were complex and context-
544 dependent. Climbing influenced species and community composition under certain habitat conditions
545 rather than causing an overall loss of vegetation cover or decline in species richness. Low-intensity
546 climbing was associated with higher biodiversity under highly heterogeneous microtopography,
547 potentially through reduced competitive dominance and increased representation of ruderal
548 strategies. However, we also observed a lower abundance of stress-tolerant and vulnerable plant
549 species in climbed areas.

550 Vegetation was further shaped by microhabitat availability determined by abiotic cliff characteristics.
551 This indicates that more detailed analyses of environmental factors, such as microtopography and
552 microclimate, are needed to fully understand the underlying ecological processes. This abiotic
553 importance and site-specific results were consistent with previous research.

554 These results have important implications for the conservation and management of cliff habitats and
555 climbing areas. They highlight the conservation value of unclimbed cliffs, which support typical stress-
556 tolerant and red-list species and should therefore be prioritised for protection to safeguard these
557 vulnerable populations. Previous studies made similar recommendations for the protection of
558 climbing-free zones ([Harrison et al. 2022](#); [Müller et al. 2004](#); [Lorite et al. 2017](#); [Camp and Knight 1998](#);
559 [McMillan and Larson 2002](#)).

560 Overall, these findings suggest that recreational rock climbing can be compatible with biodiversity
561 conservation when appropriately managed, while emphasising the importance of maintaining some
562 cliffs free from climbing to protect sensitive vegetation.

563 REFERENCES

- 564 27 Craggs. n.d. "Freyr." Accessed February 16, 2026. <https://27craggs.com/craggs/freyr>.
- 565 Aubel, Olivier, and Brice Lefèvre. 2022. "What Climbing Means...." *Journal of Outdoor Recreation and*
566 *Tourism* 40 (100585): 100585.
- 567 Bartlett, R. M., U. Matthes-Sears, and D. W. Larson. 1990. "Organization of the Niagara Escarpment
568 cliff community. II. Characterization of the physical environment." *Canadian journal of botany. Journal*
569 *canadien de botanique* 68 (9): 1931–1941.
- 570 Bates, Douglas, Martin Mächler, Ben Bolker, and Steve Walker. 2015. "Fitting Linear Mixed-Effects
571 Models Using lme4." *Journal of Statistical Software* 67 (1). <https://doi.org/10.18637/jss.v067.i01>.

572 Boggess, Laura M., Georgia R. Harrison, and Giovanna Bishop. 2021. "Impacts of Rock Climbing on Cliff
573 Vegetation: A Methods Review and Best Practices." *Applied Vegetation Science* 24 (2).
574 <https://doi.org/10.1111/avsc.12583>.

575 Boggess, Laura M., Gary L. Walker, and Michael D. Madritch. 2017. "Cliff Flora of the Big South Fork
576 National River and Recreation Area." *Natural Areas Journal* 37 (2): 200.

577 Bott, M. 2022. Freyr.

578 Bramer, Isobel, Barbara J. Anderson, Jonathan Bennie, et al. 2018. "Advances in Monitoring and
579 Modelling Climate at Ecologically Relevant Scales." In *Next Generation Biomonitoring: Part 1. Advances*
580 *in Ecological Research*. Elsevier.

581 Brooks, Mollie E., Kasper Kristensen, Koen J. van Benthem, et al. 2017. "glmmTMB Balances Speed and
582 Flexibility Among Packages for Zero-Inflated Generalized Linear Mixed Modeling." *The R Journal* 9 (2):
583 378–400.

584 Calvin, Katherine, Dipak Dasgupta, Gerhard Krinner, et al. 2023. IPCC, 2023: Climate Change 2023:
585 Synthesis Report. Contribution of Working Groups I, II and III to the Sixth Assessment Report of the
586 Intergovernmental Panel on Climate Change [core Writing Team, H. Lee and J. Romero (eds.)]. IPCC,
587 Geneva, Switzerland. Paola Arias et al. Intergovernmental Panel on Climate Change.
588 <https://doi.org/10.59327/ipcc/ar6-9789291691647>.

589 Camp, Richard J., and Richard L. Knight. 1998. "Effects of Rock Climbing on Cliff Plant Communities at
590 Joshua Tree National Park, California." *Conservation Biology: The Journal of the Society for*
591 *Conservation Biology* 12 (6): 1302–1306.

592 Capó, Miquel, Iván Cortés-Fernández, and Joshua Borràs. 2023. "Spatial Distribution of Insular Cliff
593 Vegetation and Future Scenarios in a Climate Change Perspective." In *bioRxiv*. November 13.
594 <https://doi.org/10.1101/2023.11.09.566357>.

595 Cartwright, Jennifer. 2019. "Ecological Islands: Conserving Biodiversity Hotspots in a Changing
596 Climate." *Frontiers in Ecology and the Environment* 17 (6): 331–340.

597 Clark, Peter, and Amy Hessler. 2015. "The Effects of Rock Climbing on Cliff-face Vegetation." *Applied*
598 *Vegetation Science* 18 (4): 705–715.

599 Connell, J. H. 1978. "Diversity in Tropical Rain Forests and Coral Reefs." *Science (New York, N.Y.)* 199
600 (4335): 1302–1310.

601 Datar, Mandar N., and Aparna V. Watve. 2018. "Vascular Plant Assemblage of Cliffs in Northern
602 Western Ghats, India." *Journal of Threatened Taxa* 10 (2): 11271.

603 De Cáceres, Miquel, and Pierre Legendre. 2009. "Associations between Species and Groups of Sites:
604 Indices and Statistical Inference." *Ecology* 90 (12): 3566–3574.

605 De Frenne, Pieter, Rémy Beugnon, David Klinges, et al. 2025. "Ten Practical Guidelines for Microclimate
606 Research in Terrestrial Ecosystems." *Methods in Ecology and Evolution* 16 (2): 269–294.

607 De Frenne, Pieter, Francisco Rodríguez-Sánchez, David Anthony Coomes, et al. 2013. "Microclimate
608 Moderates Plant Responses to Macroclimate Warming." *Proceedings of the National Academy of
609 Sciences of the United States of America* 110 (46): 18561–18565.

610 De Pryck, Kari. 2021. "Intergovernmental Expert Consensus in the Making: The Case of the Summary
611 for Policy Makers of the IPCC 2014 Synthesis Report." *Global Environmental Politics* 21 (1): 108–129.

612 Dobrowski, Solomon Z. 2011. "A Climatic Basis for Microrefugia: The Influence of Terrain on Climate."
613 *Global Change Biology* 17 (2): 1022–1035.

614 Ellenberg, Heinz H. 2009. *Vegetation Ecology of Central Europe*. 4th ed. Gordon K. Strutt. Cambridge
615 University Press.

616 Farris, Michael A. 1998. "The Effects of Rock Climbing on the Vegetation of Three Minnesota Cliff
617 Systems." *Canadian Journal of Botany. Journal Canadien de Botanique* 76 (12): 1981–1990.

618 Fitzsimons, James A., and Damian R. Michael. 2017. "Rocky Outcrops: A Hard Road in the Conservation
619 of Critical Habitats." *Biological Conservation* 211 (July): 36–44.

620 Fox, John F. 1981. "Intermediate Levels of Soil Disturbance Maximize Alpine Plant Diversity." *Nature*
621 293 (5833): 564–565.

622 "Freyr • Belgium - Sportclimbing - Vertical-Life Climbing Crags." n.d. Accessed February 16, 2026.
623 <https://www.8a.nu/crags/sportclimbing/belgium/freyr/routes>.

624 García, María B., Darío Domingo, Manuel Pizarro, Xavier Font, Daniel Gómez, and Johan Ehrlén. 2020.
625 "Rocky Habitats as Microclimatic Refuges for Biodiversity. A Close-up Thermal Approach."
626 *Environmental and Experimental Botany* 170 (103886): 103886.

627 Geiger, Rudolf, Robert H. Aron, and Paul Todhunter. 2012. *The Climate near the Ground*.
628 Vieweg+Teubner Verlag.

629 Graham, John H., and Jeffrey J. Duda. 2011. "The Humpbacked Species Richness-Curve: A Contingent
630 Rule for Community Ecology." *International Journal of Ecology* 2011: 1–15.

631 Graham, Liza, and Richard L. Knight. 2004. "Multi-Scale Comparisons of Cliff Vegetation in Colorado."
632 *Plant Ecology* 170 (2): 223–234.

633 Grime, J. P. 1977. "Evidence for the Existence of Three Primary Strategies in Plants and Its Relevance
634 to Ecological and Evolutionary Theory." *The American Naturalist* 111 (982): 1169–1194.

635 Harrison, Georgia R., Laura M. Boggess, Jessica M. Budke, and Michael D. Madritch. 2022. "Rock-
636 climbing Shifts Cliff-face Vegetation Community Composition Based on Site Characteristics." *Applied*
637 *Vegetation Science* 25 (2). <https://doi.org/10.1111/avsc.12667>.

638 Harrison, Georgia R., Laura M. Boggess, Sarah E. McCord, and Martí March-Salas. 2024. "A Call to
639 Action for Inventorying and Monitoring of Cliff Ecosystems to Support Conservation." *Basic and*
640 *Applied Ecology* 80 (November): 31–39.

641 Hartig, Florian. 2024. "Residual Diagnostics for Hierarchical (Multi-Level / Mixed) Regression Models
642 [R Package DHARMA Version 0.4.7]." *Comprehensive R Archive Network (CRAN)*, October 18.
643 <https://CRAN.R-project.org/package=DHARMA>.

644 Holzschuh, Andrea. 2016. "Does Rock Climbing Threaten Cliff Biodiversity? - A Critical Review."
645 *Biological Conservation* 204 (December): 153–162.

646 Hooper, David U., E. Carol Adair, Bradley J. Cardinale, et al. 2012. "A Global Synthesis Reveals
647 Biodiversity Loss as a Major Driver of Ecosystem Change." *Nature* 486 (7401): 105–108.

648 IPBES. 2019. "Global Assessment Report on Biodiversity and Ecosystem Services of the
649 Intergovernmental Science-Policy Platform on Biodiversity and Ecosystem Services." Preprint, Zenodo.
650 <https://doi.org/10.5281/ZENODO.3831673>.

651 Ivanova, Natalya, and Ekaterina Zolotova. 2023. "Landolt Indicator Values in Modern Research: A
652 Review." *Sustainability* 15 (12): 9618.

653 Keppel, Gunnar, Karel Mokany, Grant W. Wardell-Johnson, Ben L. Phillips, Justin A. Welbergen, and
654 April E. Reside. 2015. "The Capacity of Refugia for Conservation Planning under Climate Change."
655 *Frontiers in Ecology and the Environment* 13 (2): 106–112.

656 Kraft, Nathan J. B., Peter B. Adler, Oscar Godoy, Emily C. James, Steve Fuller, and Jonathan M. Levine.
657 2015. "Community Assembly, Coexistence and the Environmental Filtering Metaphor." *Functional*
658 *Ecology* 29 (5): 592–599.

659 Kuntz, Kathryn Lynne, and Douglas W. Larson. 2006a. "Influences of Microhabitat Constraints and
660 Rock-Climbing Disturbance on Cliff-Face Vegetation Communities." *Conservation Biology: The Journal*
661 *of the Society for Conservation Biology* 20 (3): 821–832.

662 Kuntz, Kathryn Lynne, and Douglas W. Larson. 2006b. "Microtopographic Control of Vascular Plant,
663 Bryophyte and Lichen Communities on Cliff Faces." *Plant Ecology* 185 (2): 239–253.

664 Larson, Douglas W., Uta Matthes, and Peter E. Kelly. 2005. *Cambridge Studies in Ecology: Cliff Ecology:*
665 *Pattern and Process in Cliff Ecosystems: Pattern and Process in Cliff Ecosystems.* Cambridge Studies in
666 Ecology. Cambridge University Press.

667 Lembrechts, Jonas J., and Ivan Nijs. 2020. "Microclimate Shifts in a Dynamic World." *Science (New York,*
668 *N.Y.)* 368 (6492): 711–712.

669 Lenth, Russell V., and Julia Piaskowski. 2017. "Emmeans: Estimated Marginal Means, Aka Least-Squares
670 Means." In *CRAN: Contributed Packages.* The R Foundation, October 20.
671 <https://doi.org/10.32614/cran.package.emmeans>.

