

1 Remote mapping of foodscapes using sUAS and a low cost BG- 2 NIR sensor

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26 **Abstract**

27 The assessment of landscape condition for large herbivores, also known as
28 foodscapes, is fast gaining interest in conservation and landscape management
29 programs worldwide. Although traditional approaches are now being replaced by
30 satellite imagery, several technical issues still need to be addressed before full
31 standardization of remote sensing methods for these purposes. We present a low-cost
32 method, based on the use of a modified blue/green/near-infrared (BG-NIR) camera
33 housed on a small-Unmanned Aircraft System (sUAS), to create foodscapes for a
34 generalist Mediterranean ungulate: the Iberian Ibex (*Capra pyrenaica*) in Northeast
35 Spain. Faecal cuticle micro-histological analyses were used to assess the dietary
36 preferences of ibexes and then individuals of the most common plant species (n = 19)
37 were georeferenced to use as test samples. Because of the seasonal pattern in
38 vegetation activity, based on the NDVI (Smooth term_{Month} = 21.5, p-value < 0.01, R²
39 = 43%, from a GAM), images were recorded in winter and spring to represent
40 contrasting vegetation phenology using two flight heights above ground level (30 and
41 60 m). Additionally, the range of image pixel sizes was 3.5-30 cm with the smallest
42 pixel size representing the highest resolution. Boosted Trees were used to classify
43 plant taxa based on spectral reflectance and create a foodscape of the study area.
44 The number of target species, the sampling season, the height of flight and the image
45 resolution were analysed to determine the accuracy of mapping the foodscape. The
46 highest classification error (70.66%) was present when classifying all plant species
47 using a 30cm pixel size from acquisitions at 30 m height. The lowest error (18.7%),
48 however, was present when predicting plants preferred by ibexes, at 3.5 cm pixel size
49 acquired at 60 m height. This methodology can help to successfully monitor food
50 availability and seasonality and to identify individual species.

51

52 **Key words:** *Capra pyrenaica*, Food resources monitoring, Remote sensing, sUAS,
53 Vegetation assessment

54

55 **1. Introduction**

56 The spatiotemporal assessment of food resources for large herbivores, also called
57 foodscapes, is fast gaining interest within landscape and wildlife management, and
58 associated research agendas worldwide. Knowing the distribution and availability of
59 specific plants (i.e., used by ungulates see Espunyes et al., 2019), has become
60 essential not only to understand population dynamics of herbivores but also those of
61 their predators (Oates et al., 2019; Peters et al., 2019; Searle et al., 2007). Mapping
62 vegetation is also essential for ecologists (Moore et al., 2010) and wildlife conservation
63 agents (Schweiger et al., 2015a). Likewise, plant biodiversity conservation strongly
64 depends on the interaction between plants and large herbivores, among others,
65 (Boulanger et al., 2018; Royo et al., 2017). The modeling of these interactions is fast
66 gaining interest (Weisberg and Coughenour, 2006), particularly by exploring plant
67 communities at different spatial and temporal scales (Golodets et al., 2011; Schweiger
68 et al., 2015a). Assessment of foodscapes is also of major concern for forest managers,
69 as both wild and domestic ungulates influence forest regeneration, tree growth and
70 forest development (Bergqvist et al., 2018; Rooney et al., 2015; Valle Júnior et al.,
71 2019). Vegetation modulates both habitat and diet selection in large herbivores, which
72 in turn affects the sign (mutualist vs. antagonist), the strength of plant-herbivore
73 interactions (Gill, 1992; López-Sánchez et al., 2016; Perea et al., 2013) and their
74 associated ecosystem services and disservices. In addition, ungulate species serve
75 societal needs as game animals or subsistence foods (Carvalho et al., 2020), and can
76 also affect agricultural crops which add importance to the understanding of nutritional
77 resources and habitat use of large herbivores (Rowland et al., 2018; Duparc et al.,
78 2019).

79 Traditionally, vegetation cover mapping has been done using field-based or *in situ*
80 measurements (Karl et al., 2011). Although field studies are still required for
81 calibration, and validation of other indirect approaches (e.g., based on remote
82 sensing), they are time-consuming and often unfeasible when covering large areas.
83 This is complicated further when studying complex landscapes across different
84 seasons and years (Manousidis et al., 2016; Moore et al., 2010; Petersen et al., 2014;
85 Royo et al., 2017). Remote sensing is thus becoming the main alternative to overcome
86 the limitations of, and complement the advantages of field-based studies (Kerr and
87 Ostrovsky, 2003; Pettorelli, 2013; Sankey et al., 2019; Skidmore et al., 2010). In fact,
88 satellite derived-measurements are becoming popular for mapping vegetation cover,
89 structure, composition, and condition in wide geographic areas and over long time
90 periods (Harris et al., 2014, Wachendorf et al., 2017). However, several technical
91 issues need to be addressed before remote sensing approaches can be fully
92 established to create foodscapes. In homogeneous landscapes, for example, remote
93 sensing methods are useful for mapping specific food resources e.g., lichens in tundra
94 used by reindeer (see Falldorf et al., 2014), or linking the greenness of mixed-grass
95 communities to diet quality of alpine ungulates (Schweiger et al., 2015b; Villamuelas
96 et al., 2016). Few efforts, however, have been made to create foodscapes in complex,
97 bushy and encroached landscapes such as those common in the Mediterranean
98 region. To date, the most ambitious contributions in this area of science have achieved
99 the assessment of the nutritional quality of specific tree species commonly used by
100 African ungulates (Skidmore et al., 2010), and by Australian marsupials (Youngentob
101 et al., 2012). Implementation and evaluation of the use of remote sensing to assess
102 the availability of specific woody species used by ungulates in complex landscapes
103 are still scarce but necessary.

104 One alternative, to gain definition in such heterogeneous environments, is the use of
105 hyperspectral sensors set in unmanned aircraft systems (UAS, see Beerli et al., 2007;
106 Schweiger et al., 2015b; Skidmore et al., 2010; Youngentob et al., 2012), or in
107 combination with lidar (Insua et al., 2019; Lone et al., 2014; Pullanagari et al., 2018).
108 Lamentably, these approaches are still expensive and remain outside of the budget
109 capabilities of most organizations and companies, as well as government-funded
110 bodies. Additionally, they show complexities that require additional expertise which
111 inhibits the broader uptake and use by non-expert researchers and land managers
112 alike. Thus, timely research in this field of remote sensing is required to support
113 decision making, oriented towards the most appropriate, timely and low-cost methods
114 available while also offering suitably high levels of accuracy (Wachendorf et al., 2017).
115 Furthermore, although there are studies focused on estimating primary production and
116 nutrient content, few of them integrate quality assessment and dietary species
117 identification (food availability), and almost no studies focus simultaneously on
118 heterogeneous environments and wide-ranging feeders.

119 Several methodological issues have to be solved to create foodscapes based on
120 remote sensing. For example, when using UAS mounted sensors, it is important to
121 define the altitude above ground level (AGL) to ensure coverage of the study area
122 while also balancing against the pixel size and subsequent spatial resolution
123 requirements of the study (Tommervik, 2014). Additionally, the spectral range and
124 resolution requirements must be met by the sensor to ensure that detailed spectral
125 responses for each pixel can be obtained. This is particularly relevant to accurately
126 distinguish plant species and monitor their spectral responses and variations across
127 different seasons (Hesketh and Sánchez-Azofeifa, 2012).

128 For these reasons, this study aims to address both challenges, heterogeneous
129 environmental assessment and mixed feeder diet classification, using a simple and
130 low cost 3 spectral band camera (modified to detect Blue, Green, and Near-Infrared
131 (BG-NIR)) mounted on a small-Unmanned Aircraft System (sUAS). The study seeks
132 to create a foodscape assessment for a mixed feeder ungulate, the Iberian Ibex (*Capra*
133 *pyrenaica*), in a heterogeneous Mediterranean scrubland. Many sUAS are low-cost
134 machines capable of carrying a wide range of sensors and imaging equipment to work
135 in complex and heterogeneous scenarios (Anderson and Gaston, 2013), and similar
136 sensors have successfully been employed in other vegetation studies (e.g., Gillan et
137 al., 2019; Lu and He, 2017; Strong et al., 2017).

138 The Iberian Ibex is a mixed feeder with a diet including herbaceous and woody species
139 (Alados and Escos, 1987; Martinez et al. 1985; Martinez and Martinez, 1987; R.
140 García-González, 1992; Granados et al. 2001). This mountain ungulate shows great
141 dietary plasticity but is influenced by plant phenology (Martinez, 2014; Martínez, 1994)
142 and landscape characteristics (Martínez and Martínez, 1987; Moço et al., 2014; Perea
143 et al., 2015).