672 Lorite, Juan, Fabio Serrano, Adrián Lorenzo, Eva M. Cañadas, Miguel Ballesteros, and Julio Peñas. 2017.
673 "Rock Climbing Alters Plant Species Composition, Cover, and Richness in Mediterranean Limestone
674 Cliffs." *PloS One* 12 (8): e0182414.

675 Lüdecke, Daniel. 2018. "Ggeffects: Tidy Data Frames of Marginal Effects from Regression Models."
676 *Journal of Open Source Software* 3 (26): 772.

677 Lüdecke, Daniel, Mattan Ben-Shachar, Indrajeet Patil, Philip Waggoner, and Dominique Makowski.
678 2021. "Performance: An R Package for Assessment, Comparison and Testing of Statistical Models."
679 *Journal of Open Source Software* 6 (60): 3139.

680 Maclean, Ilya M. D., James P. Duffy, Stef Haesen, et al. 2021. "On the Measurement of Microclimate."
681 *Methods in Ecology and Evolution* 12 (8): 1397–1410.

682 March-Salas, Martí, María Begoña García, Isaac H. Lichter-Marck, Juan Lorite, and Manuel J.
683 Steinbauer. 2025. "Cliff Ecosystems: A Critical yet Uncharted Frontier for Research and Conservation."
684 *Basic and Applied Ecology* 83 (March): 36–42.

685 March-Salas, Martí, Juan Lorite, Oliver Bossdorf, and J. F. Scheepens. 2023. "Cliffs as Priority
686 Ecosystems." *Conservation Biology: The Journal of the Society for Conservation Biology* 37 (5): e14166.

687 March-Salas, Martí, Miguel Moreno-Moya, Gemma Palomar, Pablo Tejero-Ibarra, Emily Haeuser, and
688 Luis R. Pertierra. 2018. "An Innovative Vegetation Survey Design in Mediterranean Cliffs Shows

689 Evidence of Higher Tolerance of Specialized Rock Plants to Rock Climbing Activity.” *Applied Vegetation*
690 *Science* 21 (2): 289–297.

691 McMillan, Michele A., and Douglas W. Larson. 2002. “Effects of Rock Climbing on the Vegetation of the
692 Niagara Escarpment in Southern Ontario, Canada.” *Conservation Biology: The Journal of the Society*
693 *for Conservation Biology* 16 (2): 389–398.

694 Metsu, Christophe, Wouter H. Maes, Sam Ottoy, and Koenraad Van Meerbeek. 2026. “theRmalUAV:
695 An R Package to Clean and Correct Thermal UAV Data for Accurate Land Surface Temperatures.”
696 *Methods in Ecology and Evolution* 17 (2): 488–496.

697 Morales-Armijo, Felipe, Andrea Sobrevilla-Covarrubias, Eduardo Estrada-Castillón, et al. 2024.
698 “Climbing Route Development Affects Cliff Vascular Plants More than Subsequent Climbing: A Guide
699 to Evidence-based Conservation Management to Regulate Climbing.” *The Journal of Applied Ecology*
700 61 (11): 2679–2689.

701 Müller, Stefan W., Hans-Peter Rusterholz, and Bruno Baur. 2004. “Rock Climbing Alters the Vegetation
702 of Limestone Cliffs in the Northern Swiss Jura Mountains.” *Canadian Journal of Botany. Journal*
703 *Canadien de Botanique* 82 (6): 862–870.

704 Newton, Adrian C., Robin M. Walls, Duncan Golicher, Sally A. Keith, Anita Diaz, and James M. Bullock.
705 2012. “Structure, Composition and Dynamics of a Calcareous Grassland Metacommunity over a 70-
706 Year Interval.” *The Journal of Ecology* 100 (1): 196–209.

707 Nuzzo, Victoria A. 1996. “Structure of Cliff Vegetation on Exposed Cliffs and the Effect of Rock
708 Climbing.” *Canadian Journal of Botany. Journal Canadien de Botanique* 74 (4): 607–617.

709 Nyberg, Ben, Célia Bairos, Marcela Brimhall, et al. 2024. “The Conservation Impact of Botanical Drones:
710 Documenting and Collecting Rare Plants from Vertical Cliffs and Other Hard-to-Reach Areas.”
711 *Ecological Solutions and Evidence* 5 (1). <https://doi.org/10.1002/2688-8319.12318>.

712 Oksanen, J., G. Simpson, F. Blanchet, et al. 2024. `_vegan: Community Ecology Package_`. (Stier A, Ter
713 Braak C, Weedon J).

714 R Core Team (2023). `_R: A Language and Environment for Statistical Computing_`. R Foundation for
715 Statistical Computing, Vienna, Austria. <<https://www.R-project.org/>>.

716 Radujković, Dajana, Erik Verbruggen, Jan Clavel, et al. 2025. “Road Disturbance Shifts Root Fungal
717 Symbiont Types and Reduces the Connectivity of Plant-Fungal Co-Occurrence Networks in Mountains.”
718 *Molecular Ecology* 34 (20): e17771.

719 Rusterholz, Hans-Peter, Christine Verhoustraeten, and Bruno Baur. 2011. "Effects of Long-Term
720 Trampling on the above-Ground Forest Vegetation and Soil Seed Bank at the Base of Limestone Cliffs."
721 *Environmental Management* 48 (5): 1024–1032.

722 Schmera, Dénes, Hans-Peter Rusterholz, Anette Baur, and Bruno Baur. 2018. "Intensity-Dependent
723 Impact of Sport Climbing on Vascular Plants and Land Snails on Limestone Cliffs." *Biological*
724 *Conservation* 224 (August): 63–70.

725 Shannon, C. E. 1948. "A Mathematical Theory of Communication." *The Bell System Technical Journal*
726 27 (3): 379–423.

727 Speziale, Karina L., and Cecilia Ezcurra. 2015. "Rock Outcrops as Potential Biodiversity Refugia under
728 Climate Change in North Patagonia." *Plant Ecology & Diversity* 8 (3): 353–361.

729 Stein, Anke, Katharina Gerstner, and Holger Kreft. 2014. "Environmental Heterogeneity as a Universal
730 Driver of Species Richness across Taxa, Biomes and Spatial Scales." *Ecology Letters* 17 (7): 866–880.

731 Suggitt, Andrew J., Phillipa K. Gillingham, Jane K. Hill, et al. 2011. "Habitat Microclimates Drive Fine-
732 scale Variation in Extreme Temperatures." *Oikos* (Copenhagen, Denmark) 120 (1): 1–8.

733 theCrag. n.d. "Rochers de Freyr, Sport Climbing." Accessed February 16, 2026.
734 <https://www.thecrag.com/en/climbing/belgium/freyr>.

735 Wickham, Hadley. 2009. *Ggplot2: Elegant Graphics for Data Analysis*. Springer.

736 Wild, Jan, Martin Kopecký, Martin Macek, Martin Šanda, Jakub Jankovec, and Tomáš Haase. 2019.
737 "Climate at Ecologically Relevant Scales: A New Temperature and Soil Moisture Logger for Long-Term
738 Microclimate Measurement." *Agricultural and Forest Meteorology* 268 (April): 40–47.

Supplementary materials

1

2 A.1. MIREN Rocks Survey protocol

3 A.2. Climbing intensity

4 A central challenge of this study was to quantify climbing intensity in a way that allows an accurate
5 assessment of its ecological impacts. In previous studies, climbing impacts were often evaluated by
6 comparing climbed and unclimbed areas using paired or unpaired transects ([Boggess et al. 2021](#)). At
7 Freyr, however, identifying a sufficient number of comparable unclimbed control zones was either
8 impractical or unsafe. Consequently, a gradient-based measure of climbing intensity was considered
9 more appropriate than a binary climbed/unclimbed classification.

10 Exact empirical data on how much each route is climbed were not available and are difficult to collect.
11 Therefore, climbing intensity had to be estimated using indirect indicators (proxies). Several such
12 proxies have been proposed in previous research, including the Climbing Use Index (CUI) ([Clark and
13 Hessel 2015](#)), guidebook information, online ascent logs, and expert judgement. Each of these
14 approaches has inherent limitations. To address this, a multifaceted approach was adopted to evaluate
15 and compare different indicators and to identify the most suitable method for estimating climbing
16 intensity at the study site.

17 Climbing use index (CUI)

18 As a first step, the Climbing Use Index ([Clark and Hessel 2015](#)) was calculated for the routes in Freyr.
19 The CUI combines route popularity, expressed as guidebook star ratings, with walking approach time
20 (Formula X). Because the Freyr guidebook does not provide star ratings, ratings were obtained from
21 the online platform [8a.nu \(Freyr • Belgium - Sportclimbing - Ver...\)](#). Approach times were taken from
22 the most recent Freyr guidebook (Bott & Deneyer, 2022).

23 This index was developed in a large area (New River Gorge, USA) with substantial variation in
24 approaches (2-50 min on foot). In such settings, longer approach times are expected to generally
25 correspond to lower visitation. However, in Freyr, a compact site with more uniform approach times
26 (8-18 min), CUI ranged from 1.02 to 1.78 (range = 0.76) only, a substantially more narrow range than
27 in New River Gorge (0.31-2.22; range = 1.91) (fig. 3). Unlike New River Gorge, the distribution of CUI
28 values in Freyr was also not normal, as indicated by the Shapiro-Wilk test ($W=0.840$, $p < 0.001$). This
29 narrower range and non-normal distribution
30 limited its effectiveness as a climbing intensity
31 proxy. Furthermore, the star ratings, either
32 from guidebooks or 8a.nu, are a rather arbitrary
33 estimate because they reflect climber's
34 appreciation of a route rather than usage
35 intensity. The correlation between appreciation
36 and intensity is uncertain, and these ratings lack
37 clear, objective criteria as well as transparency
38 regarding who assigned them. Both
39 components of the CUI—approach time and
40 star ratings—are thus either questionable or
41 unsuitable for Freyr, necessitating alternative
42 measures of climbing intensity.

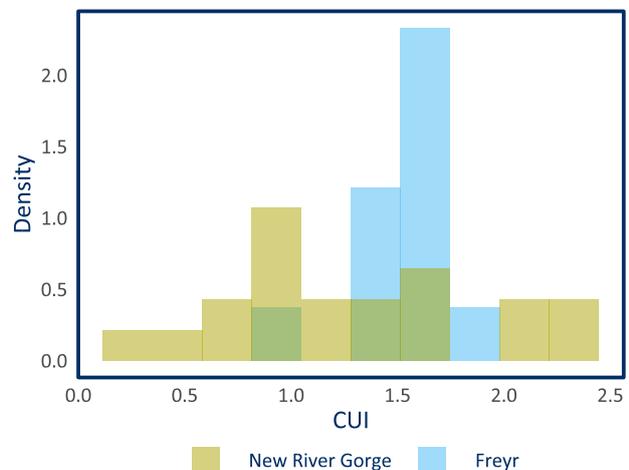


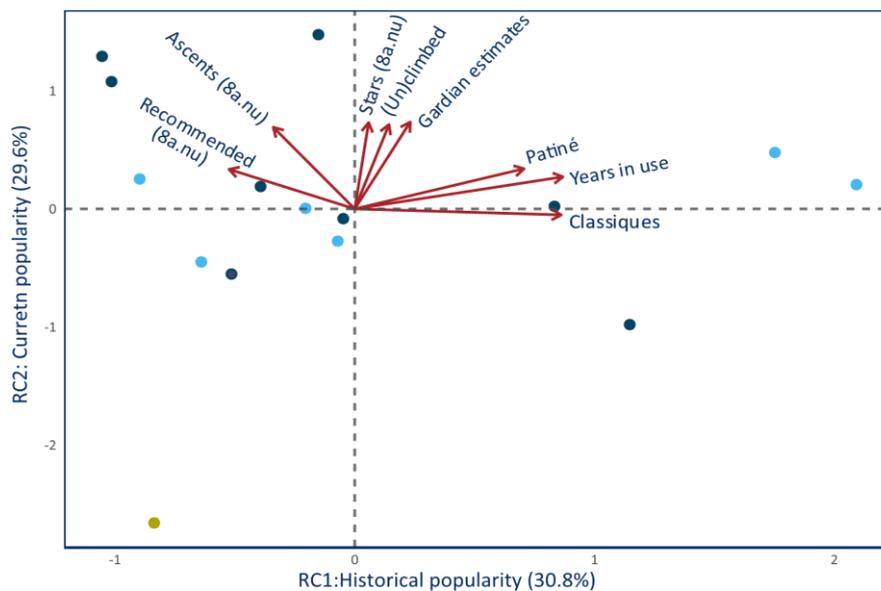
Figure A.2.1: Histogram of Climbing Use Index (CUI) distributions for New River Gorge ($n = 20$) and Freyr ($n = 14$). The data are grouped into 10 bins with a bin width of 0.2.

43
$$CUI = LOG_{10} \left([S + 1] * \left(\left[\frac{1}{T} \right] * 100 \right) \right)$$

44 Indicators for climbing intensity

45 Given the limitations of the CUI, additional indicators of climbing intensity were compiled. These
46 included guidebook information (year of route opening; classification as *classique* (indication as “must
47 climb” popular route) or *patiné* (indicating surface polishing due to frequent climbing)), ascent logs
48 from 8a.nu (number of ascents, recommendation percentage, and star ratings), and climbing intensity
49 estimates provided by the site guardian.

50 To combine these indicators, a Principal Component Analysis (PCA) was conducted to derive
51 composite measures of climbing intensity. Two main components explaining variation in climbing
52 intensity were identified (Fig. 4). The first component, interpreted as historical popularity, explained
53 30.8% of the variance and was primarily associated with years in use and classification as *classique* or
54 *patiné*. The second component, interpreted as current popularity, explained 29.6% of the variance
55 and was characterised by ascent numbers, recommendation percentages, star ratings from 8a.nu,
56 and guardian estimates. Together, these components explained 60.4% of the total variance. Variable
57 loadings are provided in Table A2.



58

59 **Figure A2.2: Biplot from Principal Component Analysis (PCA) showing two primary components explaining variation in**
60 **climbing intensity. The x-axis represents RC1 (historical popularity), while the y-axis represents RC2 (current popularity).**
61 **Arrows indicate the contribution of different variables to each principal component, and dots represent the analysed**
62 **climbing routes with colour depending on climbing intensity based on expert rank (green: unclimbed; light blue: low**
63 **intensity; dark blue: high intensity).**

64 Despite this explanatory power, the PCA approach appeared to underestimate the intensity of
65 frequently climbed low-grade routes and overestimate the intensity of higher-grade routes. For
66 example, some low-graded routes (e.g. 4c, 5c) that were ranked by experts as frequently climbed,
67 had a low PCA scores because they were not marked as *classiques* or *patiné* in the guidebook and had
68 relatively few ascent logs on 8a.nu. This could reflect a logging bias, as climbers may selectively log
69 more prestigious ascents, thus introducing a bias towards higher-graded climbs. Other similar
70 platforms to log ascents like ‘the Crag’ (www.thecrag.com) ([Rochers de Freyr, Sport climbing](#)) and
71 [27crag.com](#)) ([Freyr](#)) would suffer the same issue.

72 **Table A2: Loadings of climbing intensity indicators on the first two rotated principal component axes (RC1 and RC2) from**
73 **a Principal Component Analysis (PCA). Loadings indicate the contribution of each indicator to the rotated components,**
74 **with higher absolute values representing stronger influence on the corresponding axis.**

Variable	RC1: Historical Popularity	RC2: Current Popularity
Gardian estimates	0.347	0.347
Ascents (8a.nu)	0.119	0.517
Recommended (8a.nu)	-0.106	0.415
Stars (8a.nu)	0.315	0.356
Patiné	0.456	-0.143
Classiques	0.371	-0.419
(Un)climbed	0.396	0.280
Years in use	0.502	-0.253

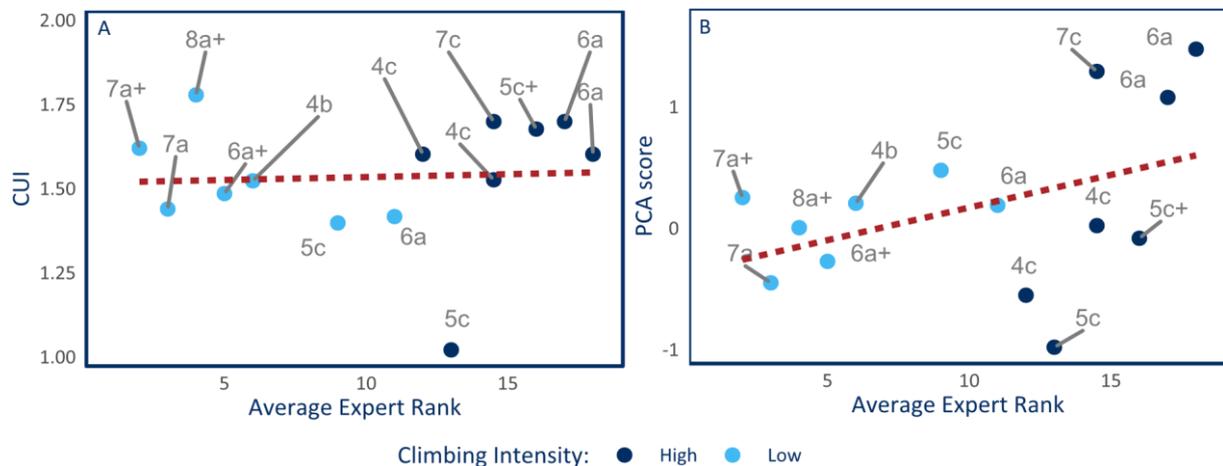
75

76 Expert-based approach

77 To validate these approaches, an expert-based ranking was obtained from eight experienced local
 78 climbers familiar with the routes in Freyr. Agreement among experts was assessed using Kendall's
 79 coefficient of concordance (W), a non-parametric statistic suitable for ordinal data. Values of W close
 80 to one indicate strong agreement, whereas values close to zero indicate weak agreement. An average
 81 rank was calculated across all expert rankings for each route.

82 To examine the relationship between expert rankings and both CUI values and PCA-derived scores,
 83 Kendall's rank correlation (τ) was used. This test is appropriate for comparing ordinal and continuous
 84 variables, particularly with small sample sizes ($n = 18$ for CUI; $n = 14$ for PCA).

85 Agreement among experts was moderate but statistically significant ($W = 0.510$, $p = 0.001$). However,
 86 correlations between expert rankings and proxy-based metrics were weak and non-significant. The
 87 correlation between expert rankings and PCA-derived current popularity scores was $\tau = 0.221$ ($p =$
 88 0.273 ; Fig. 5B), while the correlation with the CUI was $\tau = 0.156$ ($p = 0.441$; Fig. 5A). According to local
 89 experts, neither the CUI nor the PCA-based ranking accurately reflected actual climbing intensity at
 90 Freyr.



91

92 **Figure A2.3: Scatterplots showing the relationship between (A) Climbing Use Index (CUI); (B) PCA score and expert**
 93 **rankings for climbing routes. Each point represent a route, labelled whit its grade and is coloured according to its final**
 94 **climbing intensity category. Dashed correlation lines indicate the absence of a significant association between the two**
 95 **ranking methods. (n = 14)**

96 Final classification of climbing intensity
97 Given the weak correspondence between proxy-based metrics and expert judgement, climbing
98 intensity was ultimately categorised into three classes based on the average expert rankings:
99 unclimbed (n = 28), low climbing intensity (n = 116), and high climbing intensity (n = 104). This
100 categorisation allowed the inclusion of four additional routes lacking 8a.nu data, increasing the total
101 number of routes included in the analysis across the 36 subplots.
102

103 While a continuous measure of climbing intensity would retain more nuance, the categorical
104 approach was considered more robust given the limitations of the available proxies. Expert-based
105 classification also made it possible to incorporate local knowledge not captured by online platforms.
106 Although expert rankings are inherently subjective and may reflect shared perceptions within the
107 climbing community, the moderate level of agreement among experts and high agreement with the
108 final classification suggests that this approach provides a reasonable representation of relative
109 climbing intensity at Freyr. Categorisation into broader classes further helped to reduce the influence
110 of inconsistencies among individual expert estimates.

111 However, such expert-based rankings are site-specific and not directly transferable across regions or
112 climbing areas. For broader comparisons, indicators such as online ascent logs, guidebook sales, or
113 surveys about perceived crowding can be used. Future research should also consider more precise
114 monitoring, like targeted logbooks, observations or movement sensors, but these approaches have
115 to be implemented carefully to ethically handle privacy concerns.

116 A.3. Microclimate

117 Introduction

118 Cliff ecosystems are characterised by a vertical structure that generates distinctive and highly
119 heterogeneous microclimates. These fine-scale conditions can differ substantially from standardised
120 weather station measurements and are influenced by the aspect, slope, and solar exposure ([Bartlett
121 et al. 1990](#); [Bramer et al. 2018](#); [Larson et al. 2005](#)). Such fine-scale microclimatic variation will likely
122 play a key role in shaping cliff vegetation by filtering species according to their thermal and moisture
123 tolerances.

124 South-facing slopes in the Northern Hemisphere generally receive more solar radiation, making them
125 warmer and drier, while north-facing slopes are cooler and more humid ([Suggitt et al. 2011](#)). However,
126 cliff surfaces exhibit more complex radiation dynamics than horizontal or gently sloped surfaces.
127 Because of their steep angles, south-facing cliffs may receive less direct solar radiation during summer
128 in comparison to horizontal surfaces, when the sun is high in the sky, but more radiation during winter,
129 when the sun's angle is lower. In contrast, north-facing cliffs consistently receive less solar radiation
130 because they are oriented away from sun's path, resulting in persistently cooler microclimates
131 throughout the year ([Geiger et al. 2012](#); [Larson et al. 2005](#)). Radiation absorption and dissipation
132 primarily drive the energy balance at the cliff surface. With limited vegetation, evapotranspiration
133 often plays only a minor role, whereas albedo is a determining factor. Due to their low water content
134 compared to soils, rocks require less energy for heating, making them more susceptible to rapid
135 temperature fluctuations ([Larson et al. 2005](#)). The geomorphological heterogeneity of cliffs creates a
136 variety of microclimates, some of which can buffer extreme climate conditions, while others amplify
137 them. As a result, cliffs can serve as microrefugia for climate-sensitive species within a broader
138 unfavourable climate, while also potentially acting as stepping stones for warm-adapted species,
139 allowing them to extend their distribution ([De Frenne et al. 2013](#); [Dobrowski 2011](#); [Kuntz and Larson
140 2006](#); [Lembrechts and Nijs 2020](#); [Speziale and Ezcurra 2015](#)). Despite the ecological importance of cliff
141 microclimates in-situ microclimatic temperature measurements are rare. This study contributes to this
142 knowledge gap by comparing the temperature of north- and south-facing cliffs over the four seasons.

143 While the number of sensors was necessarily constrained by logistical and safety considerations, the
144 resulting data represent some of the few in situ measurements of cliff-face microclimates and offer an
145 important empirical baseline for future studies.

146 Methods

147 Microclimate data were collected by recording the temperature of air just in front of the rock (6 cm)
148 with 15 TOMST thermologgers ([Wild et al. 2019](#)) placed on different cliffs with variations in height and
149 aspect. Two sets of three sensors were placed at different heights on both a north-facing cliff
150 (Jeunesse) and a south-facing cliff (Le Mérinos). Two sensors were placed on an unclimbed cliff, which
151 was unmanaged and uncleaned, one on the north face and one on the south face. In total, six sensors
152 were placed on north faces, nine on south faces. Sensors were installed in spring (10 April 2024-24
153 June 2024) and read out in winter 2025 (20 January 2025- 21 February 2025) using TOMST's Lolly
154 software. Daily mean, minimum, and maximum temperatures were calculated for each thermologger
155 and used as the response variables in three separate linear mixed-effects models to assess differences
156 in temperature between north and south-facing cliffs across seasons. The model included season
157 (spring, summer, fall, winter) and aspect as fixed effects, along with their interaction. Aspect was
158 treated as a categorical fixed effect distinguishing between north- and south-facing cliffs, as the
159 aspects were clustered into high and low values for northness (i.e., the cosine of aspect), and variation
160 across the full compass circle was not available. Thermologger identity and date were included as
161 random effects to account for spatial and temporal variations. Afterwards, post hoc pairwise
162 comparisons were conducted using the emmeans package. Note that thermologgers are not well
163 suited for measuring temperature under direct solar radiation, due to heating of the sensor itself. We
164 thus recommend against interpreting the observed differences as absolute values ([Maclean et al.
165 2021](#)).