144 In this paper, the primary aim is to determine the feasibility of remotely classifying
145 Mediterranean plant species, in two periods of contrasting phenology, growing in a
146 high diversity and physiognomically heterogeneous plant community, with special
147 focus on the plants grazed or browsed by the Iberian Ibex diet. We do so by using a
148 novel methodology and low budget equipment and seek to set a basis for further
149 remote studies aimed at diet quality in complex environments.

150

151 2. Materials and methods

152 2.1. Study area

153 The study was undertaken in the National Game Reserve “Ports de Tortosa i Beseit”
154 (NGRPTB) in Catalonia, northeast Spain (40°46’08” N, 0°20’04” E, 450 m. a.s.l.,
155 Figure 1). The average temperature in winter is from 0.4°C to 10°C and in summer from
156 11°C to 25°C. Precipitation is concentrated in spring and autumn, with a mean rainfall
157 of 133.7mm and 116.6 mm respectively. Summer is characterized by drought
158 conditions, with an average precipitation of 29.97mm (Meteocat, 2019).

159 The study area is characterized by a Mediterranean sclerophyllous woody landscape,
160 dominated by *Quercus ilex* and a dense scrubland integrated by more than 30 woody
161 plants codominated by *Pistacia lentiscus*, *Erica multiflora*, *Quercus coccifera*,
162 *Rosmarinus officinalis*, *Genista scorpius*, and *Ulex parviflorus*. Natural and planted
163 Pine stands and isolated individuals (*Pinus nigra* and *Pinus pinaster*) were also
164 present across the study area. More specifically, the investigation was conducted in a
165 fenced scientific enclosure of 17 ha (see Figure 1), which maintained an introduced
166 herd of 18 Iberian Ibexes. The enclosure facilitated the control of Ibex population
167 numbers and represented all relevant vegetation communities and plant species
168 typical of this Mediterranean landscape.

169

170 2.2. Field data collection

171 Field data were collected in both June 2018 and March 2019. The June period was
172 selected to evaluate plants with new and well developed shoots and leaves, and the
173 March period represented an inactive period (late winter) with plants harboring old
174 shoots and leaves. It is hypothesised that spectral variations will be apparent across

175 all species on the two dates. In particular, higher NDVI values are expected to be
176 higher in winter than in spring because of the higher photosynthetically active radiation
177 of evergreen Mediterranean plants (Garbulsky et al., 2013).

178 Inside the enclosure, 11 plots of 15 x 15m were randomly distributed to obtain a
179 representative vegetation sample from within the enclosure. A total of 19
180 representative plant species (from 12 family groups) were identified and sampled in
181 the study site within the enclosure; and the height, shape, diameter of the vegetative
182 crown and phenological stage were recorded and specifically sampled (for subsequent
183 discernment in aerial imagery, Figure 2). The location of sampled individuals was
184 recorded using a differential-GPS (Leica GS07) for image matching and all species
185 coded for analysis (see Table 1). The number of individuals sampled ranged between
186 6 and 20 dependent upon species and vegetation state (see Table 1 and Figure 2). In
187 addition, plant fragments were collected for further cuticle micro-histological analysis
188 with up to 10 plants of each diet species sampled.

189 Finally, 10 fresh Ibex faecal samples were collected from around the enclosure. Fresh
190 faecal samples were placed in individual plastic bags, labelled and then transported
191 via a cool-box to the laboratory. Once at the laboratory, samples were stored in a
192 freezer, at -20°C, until further processing.

193

194 2.3. Image acquisition and processing

195 A DJI Inspire 1 sUAS (DJI, Europe) was used to capture high-resolution aerial imagery
196 of the study area. The sUAS captured imagery in consistent weather conditions
197 (temperature *circa* 25°C, wind speed < 7ms⁻¹, clear skies), and flights were conducted

198 at solar noon. On two consecutive days in June 2018, two separate flights were
199 undertaken; with a target altitude of 30 m AGL followed by 60 m AGL. On 25th of March
200 2019, one flight was undertaken with a target altitude of 30m. All flights followed a
201 cross-hatched flight plan to ensure maximum overlap (> 80%), at a 5m line spacing. A
202 low-cost payload sensor was used to collect imagery. More specifically, the payload
203 sensor was a BG-NIR modified version of the DJI X3 RGB sensor (12 megapixels,
204 DJI, Europe). The modified sensor was adjusted using a custom filter to pass infrared
205 light from the “red edge” at 680-800 nm, where plants actively reflect wavelengths,
206 and to block wavelengths over 800 nm. The filter ensured that the blue and green
207 channels only received visible light whilst allowing the detection of NIR light at 680-
208 800 nm (LDP LLC, Carlstadt, NJ, USA).

209 In addition, a total of 34 Ground Control Points (GCP) were located within the study
210 site to georeference the image in the later processing phase. A Leica GS07 dGPS was
211 used to record their coordinates. The sUAS was flown manually, and image capture
212 ranged between 83-162 images for 60 m AGL flights, and 450-563 images for 30 m
213 AGL flights in June, and between 190-1068 images in March (see table 2). All images
214 were recorded in JPEG file format and georeferenced to EXIF GPS coordinates and
215 altitude levels obtained from the DJI Inspire 1 sUAS.

216 The Pix4D Mapper® software was used to process all images. Initially, Structure-from-
217 Motion (SfM) was used to generate a Digital Surface Model of the study site (Westoby
218 et al., 2012), and then an orthomosaic generated by orthorectification of the aerial
219 imagery (Pix4D, 2018). This method removes perspective distortions from the images
220 using the Digital Surface Model.

221 To ensure consistency across all orthomosaics, the image resolution and pixel size
222 were matched to the coarsest resolution across all final images using ArcGIS
223 (v10.5.1). A nearest neighbour sampling method was used to resample discrete pixel
224 data to larger pixel sizes, in order to test the possible influence on the image
225 classification. The range of image pixel sizes was 3.5-30 cm, with the smallest pixel
226 size representing the highest resolution consistently achievable during the sUAS
227 flights.

228 Finally, using dGPS locations of the plants sampled in the field, a buffer was generated
229 (25% of the smallest measured diameter of each specific plant) and used to extract
230 the pixel values relative to each plant individually sampled. Pixel sampling of each
231 plant was conducted in this way to avoid plant edges and soil reflectance where
232 possible which may have caused disruptive pixel mixing. The resultant data provided
233 BG-NIR pixel data for each plant sampled in the field which was used to perform
234 subsequent analysis.

235

236 2.5. NDVI mapping

237 NDVI is a good indicator to reflect plant growth, quality, and phenology in
238 Mediterranean ecosystems (Ogaya et al., 2015). NDVI represents the Normalised
239 Difference Vegetation Index which is determined by calculating the difference in
240 reflectance between the NIR band and a chosen visible band which is then divided by
241 the sum of these two bands (Pettorelli, 2013). This index can be calculated for any
242 individual pixel and typically uses the red band as the visible component due to the
243 high level of absorption of this band during photosynthesis. The NIR band is used as
244 it is highly reflected by healthy plants, allowing a strong contrast with the red visible

245 band. The blue band can be similarly used to the red band but is typically restricted to
246 low altitude data acquisitions due to the negative effect of Rayleigh scattering that can
247 interfere with satellite measurements as light passes through the atmosphere. It is
248 more commonly referred to as Blue-NDVI (BNDVI) and has been used successfully in
249 other vegetation assessment studies (Beerli et al., 2007; Lu and He, 2017). NDVI
250 sensitivity to phenological stages of the plants was assessed as they manifested
251 relevant seasonal phenological changes. Initially, satellite data were obtained to
252 determine the mean NDVI values correspondent to the 2014-18 period for the entire
253 study area, and to establish seasonal trends (MOD13Q1 NDVI data extracted from the
254 MODIS repository, Moderate Resolution Imaging Spectroradiometer, provided by
255 NASA). Secondly, the Blue-NDVI was calculated from the acquired images for all the
256 plants recorded within the study area. The highest resolution images were used for
257 these comparisons: June low flight 3.5cm pixel and March low flight 3.5 cm pixel.
258 Single-band raster images were analysed and vegetation indices calculated from the
259 sUAS captured imagery. These data were used to identify differences in vegetation
260 index values and pixel values across sampled species and previously determined
261 family groups.