166 Results

167 Cliff aspect strongly influenced microclimatic temperatures, as expected. South-facing cliffs were
168 generally warmer than north-facing cliffs, but the magnitude of the difference varied seasonally. For
169 daily mean temperature, differences were most pronounced and highly significant in summer,
170 moderately significant in spring and fall, and small and non-significant in winter. Daily maximum
171 temperatures differed significantly across all seasons, and highly significant in all except winter. For
172 daily minimum temperatures, south-facing cliffs were significantly warmer than north-facing cliffs in
173 summer. See table A.3 for estimated marginal means, differences, p-values, and model R^2 .

174

175 **Table A3: Summary of linear mixed model (LMM) analysis of daily minimum, mean and maximum microclimate**
 176 **temperature. Estimated marginal means (emmeans) are shown for north- and south-facing plots across four seasons. The**
 177 **table includes the difference between aspects, corresponding p-values, and significance levels: < 0.1 (.), p < 0.05 (*), p <**
 178 **0.01 (**), p < 0.001 (***)).**

179 **Daily mean temperature:**

Season	North (°C)	South (°C)	Difference (°C)	p-value	Significance
Spring	13.08	16.79	-0.710	0.007	**
Summer	18.26	19.46	-1.197	< 0.001	***
Fall	9.34	9.97	-0.626	0.018	*
Winter	3.20	3.60	-0.398	0.133	

180 $R^2_{\text{marginal}} = 0.718, R^2_{\text{conditional}} = 0.994$

181 **Daily maximum temperature:**

Season	North (°C)	South (°C)	Difference (°C)	p-value	Significance
Spring	16.91	19.13	-2.23	< 0.001	***
Summer	22.71	26.03	-3.32	< 0.001	***
Fall	11.88	13.94	-2.05	< 0.001	***
Winter	5.57	7.03	-1.46	0.016	*

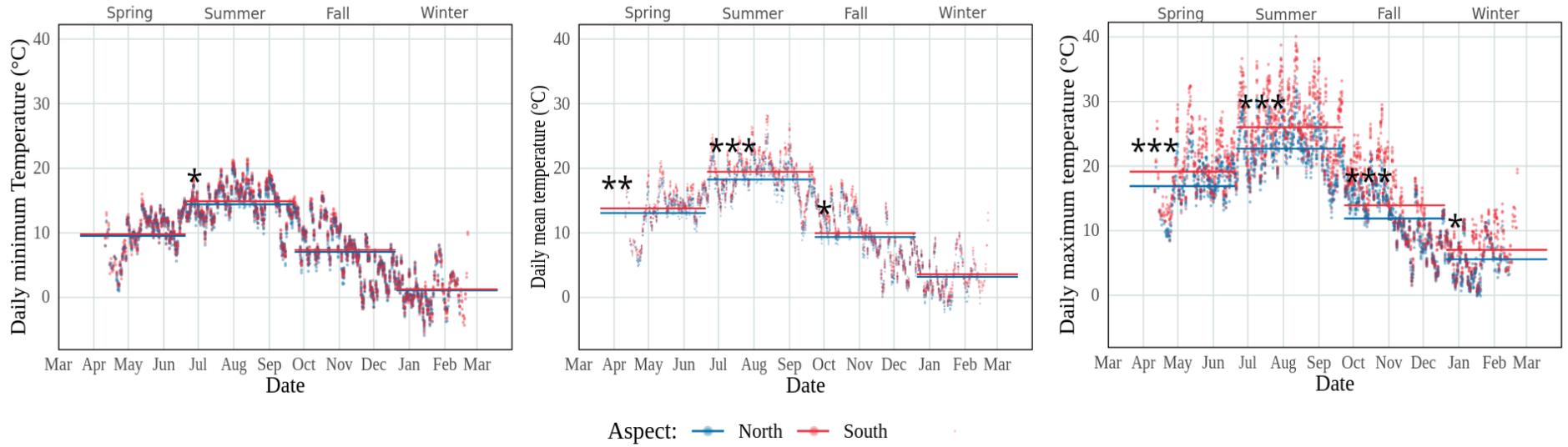
182 $R^2_{\text{marginal}} = 0.673, R^2_{\text{conditional}} = 0.73$

183 **Daily minimum temperature:**

Season	North (°C)	South (°C)	Difference (°C)	p-value	Significance
Spring	9.52	9.80	-0.279	0.2145	
Summer	14.42	14.91	-0.495	0.0274	*
Fall	7.05	7.36	-0.315	0.1609	
Winter	1.11	1.27	-0.160	0.1609	

184 $R^2_{\text{marginal}} = 0.658, R^2_{\text{conditional}} = 0.997$

185



186

187 **Figure A3: Daily minimum, mean, and max microclimate temperature over a time span of nearly one year (April 2024-February 2025) for north (blue) and south (red) faces. Dots represent**
188 **raw daily mean temperature; solid lines show estimated marginal means based on the linear mixed interaction model between aspect and season.**

189 Discussion

190 Microclimate measurements confirmed that south-facing cliffs are generally warmer than north-facing
191 cliffs, reflecting their greater exposure to direct solar radiation throughout the day ([Geiger et al. 2012](#)).
192 Temperature differences between aspects varied seasonally: in summer, long photoperiods and high
193 solar angles result in intense midday radiation on south-facing cliffs, while north-facing slopes receive
194 only oblique, low-angle sunlight during mornings and evenings. Notably, daily minimum temperatures
195 were still higher on south-facing cliffs in summer, indicating that cliffs retain heat overnight. Spring and
196 autumn showed intermediate differences, while winter differences were small and non-significant
197 ([Geiger et al. 2012](#); [Suggitt et al. 2011](#)).

198 However, the measurement of microclimate faced several limitations. First, only 15 thermologgers
199 were deployed for approximately one year, recording temperature alone. Second, as mentioned
200 above, TOMST thermologgers can overestimate temperatures under direct solar radiation, due to
201 radiative heating. Daily maximum values on exposed south-facing cliffs are affected most by it;
202 shielding reduces but does not eliminate this bias. Nevertheless, the general trends captured by the
203 microclimate model likely remain valid, as the processes causing these measurement errors - primarily
204 solar radiation - are also the main drivers of actual temperature differences. Still, specific quantitative
205 values may be unreliable. Future research could address this limitation by using ultra-fine wire
206 thermocouples, which provide more accurate measurements due to their small size, high reflectivity,
207 and thermally isolated data storage and housing ([Maclean et al. 2021](#)).

208 Third, due to the small sample size, the linear mixed model included only a simple interaction between
209 aspect and season. Future studies with expanded sensor networks could include additional variables
210 such as radiation, humidity, and wind. Such studies would also benefit from employing IoT systems for
211 efficient data collection in difficult-to-access cliff environments ([De Frenne et al. 2025](#)), although to
212 our knowledge no accurate, low-cost, robust IoT-sensor exists at this point. High-resolution UAV-based
213 thermal imagery and digital elevation models (DEMs) could further improve spatial modelling and
214 integration with topography and vegetation ([Bramer et al. 2018](#)). Comparative analyses across MIREN
215 Rocks sites worldwide would allow exploration of microclimatic patterns across latitudes, elevations,
216 and climates.

217 A.4. GLMM Species richness without high microtopography subplots (max. 6)
 218

Predictor	Species richness with all data points (n=248)	Species richness without high microtopography subplots above 6 (n=237)
<i>Intercept</i>	1.440 (p=0.007)	1.580 (p=0.007)
<i>Microtopography (linear)</i>	0.880 (p<0.001)	0.741 (p<0.001)
<i>Microtopography (quadratic)</i>	-0.276 (p<0.001)	-0.495 (p<0.001)
<i>Slope</i>	-0.458 (p=0.002)	-0.425 (p=0.004)
<i>Aspect (North/South)</i>	-0.730 (p=0.18)	-0.780 (p=0.19)
<i>Low climbing intensity</i>	-0.290 (p=0.30)	-0.256 (p=0.37)
<i>Unclimbed intensity</i>	0.372 (p=0.34)	0.392 (p=0.35)
<i>Microtopography : low climbing</i>	0.331 (p=0.04)	0.559 (p=0.005)
<i>Microtopography : unclimbed</i>	-0.573 (p=0.03)	-0.189 (p=0.52)
<i>R²_{Marginal}</i>	0.542	0.574
<i>R²_{Conditional}</i>	0.797	0.822

219
 220 Predictor effects were largely comparable between models. Both the direction and significance of the
 221 main effects remained unchanged. The only notable difference was the interaction between
 222 microtopography and unclimbed plots, which was no longer significant after excluding high
 223 microtopography subplots.

224 Explained variance was only slightly higher in the GLMM without the high microtopography subplots.

225 Overall, these results indicate that the small number of subplots with very high microtopography
 226 values did not drive or inflate the observed trends, despite their low representation in the dataset.

227 A.5. Simpson index

228 Simpson’s diversity index (D) incorporated species richness and relative abundance. The *vegan* package
 229 was also used, based on the following formula (Simpson, 1949): $D = 1 / \sum_{i=1}^S P_i^2$ where P_i is the
 230 proportional cover of species i in each subplot. Since Simpson’s diversity index is a proportion bounded
 231 between zero and one but includes a substantial number of zeros, a 16 zero-inflated beta regression
 232 generalised linear mixed model (GLMM) was fitted. The model used the same fixed-effects and full
 233 random-effects structure as described for the species richness model, except that it excluded the
 234 quadratic term for microtopography.

235 Simpson’s diversity index (D) incorporated species richness and relative abundance. The *vegan*
 236 package was also used, based on the following formula (Simpson, 1949):

237
$$D = \frac{1}{\sum_{i=1}^S P_i^2}$$

 238

239 where P_i is the proportional cover of species i in each subplot. Since Simpson’s diversity index is a
 240 proportion bounded between zero and one but includes a substantial number of zeros, a zero-
 241 inflated beta regression generalised linear mixed model (GLMM) was fitted. The model used the same
 242 fixed-effects and full random-effects structure as described for the species richness model, except
 243 that it excluded the quadratic term for microtopography.

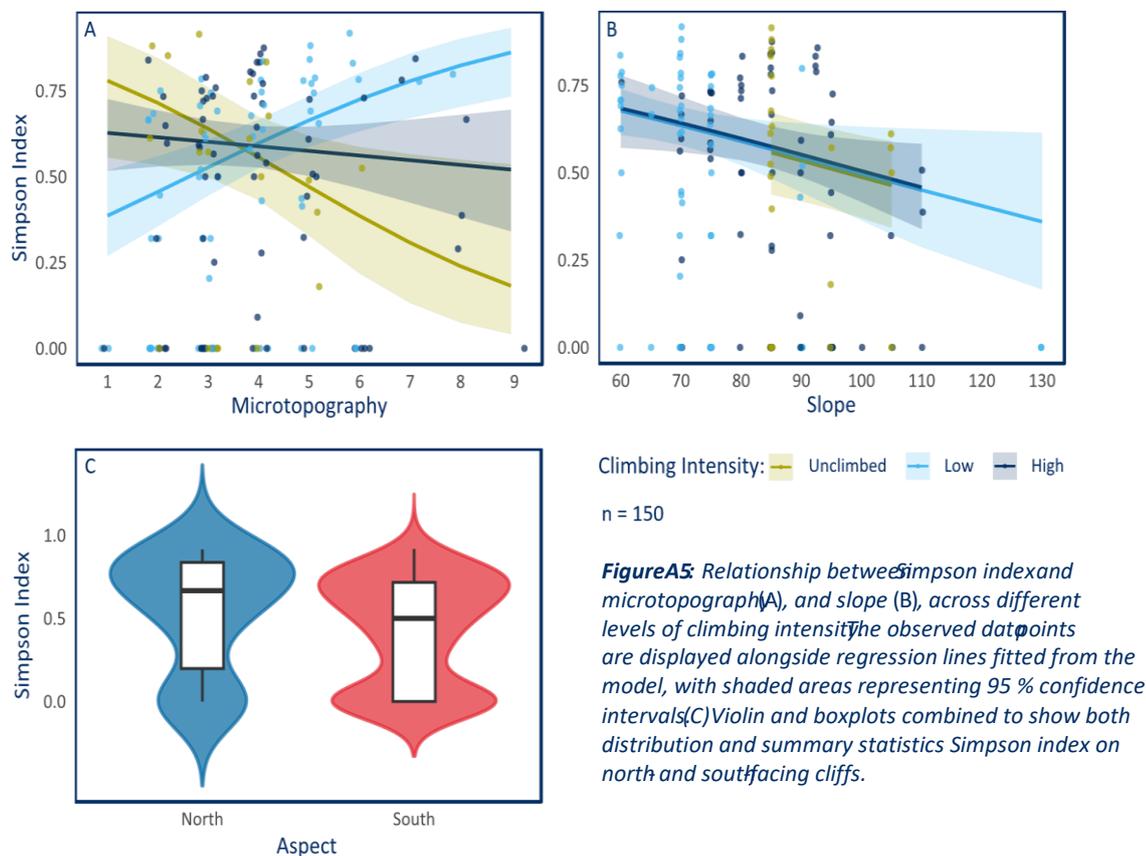
244 The generalised linear mixed model (GLMM), using a zero-inflated beta regression, revealed similar
 245 results as for Shannon index (section 4.2.3.) However, the model for Simpson index explained a lower
 246 proportion of variance ($R^2_{\text{marginal}} = 0.054$, $R^2_{\text{conditional}} = 0.078$). Additionally, a marginally non-significant
 247 negative interaction between microtopography and unclimbed subplots was observed, suggesting a
 248 negative trend between microtopography and diversity in the absence of climbing (tab. A7, fig. A7 A–
 249 C).