262

263 2.6. Diet composition

264 Faecal cuticle micro-histological analysis was used to confirm that previous studies
265 completed on the Iberian Ibex diet are suitable for this study area. A micro-histological
266 analysis was therefore performed on the field sampled data. This technique facilitated
267 the identification of plant epidermal fragments in the 10 samples collected in
268 December 2019. Samples were prepared following treatment described by Bartolome

269 et al. (1995), with minor modifications. Approximately 10 g from the milled sample were
270 placed in test tubes with 5 ml of 65% concentrated HNO₃. The test tubes were then
271 boiled in a water bath at 80° C for 2 min. After digestion in HNO₃, the samples were
272 diluted with 200 ml of water. This suspension was then passed through 1.00 mm and
273 0.125 mm filters. The 0.125-1.00 mm fraction was spread on glass microscope slides
274 in a 50% aqueous glycerine solution and cover-slips were fixed with DPX micro-
275 histological varnish. Three slides were prepared from each sample. The slides were
276 examined under a microscope at 100-400x magnifications, conducting lengthwise
277 traverses. Plant fragments were recorded and counted until 200 fragments of leaf
278 epidermis were identified from each sample.

279

280 2.7. Classification analysis and Statistics

281 A machine learning algorithm approach was used to classify and map plant species
282 across the study site. Machine learning algorithms are effective at operating on large
283 volume and multivariate datasets. They can have high accuracy, and they have been
284 successfully used when regression models are not suitable (Li et al., 2019; Marrs and
285 Ni-Meister, 2019; Van Ewijk et al., 2014). In particular, the Breiman's Random Forests
286 Model (RFM, Breiman, 2001) has been shown to be effective in other species
287 distribution studies (see Carvalho et al., 2018; Zhang et al., 2019). Digital values of
288 the three recorded bands (Blue, Green, NIR) were used as predictor variables. The
289 effect of pixel size (3.5-30 cm), flight height, the month of sampling, and the response
290 variable (e.g., all species vs diet species only) was assessed.

291 Initially, all the plant species sampled in the area were included in the RFM
292 classification. Seventy percent of the sampled individuals were used as training data,

293 whilst the remaining 30% of sampled individuals were used as test data. The inclusion
294 of the training data subset allowed independent assessment of the error in the
295 classification method, using Out Of Bag (OOB) and Prediction Test Error (PTE). A
296 confusion matrix was created, and the predictions evaluated against the independent
297 ground-truthing data. The final error matrix was then used to select the most suitable
298 image acquisition approach and processing method.

299 Following the assessment of all plant species, a further analysis was conducted to
300 discriminate between those sampled species recognised as Ibex diet species
301 according to our micro-histological results. This step was conducted in two ways.
302 Initially, species were grouped according to diet and a secondary “other” species group
303 was included; and secondly, only species recognised as Ibex diet species were
304 included within the model, and the “other” species group was omitted from the
305 analysis. As a further investigation, and to understand the source of the error and the
306 relevance of potential methodological issues, a linear model was generated that
307 predicted the percentage of error according to the pool of species included in the
308 classification and the study season when images were captured.

309 Finally, a Principal Components Analysis (PCA) was undertaken to further explore the
310 relationship between individual species, species pools and the three bands recorded
311 by the sensor (BG-NIR). In addition, PCA score values were assessed across dietary
312 species and used to discern differences in mean scores through an ANOVA post-hoc
313 analysis.

314

315 2.8. Foodscape mapping

316 To complete the study a foodscape of the distribution of the main Iberian ibex
317 resources in the study area was created. The original digital values of the already
318 georeferenced pixels making up the orthomosaic were replaced by the predicted
319 categories based on the RFM with the lowest test error. The procedure was done using
320 the “rasterFromXYZ” function of the “Raster” package, version 3.07 (Hijmans, 2019).

321

322 3. Results

323 3.1. NDVI

324 The traditional red band NDVI obtained from MODIS for the study area for the 2014-
325 2018 period presents a clear seasonal pattern. In fact, 43% of the observed NDVI
326 variability was explained by the effect of months (Smooth term_{Month} = 21.5, p-value <
327 0.01, $R^2 = 43\%$, from a GAM). A peak in primary production was shown in late winter
328 (March) and a minimum in summer (June, Fig. 3A). March and June presented
329 statistically significant differences between their NDVI values. The spatial resolution
330 of the MODIS analysis provided a preliminary indicator of general NDVI trends in
331 response to phenology which was then examined more specifically using the finer
332 spatial resolution BG_NIR sensor.

333 In the absence of the red band, the BNDVI was used for the aerial data collection via
334 the sUAS when analysing the NDVI values obtained for each recorded plant. Similar
335 trends were obtained, across the two months studied using the sUAS mounted sensor,
336 as found in the MODIS data. The differences between the values in both recorded
337 months are shown in Fig. 3. Almost all species, 16 out of the 19 studied, have
338 statistically significant differences in NDVI value between June and March (t-student

339 test with p -value < 0.05), except *Brachypodium retusum* (BR, p -value=0.1658),
340 *Helianthemum marifolium* (H, p -value=0.9869) and *Pinus pinaster* (PP, p -
341 value=0.1182).

342

343 3.2. Faecal cuticle micro-histological analysis

344 Faecal cuticle micro-histological analyses clearly show that diet composition of Ibexes
345 within the enclosure agrees with that in the base work (Martínez, 1994). Diet of Ibexes
346 was mainly based on non-legume wood species (ONLW), *Erica multiflora* (EM),
347 graminoids (G, such as *Brachypodium phoenicoides* and *B. retusum*), Labiatae-
348 Asteraceae plants (L-A, e.g., *Rosmarinus officinalis*), and Fagaceae (e.g., *Quercus*
349 *ilex*). No plant represented more than 14% of use (Fig. 4), indicating the generalist
350 feeding behaviour of the species (Granados et al., 2001). As a result, species were
351 classified into five broad family groups with the species consistent in the diet across
352 all the year. The five groups were: *Quercus* spp. as the family Fagaceae (Group F),
353 *Rosmarinus officinalis* and *Thymus vulgaris* as the family Labiatae (Group L), *Erica*
354 *multiflora* (Group E), *Cistus albidus* (Group C), *Brachypodium phoenicoides*, *B.*
355 *retusum*, and other grass-like plants as Graminoids (Group G). The remaining species
356 were then placed into the category 'Others' that were excluded from the statistical
357 analysis.

358

359 3.3. Image capture and processing

360 The results showed that the orthomosaic images were created and georeferenced with
361 an error of between 3 cm and 28 cm (see table 2), and the resultant single-band raster
362 images used to calculate vegetation indices.

363

364 3.4. Random Forest Modelling.

365 The RFM was applied to all sUAS imagery collected in both June 2018 and March
366 2019 and assessed for classification error. The results from the RFM model indicate
367 a consistent improvement in model performance using small pixel size (e.g., error
368 reduction of 42.02% in predictions based on June flights at 60 m height and 3.5 cm
369 pixel size see Table 3). In all RFM groups, the pixel size of 3.5 cm outperformed all
370 other pixel size models. Moreover, it was found that as pixel size increased (up to 30
371 cm) model performance progressively deteriorated (Table 3).

372 Conversely, when flight height (60 m and 30 m) was compared (for similar pixel
373 resolution), the RF model performance increased when the sUAS imagery was
374 captured at a higher flight height. Although, comparatively speaking, the effect of this
375 was not as substantial as the effect of pixel size on model performance (60 m flight vs
376 30 m flight; 3.5 cm pixel size; 6.6% increase in OOB error).

377 Further differences in model performance were observed when comparisons were
378 made between survey season (June vs March). The model performance was better in
379 June 2018 than in March 2019. The June 60 m flight outperformed the March 60 m
380 flight, with a 9.5% reduction in OOB error rates (see Table 3).

381 Finally, the best performance by the RF model was observed when the response
382 variable was changed to focus on specific Ibex diet species. Simplification of the

383 analysis to include only a 'diet species' group showed that the RF model's OOB error
384 rate dropped to only 18.4%; meaning a correct prediction of plants in 81.6% of
385 instances. This pattern of error reduction was apparent in all flights using the diet
386 response variables.

387 The above analysis is supported by the linear modelling undertaken. The analysis
388 showed that both pools of species (e.g., all species and diet ones) and month (season)
389 were significant variables. Interestingly, and in support once again, the species pool
390 was shown to be the variable that most affects the error (p -value = 0.001, for the pool,
391 and p -value = 0.03 for month). However, it is worth noting that the error calculated for
392 the March 2019 surveys is higher in all pool scenarios than for June 2018 (Table 3,
393 Fig. 1. Supplementary material).

394

395 3.5. Principal Component Analysis

396 Principal Component Analysis (PCA) was performed on the image data that provided
397 the best classification results (June 2018, 30 m high flight, 3.5 cm pixel size). PCA was
398 used to understand the behaviour of bands and vegetation indices, and to identify the
399 different band responses for each species and their natural groupings. Evaluating the
400 responses of the three recorded bands for all the recorded species provided the
401 following results. The PCA axis 1 and 2 accounted for 97.9% of the variation in the
402 dataset for the three spectral bands (Blue, Green, and Near Infrared). Axis 1
403 accounted for 82% of this variation with an associated elevated eigenvalue of 2.46.
404 Axis 2 explained a further 15.9% (eigenvalue = 0.48) of the variation (Figure 2,
405 supplementary material). In addition, the score values of each species were calculated
406 (Figure 5) and demonstrated that axis 1 scores showed the greatest potential to

407 represent the variation of the spectral responses to the three bands. ANOVA test and
408 corresponding post-hoc analyses showed statistically significant differences between
409 the reflectance values of all plant species. However, it is worth noting that, this could
410 be a false discovery rate due to effect size; as there were many sampled individuals
411 (246) and therefore many pixels (up to 66264, in the data set of June 2018, 30 m flight
412 height, and 3.5 cm pixel size, high flight, and June). To further understand the
413 variability within species the above analysis was repeated accounting for solely the
414 dietary species.

415 Evaluating the spectral response of each individual to the three recorded bands
416 provided the following results. The PCA axis 1 and 2 accounted for the same
417 proportion (97.9%) of the variation observed in the spectral responses across all
418 individuals. Axis 1 accounted for 82% of the variation (eigenvalue = 2.50), and axis 2
419 explained a further 15% (eigenvalue = 0.44) of the variation (Figure 3, supplementary
420 material). In addition, the score values of each plant group were calculated to illustrate
421 the differences. Post-hoc analysis revealed significant differences between all dietary
422 groups except for the pair *Labiatae* and *Erica multiflora* (p-value = 0.0628, Figure 6).

423

424 3.6. Foodscape mapping

425 The foodscape map on Fig. 7 predicts that 13.04% of the study area is covered by
426 Fagaceae plants (Group F), 4.08% by graminoids (Group G), 2.23% by Labiateae
427 (e.g., *Rosmarinus* and *Thymus*, Group L), 1.77% by *Erica multiflora* (Group E) and
428 0.02% by *Cistus albidus* (Group C). Most of the area (78.06%), however, included
429 other plants not found in the Ibex diet, bare soil and rocks.

430

431 4. Discussion

432 This study has successfully explored the application of a modified BG-NIR sensor,
433 mounted on a low cost sUAS, to create foodscapes for a mixed feeder in a
434 Mediterranean scrubland, rich in woody species with very different physiognomies.

435 Analysis has shown successful discrimination of species within a species rich and
436 physiognomically heterogeneous Mediterranean scrubland. Plant prediction was
437 correct in 81.6% of instances (prediction error 18.4%) when using a training data set
438 of 173 individuals (73 in the test dataset) representing all the most abundant species
439 at the study site. Plant prediction improved further to 88.2% when focussing on the
440 Ibex diet species alone (prediction error 11.8%) using a training data set of 97
441 individuals (and a test dataset of 41). Such errors are similar or, in some case, lower
442 than those obtained by using hyperspectral methodologies in complex ecosystems
443 (e.g., Clark et al., 2005; Ferreira et al., 2016), including Mediterranean ones (e.g.,
444 (Manevski et al., 2012).

445 Our classification results along with those of the PCA analysis, show a wide range of
446 spectral responses across the species investigated, but also a useable level of
447 homogeneity within species groups; when considering both 'all species' and only those
448 'within the Ibex diet'. These results indicate that successful classification is possible.
449 The success of using only 3 spectral bands to identify significantly different spectral
450 responses across species and similarities within species groups, coupled with the high
451 level accuracy of classification, outline the feasibility and potential for such a low-cost
452 application to develop foodscapes and facilitate their long-term management.

453 The success in discriminating Ibex diet species remotely provides a strong foundation
454 from which to discuss the technical issues in the classification process. Initially, the
455 results indicate that small pixel size provides better classification results; as previously
456 reported in other studies (Hsieh et al., 2001; Hu et al., 2019) which discuss the
457 relevance of pixel size in classification performance and the effect of variations in plant
458 size and ground coverage. However, for the same pixel size, higher flights have shown
459 to perform better than lower flights, which is similar to other methodological studies
460 (e.g., Mesas-Carrascosa et al., 2015). The study by Mesas-Carrascosa et al. (2015)
461 indicated that there was greater discrimination between species at higher flights. In
462 addition, higher altitudes typically have less perspective distortion as the ratio of the
463 topographic change to flying altitude is smaller; hence, the accuracy of the orthophoto
464 can be higher; as is reflected in the work of Nesbit and Hugenholtz, (2019). Therefore,
465 when choosing the ideal flight parameters (as pixel size is determined by both sensor
466 type and flight height) a balance between both must be found; taking into account the
467 area to evaluate, the flight platform employed, and the financial and time costs.

468 This study has also shown that season is a relevant factor when working with plant
469 species in temperate regions since they manifest different physiological status and
470 even morphologies throughout the year (see also Sperlich et al., 2014., Vogt and
471 Gerhard Gul, 1994). Our results show that the most suitable season to perform the
472 classification study at this site is June (vs March) when most of the Mediterranean
473 woody taxa end their growth and flowering. Thus, plants have a large part of their
474 vegetative structure (not woody) renewed, with its characteristic morphology. At a
475 tissue and chemical level, the Mediterranean woody plants are fully constituted to be
476 able to withstand the water deficit and the severe heat stroke of the summer that starts
477 in June (Fernández-Marín et al., 2017). All these characteristics along with net inter-

478 species differences in water content (related to drought adaptations) determine the
479 optical properties of the plants (Manevski et al., 2012). Due to its influence on the
480 classification results the importance of prior knowledge of the phenological pattern of
481 the area and the present plant species could be crucial to applying this methodology
482 over a wider area and for it to not be restricted by site specific conditions.

483 In our study the classification results improved when focusing on the target diet
484 species (Prediction error reduced to 11.8%). The result showed that the pool of training
485 species influenced classification success with accuracy reported of 88.2%. This clearly
486 indicates that an informed knowledge of the target diet species for the study area is of
487 great value to maximise classification accuracy. Our results show that the
488 classification accuracy can be improved by over 6%. Moreover, the importance of the
489 micro-histological analysis or other related diet studies is evident in this process.
490 However, caution is needed here given the associated reduction in both test and
491 training data samples. The micro-histological analysis has served as an effective tool,
492 focussing and reducing the pool of input species identified as diet species and
493 subsequently improving the relevance and overall classification performance.
494 Reducing the number of diet species classes by combining similar species, for
495 example logically grouping *Pinus pinaster* and *P. nigra* as Pines, and *Brachypodium*
496 *phoenicoides* and *B. restusum* as Graminoids offered further improvement in
497 classification accuracy but was limited to an increase of 0.5-2%, with no significant
498 improvement given the reduced specificity of the classification.

499 Regarding the obtained error, it is also relevant to consider the size (and biotype) of
500 the plants. This seems to be case of *Helianthemum marifolium*, a representative taxon
501 of typical prostrated chamaephytic communities on Mediterranean bedrocks. These
502 small prostrated plants, with diameters around the calibration error (*circa* 20 cm) and

503 loose branching on bare soils, appeared to yield higher errors in classification
504 compared to those of bigger and thicker plants. These finding are similar to other
505 recent, and related, studies in rangeland ecosystems where small shrub/subshrub
506 species, with low abundance, were found to have reduced classification success
507 (Sankey et al., 2019).

508 However, with the relative successes of our study, when focussing on target diet
509 species, areas for improvement are largely focussed on the sensor rather than on the
510 methodology adopted. Improvements are most likely to be achieved by increasing the
511 sensitivity of spectral responses to plant species variation, and taking advantage of
512 the increased spectral resolution of more expensive cameras. Although the low-cost
513 aspect of this study is a unique selling point, upgrading to a more expensive
514 hyperspectral camera could offer greater flexibility, and potentially accuracy, during
515 the classification process (Manevski et al., 2012; Sankey et al., 2019). Known as
516 hyperspectral sensors these cameras can offer a greater quantity of spectral bands
517 but also bands of finer spectral resolution and have been used advantageously in
518 several other works (Beeri et al., 2007; Lone et al., 2014). However, with this
519 consideration, one of the relevant achievements of this work should not be
520 compromised. This being the creation of an accessible and affordable tool for land
521 managers and researchers to be applied in heterogeneous landscapes and on animals
522 with species-specific diet accounting for the general ecosystem characteristics
523 including plant species composition.