250 **Table A7: Summary of the fixed effects from the zero-inflated beta regression generalised linear mixed model (GLMM)**
 251 **testing the effects of climbing and abiotic factors on Simpson index per subplot on cliffs. Displayed are the estimated**
 252 **coefficients, p values, and significance levels for the : $p < 0.1$ (.), $p < 0.05$ (*), $p < 0.01$ (**), $p < 0.001$ (***)).**

Predictor	Estimate	p-value	Significance
Intercept	1.046	< 0.001	***
Microtopography	-0.093	0.425	
Slope	-0.261	0.032	*
Aspect (north/south)	-0.656	0.011	*
Low climbing intensity	-0.240	0.270	
Unclimbed intensity	0.108	0.715	
Microtopography × Low climbing	0.582	< 0.001	***
Microtopography × Unclimbed	-0.496	0.095	.

253

254



255

256 A.6. Results table indicator species analysis

257 **Table A6: Species significantly associated with each climbing-intensity level identified by indicator species**
 258 **analysis. Locally threatened (Red List) species are shown in bold. The indicator statistic reflects the strength of**
 259 **association between species occurrence and a given climbing-intensity level, and p-values indicate statistical**

260 *significance based on permutation tests. The KS value from Flora Indicativa represents the species' ecological*
 261 *strategy according to the CSR framework (competitor–stress tolerator–ruderal), indicating relative dominance*
 262 *along this gradient (Ivanova and Zolotova 2023).*

Group	Species	Indicator Statistic	P-value	KS (Flora indicativa)
Unclimbed	Umbilicus rupestris	0.316	0.001***	css
	Saxifraga tridactylites	0.305	0.001***	rrs
	Carex humilis	0.272	0.001***	css
	Erysimum cheiri	0.256	0.001***	rss
	Jacobaea vulgaris	0.250	0.002**	NA
	Scabiosa columbaria	0.172	0.04*	csr
	Unknown shrub	0.221	0.002**	NA
	Cardamine sp.	0.221	0.02*	NA
	Hieracium murorum	0.218	0.009**	crs
	Festuca pallens	0.205	0.009**	css
	Asplenium trichomanes	0.205	0.01*	rrs
	Sedum album	0.198	0.008**	sss
	Geranium robertianum	0.196	0.01*	crr
	Sesleria caerulea	0.190	0.03*	ccs
	Silene nutans	0.168	0.05*	ccs
	Centaurea scabiosa	0.153	0.05*	ccs
	Low	Seseli libanotis	0.194	0.02*
High	Arenaria serpyllifolia	0.170	0.03*	rrs
	Taraxacum sp.	0.154	0.04*	csr

263

264 A.7. Community-weighted means Grime plant strategies

265 Detailed results on how the occurrence of Grime plant strategies differs based on a linear model.

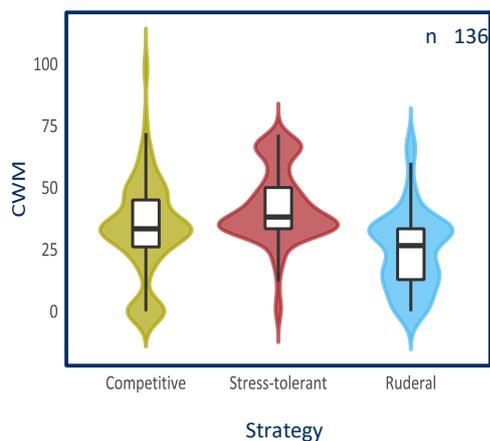


Figure Violin and boxplots combined to show both distribution and summary statistics of community weighted mean (CWM) values across the three plant strategies (competitive, stress tolerant, ruderal).

266

267

268

269

270 **Table A7: Summary of the linear model comparing CWM values among the three plant strategies, showing linear**
 271 **estimates, estimated marginal means (emmean), and p-values.**
 272

	Competitive (C)	Stress-tolerant (S)	Ruderal (R)
<i>Estimate</i>	30.5	38.9	22.7
<i>LM estimate difference</i>	30.464	8.405	-7.773
<i>Tukey-adjusted p-value</i>	< 0.001***	< 0.001***	< 0.001***

273

274 A.8. Climbing routes

275 **Table A8: Local climbing route names where vegetation was surveyed displayed with grade of climbing difficulty, amount**
 276 **of subplots (n) in each route, and category of climbing intensity based on the average expert evaluation.**

Route name	Grade	n subplots	climbing intensity
La Grune	4b	4	Low
L'Ascendante	4c	12	High
Arriba Bolivia	4c	12	High
Le Dernier Combat	5b	8	Low
Super-Vol-Au-Vent (L1)	5c	16	High
Les Anciens Belges	5c	20	Low
Le Rab Directe	5c+	20	High
The Wall	6a	12	High
Les Negresses Vertes	6a	12	High
L'avenue Chlorophyl	6a	12	High
La Lili	6a	12	Low
La Gamma	6a+	12	Low
La Tjetten	6b	12	Low
Mongolito	7a	8	Low
La cerise (L3)	7a+	8	Low
La Cerise départ Propagande	7b	4	Low
Troucage	7c	20	High
Cool Cat	8a+	16	Low

277

MIREN ROCKS

The MIREN rock survey



Pictures: Simon Dierickx

1 Overview of the MIREN rock survey protocol and contributions

The overall goal of the Mountain Invasion Research Network (MIREN) rock survey protocol is to study rock cliff vegetation around the world. We aim to (1) investigate the impact of rock climbing on these communities, (2) investigate the impact of rock climbing on the spread of alien plant species, (3) study the importance of rocks in species ranges edges, (4) compare the species composition of rock cliff vegetation with their surroundings and (5) assess the importance of the rock microclimates as microrefugia and on the distribution of rock vegetation. Data generated using the standardized protocol described in this document can be used to evaluate and quantify the processes and mechanisms shaping rock cliff communities at regional to global scales. We encourage the implementation of the protocol across the globe. We envisage revisiting the plots every five years to facilitate the investigation of long-term changes in rock vegetation. We primarily target bolted climbing routes (sport climbing, with fixed anchors), but we are open to trad(itional) climbing routes (without fixed anchors) as well. The protocol described below is developed for sport climbing. If you want to sample trad climbing routes, the protocol will be slightly different (see section 5.1). We explicitly exclude Bouldering and Via Ferrata (Or Klettersteig) routes.

If you plan to join MIREN Rocks, please let us know at miren.rocks@gmail.com!

2 Safety and permits

General safety statement:

As rock climbing inherently carries a significant level of risk, we want to stress the importance of following all rock climbing safety measures. Participation in the MIREN rock survey is therefore limited to individuals with **sufficient rock-climbing experience**. MIREN is not responsible for possible accidents during the MIREN rock survey.

Team composition:

Fieldwork must not be conducted alone. We strongly recommend a minimum of two people for sampling plots. At least one team member should be familiar with the local flora. Identifying species while hanging on the rocks can be quite time-consuming if you're not confident in your plant knowledge.

The second person is essential for both safety and efficiency. Responsibilities include:

- Conducting partner safety checks
- Belaying (if required by the chosen system)
- Monitoring safety while the climber is identifying plants or installing sensors
- Managing equipment
- Providing rapid assistance in case of an incident

Access and rope systems:

Always follow established safety standards. Ensure there is no risk of accidentally unlocking or misusing safety equipment during operations.

The most practical and controlled approach is to access rock cliffs from above, where possible.

We recommend using rope access techniques designed specifically for work at height. This includes:

- Using an additional safety rope, so fieldworkers are secured by two ropes all times
- Using semi-static ropes (never climb dynamically and fall on them)

- Using harnesses designed for height-work, which provide better comfort and stability during prolonged vertical tasks

However, all participants must use only rope access techniques in which they are properly trained, experienced, and fully competent.

Helmets must be worn at all times during fieldwork. **Rope protectors** should be used at all friction points.

Weather:

Safety in fieldwork does not rely solely on rope techniques. Rock-based fieldwork is physically demanding and often exposed to extreme environmental conditions.

- Bring ample drinking water.
- Monitor weather forecasts carefully.
- Select cooler, stable weather conditions whenever possible or prefer working during cooler parts of the day. Avoid working in extreme heat and unstable weather.
- Wear appropriate sun protection

These precautions help maintain fieldworker comfort, productivity, and reduce the risk of heat-related or weather-related incidents.

Collaboration with local climbing community and authorities:

We think it is also very important to reach out to the local climbing community and communicate openly about the goals of our research project. We encourage interested researchers to team up with local climbers since they often have invaluable information on the rock site, safety measures and the legal framework.

Lastly, be sure to contact local authorities or the organizations managing the rock climbing site to obtain the necessary permits for your fieldwork! If you install temperature sensors, make sure to communicate this to the local climbing community. You can find an example of affiche about the sensor as an appendix of the protocol.

3 Protocol methodology

Within each **Site**, sampling occurs along **transects**. Whenever feasible, we strongly encourage sampling **paired transects** (one climbed and one unclimbed). However, we acknowledge that unclimbed transects are often difficult or impossible to find. In such cases, working with unpaired transects (e.g., only climbed transects, or fewer unclimbed than climbed transects) is acceptable.

Along each transect, plant species composition will be recorded in several **Plots** of 2 by 2 m², composed out of 4 subplots of 1 m². Sites are potentially subdivided into different **Sectors** (e.g. a north facing and a south facing sector within one site). Further description of Sites, sectors, Transects and Plots can be found below. We have set the minimum amount of plots to be able to join MIREN rocks at 30 (=120 & m² subplots).

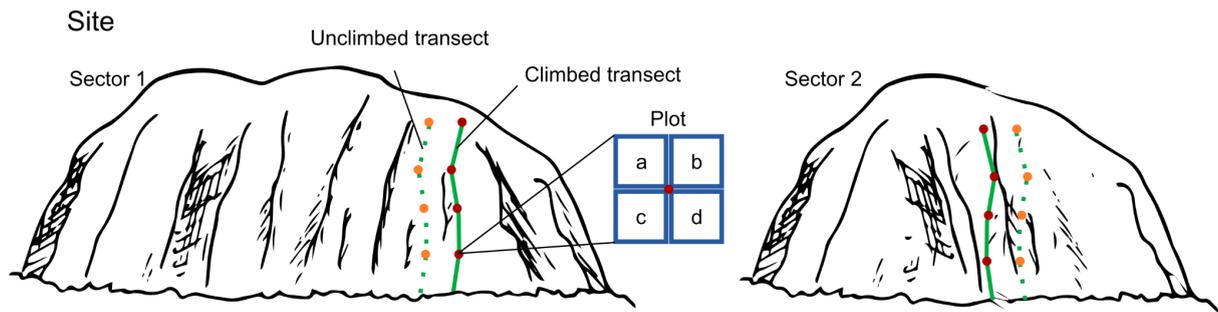


Figure 1: A rock climbing site with 2 paired transects (in total 2 climbed and 2 unclimbed transects). The vegetation is recorded in a plot of 2 m x 2 m, consisting of 4 subplots of 1 m² (a, b, c, d)

3.1 Site

In this survey, a Site is defined as a rock cliff which is used and equipped for rock climbing. The protocol is specifically designed for climbing sites with routes having fixed anchors (but see section 5), which are used to locate the plots and mark them for resurveys. It could be that a Site consists of different climbing Sectors, which are often physically isolated from each other. These sectors often get distinct names. If this is the case, the hierarchical protocol gets an additional fifth layer: Sites>Sectors>Transects>Plots>Subplots.