524

525 5. Conclusion

526 This feasibility study outlines the possibility of describing food location for a mixed
527 herbivore in a heterogeneous environment. A modified BG-NIR camera, mounted on
528 a low cost sUAS, has been shown to be successful in classifying plant species and to
529 describe the foodscapes of ungulates such as Ibex. The benefits of this study, the
530 methodology presented, and the potential for further development are applicable to
531 studies of wild and domestic herbivore distribution and welfare, predator distribution,
532 vegetation changes, and habitat biodiversity internationally (Boulanger et al., 2018;
533 Moore et al., 2010; Oates et al., 2019; Peters et al., 2019; Searle et al., 2007)

534 Further studies with hyperspectral cameras are necessary to assess the
535 improvements hypothesised, and the potential enhancement of efficiency and
536 accuracy offered by an increased quantity and refined nature of spectral information.
537 Additionally, it would be interesting to increase the number of sampling months in order
538 to fully study plant availability throughout the year. Further research should be done in
539 order to evaluate diet quality by improving and refining the affordable, accessible and
540 non-time consuming methods that would be beneficial to land managers. Moreover,
541 following the methodology describe here, we might be able to predict feeding
542 behaviour across seasons and according to relevant meteorological events and
543 landscape changes. To do this, we should also have more information about plants
544 physiology and ibex diet data across seasons. The classification results obtained in
545 this study provide a strong foundation on which to develop such diet quality studies.

546

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557 **References**

- 558 Alados, C.L., Escos, J., 1987. Relationships between movement rate, agonistic displacements
559 and forage availability in spanish ibexes (*Capra pyrenica*). *Biol. Behav.* 12, 245–255.
- 560 Anderson, K., Gaston, K.J., 2013. Lightweight unmanned aerial vehicles will revolutionize
561 spatial ecology. *Front. Ecol. Environ.* 11, 138–146. <https://doi.org/10.1890/120150>
- 562 Bartolome, J., Franch, J., Gutman, M., Seligman, N.G., 1995. Physical Factors That Influence
563 Fecal Analysis Estimates of Herbivore Diets. *J. Range Manag.* 48, 267.
564 <https://doi.org/10.2307/4002432>
- 565 Beeri, O., Phillips, R., Hendrickson, J., Frank, A.B., Kronberg, S., 2007. Estimating forage
566 quantity and quality using aerial hyperspectral imagery for northern mixed-grass prairie.
567 *Remote Sens. Environ.* 110, 216–225. <https://doi.org/10.1016/j.rse.2007.02.027>
- 568 Bergqvist, G., Wallgren, M., Jernelid, H., Bergström, R., 2018. Forage availability and moose
569 winter browsing in forest landscapes. *For. Ecol. Manage.* 419–420, 170–178.
570 <https://doi.org/10.1016/j.foreco.2018.03.049>
- 571 Boulanger, V., Dupouey, J.L., Archaux, F., Badeau, V., Baltzinger, C., Chevalier, R., Corcket,
572 E., Dumas, Y., Forgeard, F., Mårell, A., Montpied, P., Paillet, Y., Picard, J.F., Saïd, S.,
573 Ulrich, E., 2018. Ungulates increase forest plant species richness to the benefit of non-
574 forest specialists. *Glob. Chang. Biol.* 24, e485–e495. <https://doi.org/10.1111/gcb.13899>
- 575 Breiman, L., 2001. ST4_Method_Random_Forest. *Mach. Learn.* 45, 5–32.
576 <https://doi.org/10.1017/CBO9781107415324.004>
- 577 Carvalho, J., Büntgen, U.L.F., Pettorelli, N., Mentaberre, G., Boix, X.O., Eizaguirre, O., Pérez,
578 J.M., Fandos, P., Torres, R.T., Lavín, S., Fonseca, C., Serrano, E., 2020. Habitat and
579 Harvesting Practices Influence Horn Growth of Male Ibex. *J. Wildl. Manage.*
580 <https://doi.org/10.1002/jwmg.21830>
- 581 Carvalho, J., Santos, J.P.V., Torres, R.T., Santarém, F., Fonseca, C., 2018. Tree-based
582 methods: Concepts, uses and limitations under the framework of resource selection
583 models. *J. Environ. Informatics* 32, 112–124. <https://doi.org/10.3808/jei.201600352>
- 584 Clark, M.L., Roberts, D.A., Clark, D.B., 2005. Hyperspectral discrimination of tropical rain
585 forest tree species at leaf to crown scales. *Remote Sens. Environ.* 96, 375–398.
586 <https://doi.org/10.1016/j.rse.2005.03.009>
- 587 Duparc, A., Garel, M., Marchand, P., Dubray, D., Maillard, D., Loison, A., 2019. Through the
588 taste buds of a large herbivore: foodscape modeling contributes to an understanding of
589 forage selection processes. *Oikos*. <https://doi.org/10.1111/oik.06386>
- 590 Espunyes, J., Bartolomé, J., Garel, M., Gálvez-Cerón, A., Aguilar, X.F., Colom-Cadena, A.,
591 Calleja, J.A., Gassó, D., Jarque, L., Lavín, S., Marco, I., Serrano, E., 2019. Seasonal diet
592 composition of Pyrenean chamois is mainly shaped by primary production waves. *PLoS*
593 *One* 14, 1–23. <https://doi.org/10.1371/journal.pone.0210819>
- 594 Falldorf, T., Strand, O., Panzacchi, M., Tømmervik, H., 2014. Estimating lichen volume and
595 reindeer winter pasture quality from Landsat imagery. *Remote Sens. Environ.* 140, 573–
596 579. <https://doi.org/10.1016/j.rse.2013.09.027>
- 597 Fernández-Marín, B., Hernández, A., Garcia-Plazaola, J.I., Esteban, R., Míguez, F., Artetxe,
598 U., Gómez-Sagasti, M.T., 2017. Photoprotective strategies of mediterranean plants in
599 relation to morphological traits and natural environmental pressure: A meta-analytical
600 approach. *Front. Plant Sci.* 8. <https://doi.org/10.3389/fpls.2017.01051>
- 601 Ferreira, M.P., Zortea, M., Zanotta, D.C., Shimabukuro, Y.E., de Souza Filho, C.R., 2016.
602 Mapping tree species in tropical seasonal semi-deciduous forests with hyperspectral and
603 multispectral data. *Remote Sens. Environ.* 179, 66–78.
604 <https://doi.org/10.1016/j.rse.2016.03.021>

605 Garbulsky, M.F., Peñuelas, J., Ogaya, R., Filella, I., 2013. Leaf and stand-level carbon uptake
606 of a Mediterranean forest estimated using the satellite-derived reflectance indices EVI and
607 PRI. *Int. J. Remote Sens.* 34, 1282–1296. <https://doi.org/10.1080/01431161.2012.718457>

608 Gill, R.M.A., 1992. A review of damage by mammals in north temperate forests: 3. Impact on
609 trees and forests. *Forestry* 65, 363–388. <https://doi.org/10.1093/forestry/65.4.363-a>

610 Gillan, J.K., McClaran, M.P., Swetnam, T.L., Heilman, P., 2019. Estimating forage utilization
611 with drone-based photogrammetric point clouds. *Rangel. Ecol. Manag.* 72, 575–585.
612 <https://doi.org/10.1016/j.rama.2019.02.009>

613 Golodets, C., Kigel, J., Sternberg, M., 2011. Plant diversity partitioning in grazed
614 Mediterranean grassland at multiple spatial and temporal scales. *J. Appl. Ecol.* 48, 1260–
615 1268. <https://doi.org/10.1111/j.1365-2664.2011.02031.x>

616 Granados, J.E., Pérez, J.M., Márquez, F.J., Serrano, E., Soriguer, R.C., Fandos, P., 2001. LA
617 CABRA MONTÉS (Capra. Galemys 13, 3–37.