The delineation of a Site will depend on many local factors and we do not seek to constrain these. Ideally, a new Site should not be in, or part of, an existing 'MIREN rocks' site (see <https://www.mountaininvasions.org/miren-rocks> for an up-to-date map). Please contact us (miren.rocks@gmail.com) before starting the survey to check whether your region is suitable for inclusion in the MIREN rocks database.

3.2 Transects

Paired transects

As much as possible, we will sample pairs of transects, consisting of a climbed and an unclimbed transect. A climbed transect is a climbing route, which is equipped with fixed anchors for climbing (e.g. bolts, pitons). These climbing routes should have a minimum length of 15 m. We did not set a maximum length. Unclimbed transects have no evidence of climbing (no chalk or permanent equipment), are not mentioned in local guidebooks and occur at least 2 m from established climbing routes (Harrison et al, 2022, see Figure 2 - Option 1). Given our 2 m by 2 m plots, centered around the bolted anchor, the plots of the climbed and unclimbed transect will touch at the sides. Please consider a wider spacing between climbed and unclimbed transects if you notice that the impact of climbing goes beyond 1 meter at either side of the bolted anchor. It is important that paired climbed and unclimbed transects have similar environmental conditions (e.g. slope, aspect, microenvironment) and only differ in terms of climbing use.

Unpaired transects

In climbing sites where a dense network of routes does not allow the placement of unclimbed transects directly next to the climbed transects as described above, the unclimbed transects could also be placed in a separate, unclimbed part of the rock site (Figure 2 - Option 2). Since bolting is missing here (or climbing is not allowed), we advise reaching the transect via the top of the rock (rappelling down). Also in option 2, paired climbed and unclimbed transects should have similar

environmental conditions, if possible. Consider contrasting highly popular with rarely used transects to get contrasting climbing intensity.

For both in paired and unpaired transects, we are interested in climbing intensity. Empirical count data on climbing frequency are often difficult to obtain. In a previous case study, we explored several approaches, including online ascent logs (e.g., 8a.nu or TheCrag), the Climbing Use Index (Clark & Hessel, 2015), and information from local guidebooks, as well as **surveys of local climbers**.

We ask you to conduct a similar survey at your study site. This involves listing all routes where inventories were conducted and sending this list to at least 10 climbers who are very familiar with the area and its routes. Ask them to rank the routes from least frequently climbed to most frequently climbed, based on their own experience, perception, and observations. An example of such a survey is provided in the appendix 6.7 of this protocol.

There is no minimum amount of transects to survey to be included in the database, since transect length will differ considerably among regions. However, we estimate that at least 30 plots (15 climbed and 15 unclimbed if using paired transects) would be needed to have sufficient regional explanatory power. Note that 30 plots are 120 1 m² subplots. More might be necessary if the diversity and heterogeneity in microenvironments and plant communities are high, or if rock faces are very smooth and vegetation presence is low in general. In that case, consider monitoring across a range of climbing grades, as vegetation is usually more present on easier climbs. Of course, we want to stimulate you to sample as many plots as possible.

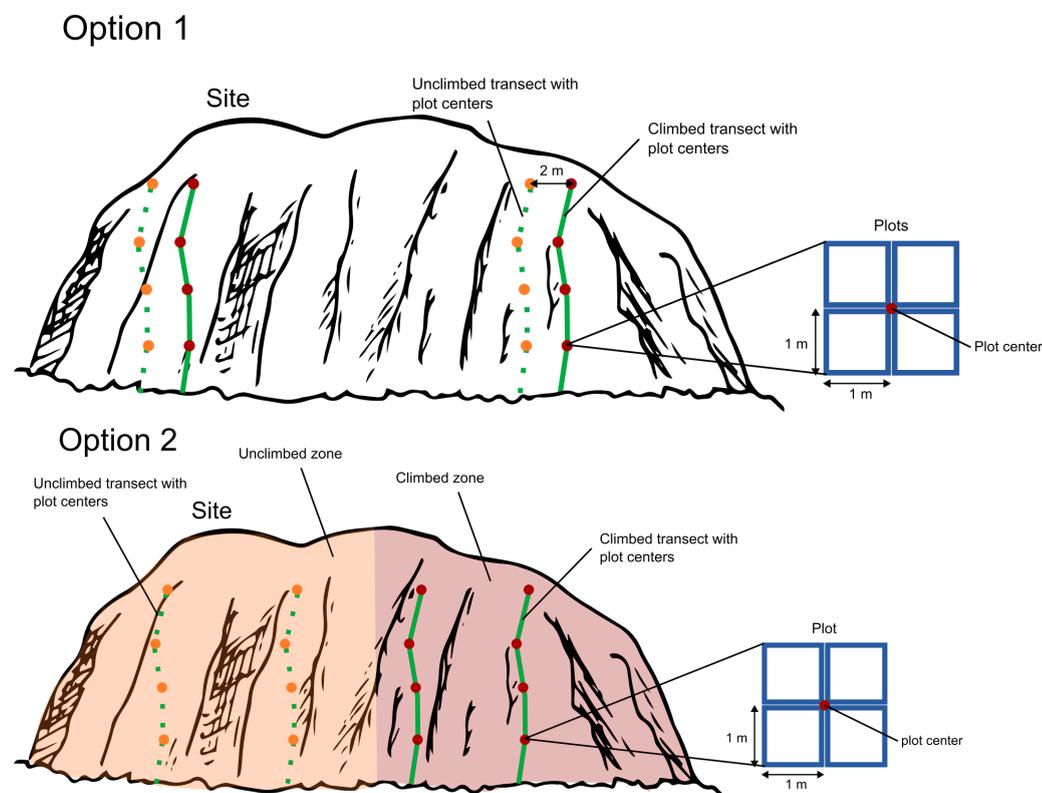


Figure 2: two options to position paired climbed and unclimbed transects within a rock site

3.3 Plots

Along each transect, we will sample the vegetation and environmental conditions in squared plots of 2 m by 2 m, consisting of 4 subplots of 1 m². For the climbed transect, the squared plots will be centered around the fixed anchors (e.g. fixed bolts and pitons). These fixed anchors serve as permanent markings and will allow resurveys in the future. The minimum separation between the plot centers of two consecutive plots should be 3 m. You should aim to have at least 3 plots per transect.

For option 1 (Figure 2), unclimbed transects will be chosen at either side of a climbed transect. The plot centers of the unclimbed transect will be positioned at the same height as the plot centers of the climbed transect and at a minimum distance of 2 m from the plot centers of the climbed transect. At this minimum distance of 2 m, the subplots of the climbed and unclimbed transect touch each other at the sides. Paired plots must have similar conditions (heterogeneity, slope, aspect). For option 2 (Figure 2), try to position the unclimbed plots at the same height as the climbed plots of the corresponding transect. Also here, pay special attention to having similar environmental conditions in paired plots.

We will work with a fixed grid of 1 m by 1 m (e.g. consisting of plastic tubes, see Appendix 6.1) to position the subplots around a plot center. We advise using chalk to draw each of the subplots consecutively on the rock, after which you can start collecting data in each of the subplots (see section 4). Please remove the chalk after the survey.

4 Data collection

The following descriptions are based on the criteria to include a dataset in the global MIREN Rocks database. All data sheets must include the date of the survey and the names and emails of recorders to facilitate data checking. If you have any questions regarding data collection, contact miren.rocks@gmail.com.

All data should be submitted to miren.rocks@gmail.com using [this template](#). For the northern hemisphere, the deadline to submit the data is October 30th 2026. For the surveys in the southern hemisphere, data should be submitted before April 30th 2026. Please contact us if you want to join, but the initiated deadlines are too soon.

Appendix 6.9 shows an A5 summary sheet of the data that must be collected in the field. Print it out and take it with you during fieldwork. You can also use it as a template for your notes.

4.1 Site

- Country/location
- Site name: Official name of climbing site
- Type of cliff: Natural cliff, quarry, ...
- Grade system of routes: Yosemite Decimal System, French Grading Scale, UIAA, Australian scale, UK system - numerical, UK system - adjective
- Is the site managed/cleaned for climbing purposes by local climbing organizations?: Yes/No
 - How is it cleaned?

- How often is it cleaned? Multiple times per year/ once per year / once every few years
- When was the last cleaning (if known?)
- If the Site has different sectors, also record the sector names
- Estimate of climbing intensity of site:
 - Numbers of climbers per year (in classes: 1: 0-100, 2: 100-500, 3: 500-1000, 4: 1000-2000, 5: >2000. Sources: climbing books, local climbing association or websites like thecrag.com or mountainproject.com)
 - Year of opening of the site
 - Climbing season length (months)
- Microclimate temperature (optional, see section 4.4)
- For communication purposes, we welcome pictures of the surveyed sites and of you when climbing and conducting the research protocol

4.2 Transect

- Geolocation: Coordinates should be recorded at the base of each transect and reported as latitude and longitude in decimal degrees using the WGS 84 datum; please check that the mapped locations are correct using GIS or Google Earth before submitting data.
- Elevation at the base of the transect: in meters above sea level; obtained using a digital elevation model or Google Earth; at the same locations as geolocation is recorded.
- Overall aspect/orientation (°) using a compass
- Overall slope (°) - overhanging transect (>90°), slab transects (<90°) using an inclinometer
- Type of rock substrate: we will follow the classification system of thecrag.com (see Appendix 6.2)
- Total length of the transect (m)
- Date of survey: dd/mm/yyyy
- For climbed transects:
 - Local name of the route
 - Year the route opened
 - Multipitch: Yes/No
 - Grade of route
 - Popularity of the individual climb (0-5; relative within the site) inferred by the number of stars or popularity (source: climbing books, local climbing association or websites like thecrag.com or mountainproject.com)
 - Time to approach the transect (in minutes)
- For unclimbed transects
 - Position compared to the paired climbed transect: Left/Right
 - Distance to paired climbed transect (in m)

4.3 Plot

The following is to be recorded in each 2 m x 2 m plot:

- Plot code: Unique identifier for each plot, recorded in the form **Country.Site.Transect.PlotSubplot** (e.g. BE.FRE.C6.2a).
 - Country is the official two-letter country code (e.g. Belgium = BE).

- Site should be three letters (e.g. FRE = Freyr). This code will be given to you by MIREN. If there are different sectors in one site, please number them (eg. FRE1, FRE2, ...)
- Transect consists of a letter (C1-C... for climbed routes, U1-U... for unclimbed routes) and a number. Transects are numbered from 1 onwards. Pairs of transects get the same transect number (e.g. C6 and U6 = 6th climbed and unclimbed transect).
- Plots are numbered from 1 onwards, with 1 being the lowest plot. Paired plots should get the same number.
- Subplots are labelled a (top left), b (top right), c (bottom left), d (bottom right) (see Figure 1)
- Name and cover of each species: see section 3.3.1 for species nomenclature information and see section 3.3.2 for the ordinal projective foliage cover estimates to be used.
- Height of the plot center above the base of the route (m)
- Anchor number (measured from bottom)
- Aspect (°) - measured using a compass
- Slope (°) - measured at plot centre using (in)clinometer
- At the subplot level: surface heterogeneity in classes (see Appendix 6.3)
- At the subplot level: total vegetation cover (by vascular plants, in %)
- Take a picture of every subplot

4.3.1 Species information

Vascular plants

All angiosperms, gymnosperms, and ferns, both native and non-native, that create foliar cover in a subplot are to be recorded.

- If a species can only be identified to the family or genus level, assign a region-specific name. This name should combine the taxonomic level, country code, site code, and a unique identifier (e.g., *Acacia BE.FRE1* or *Asteraceae BE.FRE2*).
- If a species cannot be identified even to the family level, use a name consisting of its life form (e.g., graminoid, herb, fern, vine, shrub, or tree), the region code, and a unique identifier (e.g., *shrub BE.FRE3*).