618 Harris, A., Carr, A.S., Dash, J., 2014. Remote sensing of vegetation cover dynamics and
619 resilience across southern Africa. *Int. J. Appl. Earth Obs. Geoinf.* 28, 131–139.
620 <https://doi.org/10.1016/j.jag.2013.11.014>

621 Hesketh, M., Sánchez-azofeifa, G.A., 2012. Remote Sensing of Environment The effect of
622 seasonal spectral variation on species classification in the Panamanian tropical forest.
623 *Remote Sens. Environ.* 118, 73–82. <https://doi.org/10.1016/j.rse.2011.11.005>

624 Hijmans, R.J., 2019. Introduction to the 'raster' package (version 3.0-7) 1–250.

625 Hsieh, P.F., Lee, L.C., Chen, N.Y., 2001. Effect of spatial resolution on classification errors of
626 pure and mixed pixels in remote sensing. *IEEE Trans. Geosci. Remote Sens.* 39, 2657–
627 2663. <https://doi.org/10.1109/36.975000>

628 Hu, P., Guo, W., Chapman, S.C., Guo, Y., Zheng, B., 2019. Pixel size of aerial imagery
629 constrains the applications of unmanned aerial vehicle in crop breeding. *ISPRS J.*
630 *Photogramm. Remote Sens.* 154, 1–9. <https://doi.org/10.1016/j.isprsjprs.2019.05.008>

631 Insua, J.R., Utsumi, S.A., Basso, B., 2019. Estimation of spatial and temporal variability of
632 pasture growth and digestibility in grazing rotations coupling unmanned aerial vehicle
633 (UAV) with crop simulation models. *PLoS One* 14, 1–13.
634 <https://doi.org/10.1371/journal.pone.0212773>

635 Karl, J.W., Colson, K., Swartz, H., 2011. Rangeland Assessment and Monitoring Methods
636 Guide: An interactive tool for selecting methods for assessment and monitoring.
637 *Rangelands* 33, 48–54. <https://doi.org/10.2111/1551-501X-33.4.48>

638 Kerr, J.T., Ostrovsky, M., 2003. From space to species: Ecological applications for remote
639 sensing. *Trends Ecol. Evol.* 18, 299–305. [https://doi.org/10.1016/S0169-5347\(03\)00071-5](https://doi.org/10.1016/S0169-5347(03)00071-5)

640

641 Li, Z., Zan, Q., Yang, Q., Zhu, D., Chen, Y., Yu, S., 2019. Remote Estimation of Mangrove
642 Aboveground Carbon Stock at the Species Level Using a Low-Cost Unmanned Aerial
643 Vehicle System. *Remote Sens.* 11, 1018. <https://doi.org/10.3390/rs11091018>

644 Lone, K., Van Beest, F.M., Mysterud, A., Gobakken, T., Milner, J.M., Ruud, H.P., Loe, L.E.,
645 2014. Improving broad scale forage mapping and habitat selection analyses with airborne
646 laser scanning: The case of moose. *Ecosphere* 5, 1–22. <https://doi.org/10.1890/ES14-00156.1>

647

648 López-Sánchez, A., Perea, R., Dirzo, R., Roig, S., 2016. Livestock vs. wild ungulate
649 management in the conservation of Mediterranean dehesas: Implications for oak
650 regeneration. *For. Ecol. Manage.* 362, 99–106.
651 <https://doi.org/10.1016/j.foreco.2015.12.002>

652 Lu, B., He, Y., 2017. Species classification using Unmanned Aerial Vehicle (UAV)-acquired
653 high spatial resolution imagery in a heterogeneous grassland. *ISPRS J. Photogramm.*
654 *Remote Sens.* 128, 73–85. <https://doi.org/10.1016/j.isprsjprs.2017.03.011>

655 Manevski, K., Manakos, I., Petropoulos, G.P., Kalaitzidis, C., 2012. Spectral discrimination of
656 mediterranean maquis and phrygana vegetation: Results from a case study in Greece.
657 IEEE J. Sel. Top. Appl. Earth Obs. Remote Sens. 5, 604–616.
658 <https://doi.org/10.1109/JSTARS.2012.2190044>

659 Manousidis, T., Kyriazopoulos, A.P., Parissi, Z.M., Abraham, E.M., Korakis, G., Abas, Z.,
660 2016. Grazing behavior , forage selection and diet composition of goats in a
661 Mediterranean woody rangeland. Small Rumin. Res.
662 <https://doi.org/10.1016/j.smallrumres.2016.11.007>

663 Marrs, J., Ni-Meister, W., 2019. Machine Learning Techniques for Tree Species Classification
664 Using Co-Registered LiDAR and Hyperspectral Data. Remote Sens. 11, 819.
665 <https://doi.org/10.3390/rs11070819>

666 Martínez, T., 2014. Diet selection by Spanish ibex in early summer in Sierra Nevada. Acta
667 Theriol. (Warsz). 45, 335–346. <https://doi.org/10.4098/at.arch.00-33>

668 Martínez, T., 1994. Dieta estacional de la cabra montés (*Capra pyrenaica*) en los Puertos de
669 Tortosa y Beceite (área Mediterránea del Nordeste de España). Ecología 8, 373–380.

670 Martínez, T., Martínez, E., 1987. in spring and summer at the Sierra de Credos , Spain.

671 Martínez, T., Martínez, E., Fandos, P., 1985. Composition of the food of the Spanish Wild Goat
672 in Sierras de Cazorla and Segura, Spain. Acta Theriol. (Warsz). 30, 461–494.
673 <https://doi.org/10.4098/at.arch.85-31>

674 Mesas-Carrascosa, F.J., Torres-Sánchez, J., Clavero-Rumbao, I., García-Ferrer, A., Peña, J.M.,
675 Borra-Serrano, I., López-Granados, F., 2015. Assessing optimal flight parameters for
676 generating accurate multispectral orthomosaicks by uav to support site-specific crop
677 management. Remote Sens. 7, 12793–12814. <https://doi.org/10.3390/rs71012793>

678 Meteocat, 2019. Anuari de dades meteorològiques - PN dels Ports, X5 (Baix Ebre). Serv.
679 Meteorològic Catalunya. <http://www.meteo.cat/wpweb/climatologia/serveis-i-dades-climatiques/anuaris-de-dades-meteorologiques/xarxa-destacions-meteorologiques-automatiques/>.

682 Moço, G., Serrano, E., Guerreiro, M., Ferreira, A.F., Petrucci-Fonseca, F., Santana, D., Maia,
683 M.J., Soriguer, R.C., Pérez, J.M., 2014. Does livestock influence the diet of Iberian ibex
684 *Capra pyrenaica* in the Peneda-Gerês National Park (Portugal)? Mammalia 78, 393–399.
685 <https://doi.org/10.1515/mammalia-2013-0139>

686 Moore, B.D., Lawler, I.R., Wallis, I.R., Beale, C.M., Foley, W.J., 2010. Palatability mapping:
687 A koala’s eye view of spatial variation in habitat quality. Ecology 91, 3165–3176.
688 <https://doi.org/10.1890/09-1714.1>

689 Nesbit, P.R., Hugenholtz, C.H., 2019. Enhancing UAV-SfM 3D model accuracy in high-relief
690 landscapes by incorporating oblique images. Remote Sens. 11.
691 <https://doi.org/10.3390/rs11030239>

692 Oates, B.A., Merkle, J.A., Kauffman, M.J., Dewey, S.R., Jimenez, M.D., Vartanian, J.M.,
693 Becker, S.A., Goheen, J.R., 2019. Antipredator response diminishes during periods of
694 resource deficit for a large herbivore. Ecology 100, 1–8. <https://doi.org/10.1002/ecy.2618>

695 Ogaya, R., Barbeta, A., Başnou, C., Peñuelas, J., 2015. Satellite data as indicators of tree
696 biomass growth and forest dieback in a Mediterranean holm oak forest. Ann. For. Sci. 72,
697 135–144. <https://doi.org/10.1007/s13595-014-0408-y>

698 Perea, R., Delibes, M., Polko, M., Suárez-Esteban, A., Fedriani, J.M., 2013. Context-dependent
699 fruit-frugivore interactions: Partner identities and spatio-temporal variations. Oikos 122,
700 943–951. <https://doi.org/10.1111/j.1600-0706.2012.20940.x>

701 Perea, R., Perea-García-Calvo, R., Díaz-Ambrona, C.G., San Miguel, A., 2015. The
702 reintroduction of a flagship ungulate *Capra pyrenaica*: Assessing sustainability by
703 surveying woody vegetation. Biol. Conserv. 181, 9–17.
704 <https://doi.org/10.1016/j.biocon.2014.10.018>

705 Peters, W., Hebblewhite, M., Mysterud, A., Eacker, D., Hewison, A.J.M., Linnell, J.D.C.,
706 Focardi, S., Urbano, F., De Groeve, J., Gehr, B., Heurich, M., Jarnemo, A., Kjellander,
707 P., Kröschel, M., Morellet, N., Pedrotti, L., Reinecke, H., Sandfort, R., Sönnichsen, L.,
708 Sunde, P., Cagnacci, F., 2019. Large herbivore migration plasticity along environmental
709 gradients in Europe: life-history traits modulate forage effects. *Oikos* 128, 416–429.
710 <https://doi.org/10.1111/oik.05588>