Bryophytes

Bryophytes (mosses, liverworts, and hornworts) present in each subplot will also be recorded. Since field identification of these groups is challenging, we will employ the following dual approach:

1. **Morphotype classification**
 - Bryophytes will be classified into morphotypes instead of species, following the guidelines in Appendix 6.4.
2. **Bryophyte harvesting**
 - In each subplot, carefully remove a small, representative fragment of each moss cushion using a knife or similar tool, minimizing damage to the remaining

cushion. Place the collected samples in a pre-labeled paper bag (with the subplot code).

- Dry the samples at room temperature for three days, then send them for identification to:

Alain Vanderpoorten

**BAT. B22 Biologie de l'évolution et de la
conservation-aCREA-Ulg**

Quartier Vallée 1

chemin de la Vallée 4

4000 Liège 1

Belgium

- Please notify us by email once the moss samples have been shipped.

Lichens

Lichens will also be classified into **morphotypes** instead of species, following the guidelines in Appendix 6.4. **Harvesting lichens for the MIREN Rocks protocol is not needed.**

If in-house expertise is available for identifying bryophytes and/or lichens to the species level, such data is encouraged in addition to morphotype classifications. These observations should be recorded in the same form and using the same methods as those for vascular plants.

4.3.2 Species cover estimates

Estimate total plant cover (in %) of each plant species. Plant cover is defined as the proportion of the rock that is shaded by vegetation foliage when projecting it perpendicularly on the rock surface. This value needs to include cover from all locations of each species within the plot boundaries. Note that the sum of all species-specific cover estimates can exceed 100 % due to overlap in coverage. In order to help the estimation of the plant cover, you can find physical representations of 1, 2 and 5% as a reference in Appendix 6.5. We urge you to take these examples with you in the field.

Estimates are at the 1% resolution, yet for small plants, covers of 0.1 (~3 x 3 cm) and 0.5% (~7 x 7 cm) could be used. It is important to calibrate cover assessment among a survey team, and an accuracy assessment should be conducted (see Appendix 6.6). Surveying should only begin once accuracy standards have been met.

4.4 Microclimate measurements

The measurements of the microclimate temperature are optional. The goal is to measure microclimate temperature at the rock site level (or sector level) and compare it to the microclimate temperature of an open (non-shaded), flat habitat near the rock site. More specifically, we ask you to install (1) one temperature logger at the rock face of every site or at every sector within a rock site. Make sure to cover the different rock cliffs facing different directions, since we expect the microclimate to differ strongly between different directions, and (2) one temperature logger in an open (non-shaded), flat habitat near every rock site which serves as a control. The data should be delivered in the SoilTemp data format: <https://www.soiltempproject.com/how-to-submit-data/>.

Here, we describe the specifics of the temperature measurements:

- Temporal specifications of the measurements:

- We ask you to measure temperature during the growing season for at least 3 months
- Measure at a minimal temporal resolution of 1 measurement every 30 minutes.
- Installation location:
 - The logger in the rocks should also be installed above the trees and away from shrubs to avoid their shade. Being exposed to the sun, it is absolutely necessary to shield the sensors from the sun!
 - The logger in the open habitat should be installed just above the soil surface (e.g. between 5 and 15 cm above the surface) at a non-shaded location, far from trees or other shade-casting objects. Being exposed to the sun, it is absolutely necessary to shield the sensors from the sun! If there is a lot of wildlife in the region, it could be necessary to protect the logger with a wire cage (e.g. a [shopping basket](#)).
- Recommended loggers:
 - We recommend using the [TOMST Thermologger](#) on the rock face, because it is very accurate and it comes with a handy screw to attach it to the rock (Figure 3). Best use the included screw and a plug (not included) to attach it to the rock. Please make sure to order the triple radiation shield from TOMST to shield this logger from the sun (per default it comes with a single shield). This logger can also be used to measure the temperature in the open, flat habitat. [Maclean et al. \(2021\)](#) have shown that a single radiation shield is not enough.
 Note: installing the microclimate sensors on the rock face can be challenging. Some cliffs have cracks that you can utilize, while others don't. Some participants have used drills, but that can add extra complexity to the process.
 - For the open flat habitat, you can also use the Thermologger, or you can use the (more expensive) [TOMST TMS-4](#), which is specifically designed to be installed in the soil. Please make sure to order the triple radiation shield from TOMST to shield this logger from the sun (per default it comes with a single shield). If these TOMST sensors are not available in your country, we recommend using Lascar or Hobo loggers (or another brand with similar accuracy), installed with a Gill-style radiation shield (which can be self-made, see Figure 4).



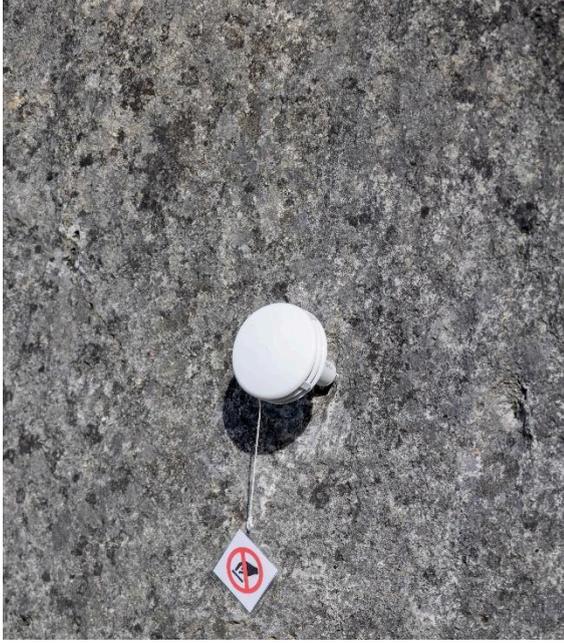


Figure 3: TOMST Thermologger can be used to measure the temperature at the rockface, as it comes with a handy screw and triple radiation shield. Pictures from <https://tomst.com/>, Jonas Lembrechts and Kobe Burdack



Figure 4: Lascar logger with a self-made double radiation shield, made from two white funnels. The inner funnel has holes to stimulate air flow. Pictures from Pieter De Frenne

4.5 Surrounding vegetation

To answer research questions 3, 4 and 5, we would need information on plant species occurring in open habitats in the region surrounding the surveyed rock sites. Therefore, we ask you to provide a list of plant species that occur in open natural habitats in the region surrounding the surveyed rock sites. This list can be expert-based or extracted from databases, like GBIF. If you have the time or just happen to have data, community data (i.e. plot-level data) are welcome as well (but not mandatory).

5 Deviations from standard protocol

5.1 Traditional climbing

The protocol above is developed for sport climbing routes with fixed anchors. If you want to join with a traditional climbing site, there are a few things to consider:

- If you aim to compare climbed and unclimbed transects (but see 5.2), make sure the climbed transect is being climbed (e.g. existing route in a climbing book, signs of chalk). The same for unclimbed transects: make sure it is not climbed.
- Since it will be hard to have permanent markings to facilitate resurveys, we don't consider these sites for future resurveys
- Data on anchor numbers are not needed
- Be extra careful!!

6. Appendices

6.1 Fixed frame

We advise using a frame consisting of two plastic tubes of 1 m, connected with 1 m ropes (Figure 5). Best to hang the frame to a second rope with a prusik knot for easy vertical movement along the wall. A spirit level attached to one of the tubes allows to level the frame horizontally. Adding coloured tape could aid the estimation of the species cover (Figure 6).



Figure 5: Fixed frame connected with a prusik knot to a second rope

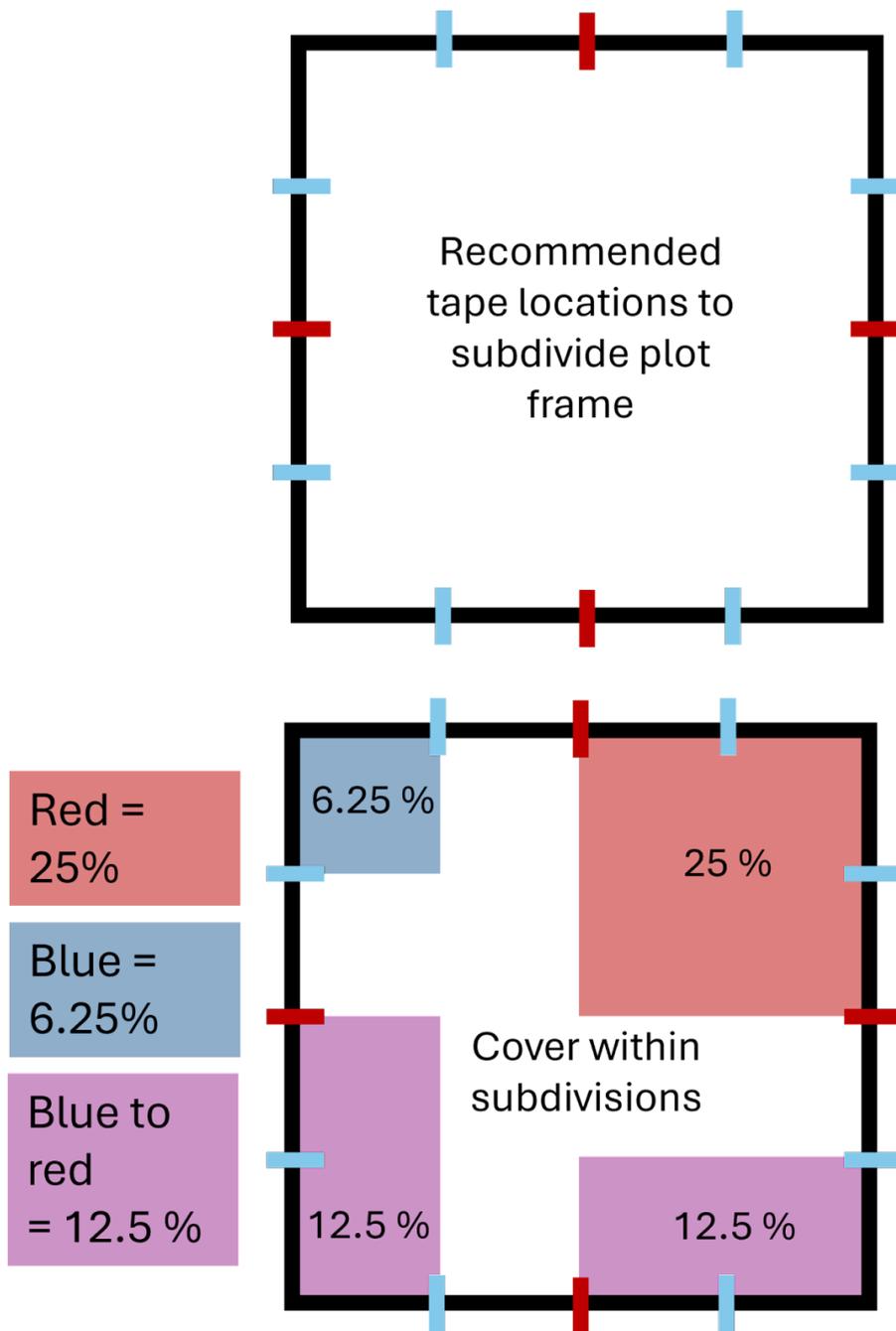


Figure 6: Recommendations for plot subdivisions to assess cover using colored guides (tape). Figure from Georgia Harrison & Laura Boggess

6.2 Geological substrates

We will follow the classification system of thecrag.com, which can be downloaded [here](#). For more information, you can visit thecrag.com

6.3 Surface heterogeneity classification (Protocol from Georgia Harrison & Laura Boggess)

Surface heterogeneity describes the abundance and size of surface features on cliff faces such as ledges, pockets, and cracks. These features can capture soil, water, and propagules so quantifying their abundance is key to understanding cliff-face vegetation. In addition, overhanging features can impact the vegetative community below, and their presence should be considered.

Assign a surface heterogeneity value between 0-10 for each plot. 0 is smooth rock, 10 is highly fissured. Score only the rock, so if vegetation is present, try to picture the plot without any vegetation or soil, and score only the rock. See photos (Figure 7) below for proposed examples.

Only quantify features **within the plot**. Exclude features that are not inside the plot boundary. Note if a plot is below an overhang. Overhangs should be at least 0.5 m deep.