711 Petersen, C.A., Villalba, J.J., Provenza, F.D., Petersen, C.A., Villalba, J.J., Provenza, F.D.,
712 2014. Influence of Experience on Browsing Sagebrush by Cattle and Its Impacts on Plant
713 Community Structure Influence of Experience on Browsing Sagebrush by Cattle and Its
714 Impacts on Plant Community Structure 67, 78–87. <https://doi.org/10.2111/REM-D-13-00038.1>

716 Pettorelli, N., 2013. Satellite Data-Based Indices to Monitor Land Use and Habitat Changes,
717 in: *Biodiversity Monitoring and Conservation: Bridging the Gap between Global*
718 *Commitment and Local Action*. pp. 95–119. <https://doi.org/10.1002/9781118490747.ch5>

719 Pullanagari, R.R., Kereszturi, G., Yule, I., 2018. Integrating airborne hyperspectral,
720 topographic, and soil data for estimating pasture quality using recursive feature
721 elimination with random forest regression. *Remote Sens.* 10.
722 <https://doi.org/10.3390/rs10071117>

723 R. García-González, P.C., 1992. 37_Feeding_Strat.Pdf.

724 Rooney, T., 2015. Integrating Ungulate Herbivory into Forest Landscape Restoration.

725 Rowland, M.M., Wisdom, M.J., Nielson, R.M., Cook, J.G., Cook, R.C., Johnson, B.K., Coe,
726 P.K., Hafer, J.M., Naylor, B.J., Vales, D.J., Anthony, R.G., Cole, E.K., Danilson, C.D.,
727 Davis, R.W., Geyer, F., Harris, S., Irwin, L.L., Mccoy, R., Pope, M.D., Sager-fradkin, K.,
728 Vavra, M., Hellgren, E.C., 2018. Modeling elk nutrition and habitat use in Western
729 Oregon and Washington. *Wildl. Monogr.* 199, 1–69.

730 Royo, A.A., Kramer, D.W., Miller, K. V., Nibbelink, N.P., Stout, S.L., 2017. Spatio-temporal
731 variation in foodscapes modifies deer browsing impact on vegetation. *Landsc. Ecol.* 32,
732 2281–2295. <https://doi.org/10.1007/s10980-017-0568-x>

733 Sankey, T.T., Leonard, J.M., Moore, M.M., 2019. Unmanned Aerial Vehicle – Based
734 Rangeland Monitoring: Examining a Century of Vegetation Changes. *Rangel. Ecol.*
735 *Manag.* 72, 858–863. <https://doi.org/10.1016/j.rama.2019.04.002>

736 Schweiger, A.K., Risch, A.C., Damm, A., Kneubühler, M., Haller, R., Schaepman, M.E.,
737 Schütz, M., 2015a. Using imaging spectroscopy to predict above-ground plant biomass in
738 alpine grasslands grazed by large ungulates. *J. Veg. Sci.* 26, 175–190.
739 <https://doi.org/10.1111/jvs.12214>

740 Schweiger, A.K., Schütz, M., Anderwald, P., Schaepman, M.E., Kneubühler, M., Haller, R.,
741 Risch, A.C., 2015b. Foraging ecology of three sympatric ungulate species - behavioural
742 and resource maps indicate differences between chamois, ibex and red deer. *Mov. Ecol.*
743 3, 6. <https://doi.org/10.1186/s40462-015-0033-x>

744 Searle, K.R., Hobbs, N.T., Gordon, I.J., 2007. It’s the “Foodscape”, not the Landscape: Using
745 Foraging Behavior to Make Functional Assessments of Landscape Condition. *Isr. J. Ecol.*
746 *Evol.* 53, 297–316. <https://doi.org/10.1560/ijee.53.3.297>

747 Skidmore, A.K., Ferwerda, J.G., Mutanga, O., Van Wieren, S.E., Peel, M., Grant, R.C., Prins,
748 H.H.T., Balcik, F.B., Venus, V., 2010. Forage quality of savannas - Simultaneously
749 mapping foliar protein and polyphenols for trees and grass using hyperspectral imagery.
750 *Remote Sens. Environ.* 114, 64–72. <https://doi.org/10.1016/j.rse.2009.08.010>

751 Sperlich, D., Chang, C.T., Peñuelas, J., Gracia, C., Sabaté, S., 2014. Foliar photochemical
752 processes and carbon metabolism under favourable and adverse winter conditions in a
753 Mediterranean mixed forest, Catalonia (Spain). *Biogeosciences Discuss.* 11, 9697–9759.
754 <https://doi.org/10.5194/bgd-11-9697-2014>

755 Stout, S.L., Kramer, D.W., Miller, K. V, Nibbelink, N.P., 2018. Spatio-temporal variation in
756 foodscapes modifies deer browsing impact on vegetation Spatio-temporal variation in
757 foodscapes modifies deer browsing impact on vegetation. *Landsc. Ecol.*
758 <https://doi.org/10.1007/s10980-017-0568-x>

759 Strong, C.J., Burnside, N.G., Llewellyn, D., 2017. The potential of small-Unmanned Aircraft
760 Systems for the rapid detection of threatened unimproved grassland communities using
761 an Enhanced Normalized Difference Vegetation Index. *PLoS One* 12, 1–16.
762 <https://doi.org/10.1371/journal.pone.0186193>

763 Tommervik, H., 2014. Use of Unmanned Aircraft Systems (Uas) in a Multi-Scale Vegetation
764 Index Study 75, 47–52. <https://doi.org/10.12760/02-2014-1-09>

765 Valle Júnior, R.F. do, Siqueira, H.E., Valera, C.A., Oliveira, C.F., Sanches Fernandes, L.F.,
766 Moura, J.P., Pacheco, F.A.L., 2019. Diagnosis of degraded pastures using an improved
767 NDVI-based remote sensing approach: An application to the Environmental Protection
768 Area of Uberaba River Basin (Minas Gerais, Brazil). *Remote Sens. Appl. Soc. Environ.*
769 14, 20–33. <https://doi.org/10.1016/J.RSASE.2019.02.001>

770 Van Ewijk, K.Y., Randin, C.F., Treitz, P.M., Scott, N.A., 2014. Predicting fine-scale tree
771 species abundance patterns using biotic variables derived from LiDAR and high spatial
772 resolution imagery. *Remote Sens. Environ.* 150, 120–131.
773 <https://doi.org/10.1016/j.rse.2014.04.026>

774 Villamuelas, M., Fernández, N., Albanell, E., Gálvez-Cerón, A., Bartolomé, J., Mentaberre,
775 G., López-Olvera, J.R., Fernández-Aguilar, X., Colom-Cadena, A., López-Martín, J.M.,
776 Pérez-Barbería, J., Garel, M., Marco, I., Serrano, E., 2016. The Enhanced Vegetation
777 Index (EVI) as a proxy for diet quality and composition in a mountain ungulate. *Ecol.*
778 *Indic.* 61, 658–666. <https://doi.org/10.1016/j.ecolind.2015.10.017>

779 Vogt, T., Gerhard Gul, P., 1994. Accumulation of flavonoids during leaf development in *Cistus*
780 *laurifolius*. *Phytochemistry* 36, 591–597. [https://doi.org/10.1016/S0031-9422\(00\)89780-](https://doi.org/10.1016/S0031-9422(00)89780-0)
781 0

782 Wachendorf, M., Fricke, T., Möckel, T., 2017. Remote sensing as a tool to assess botanical
783 composition, structure, quantity and quality of temperate grasslands. *Grass Forage Sci.*
784 73, 1–14. <https://doi.org/10.1111/gfs.12312>

785 Weisberg, P.J., Coughenour, M.B., 2006. Modelling of large herbivore – vegetation
786 interactions in a landscape context.