Table 1. Example Table of plot-level abiotic information to be collected:

Subplot ID	Surface heterogeneity (0-10)
BE.FRE.C6.2a	2
BE.FRE.C6.2b	4

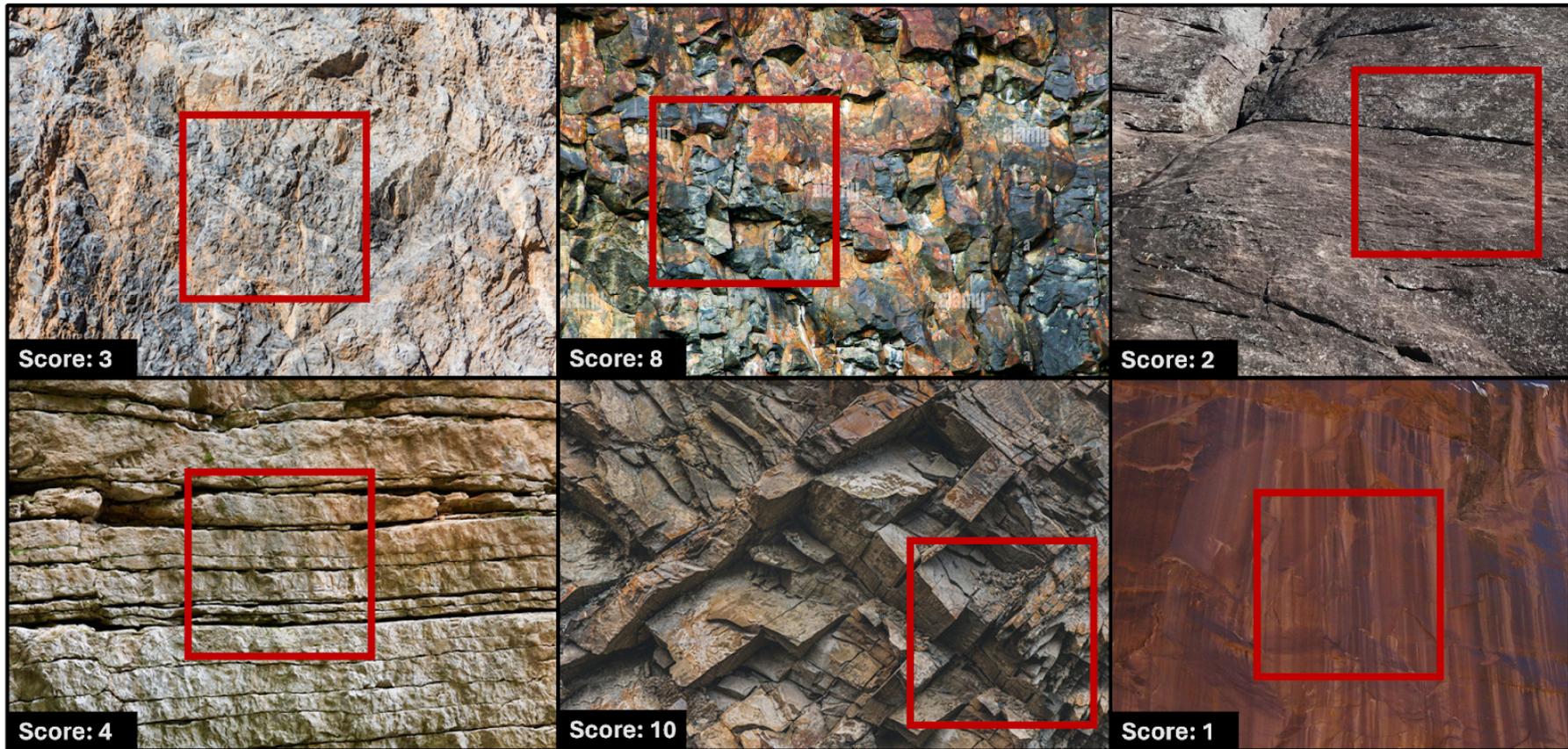


Figure 7: Example cliff face photos to score surface heterogeneity. Red box is plot frame, only score surface heterogeneity within plot frames. Not included in photos is the distinction between overhanging or non-overhanging features.

**Note: these photos are just found from the internet, but it may be useful to create a guide with example photos and their scores with photos from a specific site.*

6.4 Bryophyte and lichens classification (Protocol from Georgia Harrison & Laura Boggess)

Lichens and bryophytes (mosses, liverworts, and hornworts) are the most abundant and diverse taxa on cliffs. While identification can be challenging, collecting at least some data for these groups is essential to understanding ecological processes on cliffs. When species identification is not possible, recording cover for lichen and bryophyte morphotypes can still provide valuable information. Here, we use this morphotype identification in addition to species-level identification.

6.4.1 Lichens

Lichens are symbiotic organisms composed of fungi and algae or cyanobacteria. They are phenotypically plastic and the same species can look different under different environmental conditions, making them difficult to identify in the field. To meet this challenge, we recommend assigning a unique morphotype to each unidentified species of lichen.

Lichens are generally categorized into three main growth forms (Figure 8):

1. Crustose: These lichens form a crust adhering tightly to the substrate. They cannot be removed without damaging the substrate. Crusts are the most common morphotype in cliff plots.
 - a. *optional*: Powder lichens are a subtype of crusts whose bodies (thalli) are covered in powdery lichenized propagules and tend to occupy ecologically different niches from non-powder crustose lichens. To distinguish between powder or non-powder lichens, rub a finger on top of the thallus and if dust or powder comes off, this organism can be classified as a powder lichen.
2. Foliose: These are leaf-like lichens that are attached at some points, allowing some parts to lift away from the substrate. They generally have a flat, leafy appearance and a distinct upper and lower surface.
3. Fruticose: These lichens are bushy or hair-like and grow away from the substrate in all directions. They are often cylindrical or branched and do not have a distinct upper and lower surface.

When collecting lichen data, describe the morphotype by determining growth form, then noting the color and texture (Table 2). Overall, classifying into lichen morphotypes is still likely underestimating overall diversity. As a result, if you're not sure if two lichens are the same or different species, split the species and give it a different descriptor for morphotype so that you could recognize it in future plots. It is better to have more, distinct groups which are recognizable in future plots than fewer, coarse groups.

Table 2. Example Cover table for functional groups. For Taxon Group, L = Lichen, B = Bryophyte

Subplot ID	Taxon Group	Growth form	Color	Texture/additional notes (optional)	Cover (%)
BE.FRE.C6.2a	L	Crustose	White	bubbles	10
BE.FRE.C6.2b	B	Acrocarp moss	Light green	feathery	15



Figure 8: Three main lichen morphotypes. Powder lichens (not pictured) are a sub-type of the crustose group.

6.4.2 Bryophytes

Bryophytes are non-vascular plants, including mosses, liverworts, and hornworts. They are typically found in moist, shaded environments. First, determine the major group (Figures 9 and 10).

1. Mosses: usually have clearly differentiated stems and leaves. They grow in dense green clumps or mats.
2. Liverworts: can be leafy like mosses, but often have a flatter, more ribbon-like appearance with less obvious differentiation between stems and leaves. They can also be thalloid, which are flat and spread out like a thin crust.
3. Hornworts: These have a thalloid body similar to some liverworts but differ mainly in their reproductive structures which are elongated and horn-like.

Mosses can be further delineated into two main groups: "acrocarp" and "pleurocarp" based on growth patterns and reproductive structures (Figure 9). Acrocarp vs Pleurocarp:

1. Growth Pattern: Acrocarpous mosses grow upright and tend to form dense clumps, while pleurocarpous mosses spread out, forming mats.
2. Reproductive Placement: Acrocarpous mosses have reproductive structures at the top of vertical stems; pleurocarpous mosses bear theirs along the sides of branches.
3. Environmental Preference: Acrocarpous mosses are more adaptive to varying moisture conditions and can survive drying, whereas pleurocarpous mosses thrive in consistently wet habitats.



Figure 9: Major groups of mosses: Top row: acrocarpous moss with reproductive structures on the tip of the stalk. Bottom row: pleurocarpous moss with the reproductive structure emerging from a side branch



Figure 10: Left: Hornwort with typical horn-like reproductive structures. Right: Liverwort, here with umbrella shaped reproductive structures

6.5 Physical representations of 1, 2 and 4% cover as a reference (print on A4 format!)

1%

This square should be

10 cm x 10 cm

2%

This square should be

14.14 cm x 14.14 cm

4%

This square should be 25 cm x 16 cm

6.6 Setting accuracy standards (Protocol from Georgia Harrison & Laura Boggess)

To ensure data collection with high accuracy, conduct a calibration test and accuracy assessment for surface heterogeneity and cover (either virtually or in-person) at each site. The team leader should explain the protocol and provide a few examples (i.e. show a picture of a plot with a surface heterogeneity score of 0, 5, and 10) and justification for the score. Then, to ensure that the sampling group will be consistent, each member of the group should conduct their own assessment for standard plots. Repeat until the group is consistently scoring within accuracy standard.

We recommend using a 10% accuracy standard, and aiming for at least three plots in a row that meet the accuracy standard. Virtually, this accuracy assessment could also be conducted with a form or survey, but it is best if surveyors receive instant feedback (i.e. I scored this plot a 5, but the team agrees it is a 3).

Table 3. Example accuracy training table for surface heterogeneity. Set a standard for accuracy (i.e. within 1 surface heterogeneity unit, or within 10% cover).

Plot	Surface heterogeneity (personal)				Meeting accuracy standards?
	Team member 1	Team member 2	Team member 3	Team member 4	
A	8	2	5	6	No
B	6	4	7	5	No
C	9	8	8	9	Yes

6.7 Estimating climbing intensity with a survey

To estimate climbing intensity, prepare an Excel file (or similar spreadsheet format) and send it to around 10 local climbers who are very familiar with the site and its routes.

List all climbing routes where you conducted vegetation inventories. Ask the participants to rank the routes according to climbing intensity, from least frequently climbed to most frequently climbed, based on their own experience, observations, and perception. They do not need to have personally climbed all routes to provide a ranking.

Each respondent should provide their ranking in a separate column next to the column listing the routes.

Please submit the completed spreadsheet together with your data submission.

Below, you will find a screenshot of a similar survey used in a previous case study.

	A	B	C	D	E	F	G	H	I	J	K	L	M	N	O
1	Route Name	Order (1 Lowest, 18 Highest)													
2	Armba Bolivia														
3	Super Vol-au-vent L1														
4	La Cerise Depart Propaganda														
5	La Cerise L3														
6	La Gamma														
7	La Grune														
8	La Lili														
9	La Tjetten														
10	L' Ascendente														
11	L'Avenue Chlorophyl														
12	Le Dernier Combat														
13	Le Rab Directe														
14	Les Anciens Belges														
15	Mongolito														
16	Negresses Vertes														
17	The Wall														
18	Troucage														
19	Cool Cat														
20															

Rank these routes from least climbed (1) to most climbed (18) according to your own perception of current climbing intensity.
 Write the numbers 1 to 18 to the corresponding routes in column Order.

6.8 Example communication thermologgers

SENSOR ON THE WALL



Chers grimpeurs,
Des capteurs ont été placés sur ce rocher à des fins de recherche scientifique sur la végétation rocheuse. Ils sont placés de manière à ne pas gêner l'escalade. Ne les touchez pas et laissez les capteurs sur place. Nous vous remercions de votre coopération.

Beste klimmers,
Op deze rots zijn sensoren geplaatst voor wetenschappelijk onderzoek over rotsvegetatie. Ze zijn zo gepositioneerd dat ze het klimmen niet verstoren. Gelieve de sensoren niet aan te raken en te laten hangen. Dank voor jullie medewerking.

Dear climbers,
Sensors have been placed on this rock for scientific research on rock vegetation. They are positioned so as not to disturb climbing. Please do not touch the sensors and leave them in place. Thank you for your cooperation.

Contact: miren.rocks@gmail.com



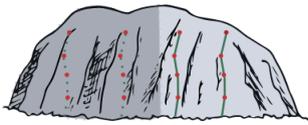
6.9 Summary sheet field data collection



Date of the survey

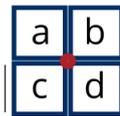
MIREN Rocks survey summary

Transect



- Length
- Coordinates
- Elevation
- Overall slope and aspect
- Climbed?
 - Y - Route (+ info)
 - N - Position & distance to route

Plot



- Anchor number
- Height
- Aspect
- Slope

Subplot



- Microtopography



Vegetation:

Vascular plants	Mosses	Lichens
% total cover	% total cover	% total cover
Species <ul style="list-style-type: none"> • identification • % cover 	Morphotype Harvest fragment from each cushion + %cover	Morphotype <ul style="list-style-type: none"> • Crustose • Foliose • Fruticose + %cover