787 Westoby, M.J., Brasington, J., Glasser, N.F., Hambrey, M.J., Reynolds, J.M., 2012. “Structure-
788 from-Motion” photogrammetry: A low-cost, effective tool for geoscience applications.
789 *Geomorphology* 179, 300–314. <https://doi.org/10.1016/j.geomorph.2012.08.021>

790 Youngentob, K.N., Renzullo, L.J., Held, A.A., Jia, X., Lindenmayer, D.B., Foley, W.J., 2012.
791 Using imaging spectroscopy to estimate integrated measures of foliage nutritional quality.
792 *Methods Ecol. Evol.* 3, 416–426. <https://doi.org/10.1111/j.2041-210X.2011.00149.x>

793 Zhang, L., Huettmann, F., Liu, S., Sun, P., Yu, Z., Zhang, X., Mi, C., 2019. Classification and
794 regression with random forests as a standard method for presence-only data SDMs: A
795 future conservation example using China tree species. *Ecol. Inform.* 52, 46–56.
796 <https://doi.org/10.1016/j.ecoinf.2019.05.003>

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799 **TABLES**

800 **Table1.** Species and number of individuals (n) sampled in NGRPTB Iberian ibex enclosure. Species
 801 are grouped by families and given a code to simplify the interpretation of the consequent analysis. Plant
 802 life-forms agree with Raunkier's classification (1934).

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Family	Specie	CODE	n	Physiognomy
Anacardiaceae	<i>Pistacia lentiscus</i>	PL	20	Evergreen microphanerophyte with compound broad-leaves
Buxaceae	<i>Buxus sempervirens</i>	BS	6	Evergreen microphanerophyte with simple broad-leaves
Cistaceae	<i>Cistus albidus</i>	CA	8	Evergreen nanophanerophyte with simple broad-leaves covered by dense white tomentous
	<i>Helianthemum marifolium</i>	H	8	Evergreen loosely branched chamaephyte with tiny (< 10x10 mm) simple and hairy broad-leaves
Cupressaceae	<i>Juniperus oxycedrus</i>	JO	20	Evergreen microphanerophyte with simple needle-leaves.
Ericaceae	<i>Erica multiflora</i>	EM	20	Evergreen nanophanerophyte with tiny (< 10 mm length) simple linear-leaves
Fagaceae	<i>Quercus coccifera</i>	QC	20	Evergreen microphanerophyte with simple broad-leaves
	<i>Quercus ilex</i>	QI	20	Evergreen mesophanerophyte with simple broad-leaves
Labiatae	<i>Rosmarinus officinalis</i>	RO	20	Evergreen nanophanerophyte with simple linear-leaves
	<i>Thymus vulgaris</i>	TV	20	Evergreen chamaephyte with tiny (< 5 mm length) simple linear-leaves
Leguminosae	<i>Genista scorpius</i>	GS	8	Evergreen thorny and promptly leafless nanophanerophyte
	<i>Ulex parviflorus</i>	UP	11	
Oleaceae	<i>Phillyrea angustifolia</i>	PA	7	Evergreen microphanerophyte with simple lanceolate broad-leaves
Pinaceae	<i>Pinus nigra</i>	PN	10	Evergreen mesophanerophyte with simple needle-leaves
	<i>Pinus pinaster</i>	PP	10	
Palmae	<i>Chamaerops humilis</i>	CH	8	Evergreen nanophanerophyte with compound fan-like leaves
Graminoids	<i>Brachypodium phoenicoides</i>	BP	10	Evergreen tussock-like hemicryptophyte
	<i>Brachypodium retusum</i>	BR	10	
	Graminoids	G	10	

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805 **Table2.** Image processing information for each flight executed in NGRPTB Iberian Ibex enclosure. The
 806 table includes season, flight height, surface recorded by the image, number of ground control points
 807 (GCP), calibration error (mean RMS error), number of recorded images and number of images used to
 808 create the model.

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Month	Flight (m)	Surface (ha)	GCP	Mean RMS error (m)	No. Images	No. Calibrated images
June	30	2.8457	18	0.04-0.28	1013	868
June	60	5.8915	16	0.23-0.08	245	245
March	30	5.1267	21	0.03-0.07	2188	2152

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824 **Table 3.** Plant species classification results obtained with a set of random forest models using digital
825 values of the three bands recorded by a BG-NIR camera (NIR, Blue and Green). Data collected using
826 a sUAS flying in July (2018) and March (2019) in the Iberian Ibex (*Capra pyrenaica*) enclosure in the
827 NGRPTB. Images were recorded at two heights (30 and 60 m) and different resolutions (3.5, 5, 10 and
828 30 cm pixel size). OOB error is the classification error according to the random forest algorithm. Test
829 error is the error obtained when comparing algorithm prediction results with test data not included in the
830 model development. We performed two kinds of classification: for the most abundant plants in the
831 enclosure (n = 19, All species), and for plants preferred by Ibexes (Diet). In the Diet group, plants have
832 been grouped in 4 types namely: *Cistus albidus*, *Erica multiflora*, Fagaceae (*Quercus coccifera* and *Q.*
833 *ilex*) and Graminoids (e.g., *Brachypodium phoenicoides*, *B. retusum* and other graminoids).

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Response variables	Flight	Month	Resolution (cm)	OOB error (%)	Test error (%)
All species	low	March	30	70.89	70.66
	high	June	30	66.49	70.28
	low	March	10	69.14	68.56
	low	June	10	55.50	56.07
	low	June	5	53.24	52.73
	low	March	5	50.89	50.43
	high	June	10	50.03	49.19
	low	March	3.5	48.38	48.57
	high	June	5	43.54	43.77
	low	June	3.5	35.39	35.64
	low	June	30	68.99	29.78
	high	June	3.5	28.77	25.26
	Diet	low	March	3.5	27.94
low		June	3.5	23.51	23.87
high		June	3.5	18.38	18.70

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845 **FIGURE CAPTIONS**

846 **Figure 1.** The study area is placed in the National Game Reserve “Ports de Tortosa i Besoit” (NGRPTB)
847 in Catalonia, northeast Spain (40°46’08” N, 0°20’04” E, 450 m. a. s. l.) marked in the upper inset. The
848 yellow line marks the Iberian Ibex enclosure. The white area indicates the flight area.

849 **Figure 2.** Field sampling in the Iberian Ibex enclosure in NGRPTB. Up to 20 individuals of the most
850 representative species of the area were marked with a dGPS. Insets present the species most
851 consumed by Iberian ibex in the enclosure.

852 **Figure 3.** A) Monthly variation of mean NDVI values recorded in the Iberian Ibex enclosure, in the
853 NGRPTB. Mean NDVI values corresponding to the 2014-18 period. Clear seasonal pattern evident with
854 peak primary production in winter and minimum in summer. The asterisk indicates statistically significant
855 differences between March and June NDVI (red bars). B) Variation of mean Blue-NDVI in the plants
856 recorded in the Ibex enclosure in March and June calculated from the images obtained by a BG-NIR
857 camera. All species have statistical differences (t-test) except for those marked with asterisks.
858 *Brachypodium retusum* (BR), *Helianthemum marifolium* (H), and *Pinus pinaster* (PP). All plant species
859 abbreviations are depicted in Table 2.

860 **Figure 4.** Diet composition assessed by a faecal micro-histological analysis of 10 faecal samples
861 collected in the Ibex enclosure in December 2019. Bars represent the mean of the proportion of each
862 plant species in our faecal samples. ONLW (Other Non-Legume Wood species), EM (*Erica multiflora*),
863 G (*Brachypodium phoenicoides*, *B. retusum* and other graminoids), ONLF (Other Non-Legume Forb
864 species), L-A (Labiatae-Asteraceae), QI (*Quercus ilex*), CA (*Cistus albidus*), TV (*Thymus vulgaris*), PL
865 (*Pistacia lentiscus*), HH (*Hedera helix*), CM (*Crataegus monogyna*), RU (*Rubus ulmifolius*), RO
866 (*Rosmarinus officinalis*), GS (*Genista scorpius*), Sasp (*Smilax aspera*), L (Laminaceae), Dsp
867 (*Dorycnium* sp.), A (Asteraceae), BS (*Buxus sempervirens*), I-O (Iridaceae-Orchidaceae), PA (*Phillyrea*
868 *angustifolia*), Psp (*Pinus* sp.), Csp (*Carex* sp.), Ssp (*Satureja* sp.), Rsp (*Rosa* sp.).

869 **Figure 5.** Mean scores from the first PCA dimension performed with NIR, Green and Blue band
870 recordings on 19 plant species sampled in the NGRPTB vegetation study. All plant species
871 abbreviations are depicted in Table 2.

872 **Figure 6.** Mean scores from the first PCA dimension performed with NIR, Green and Blue band
873 recordings on 5 plant categories consumed by Iberian ibexes in the NGRPTB. ‘C’: *Cistus albidus*; ‘G’:
874 *Brachypodium phoenicoides*, *B. retusum*, and other grass-like plants as Graminoids. ‘L’: *Rosmarinus*
875 *officinalis* and *Thymus vulgaris* as the family Labiatae; ‘E’: *Erica multiflora*; ‘F’: *Quercus* spp. as the
876 family Fagaceae.

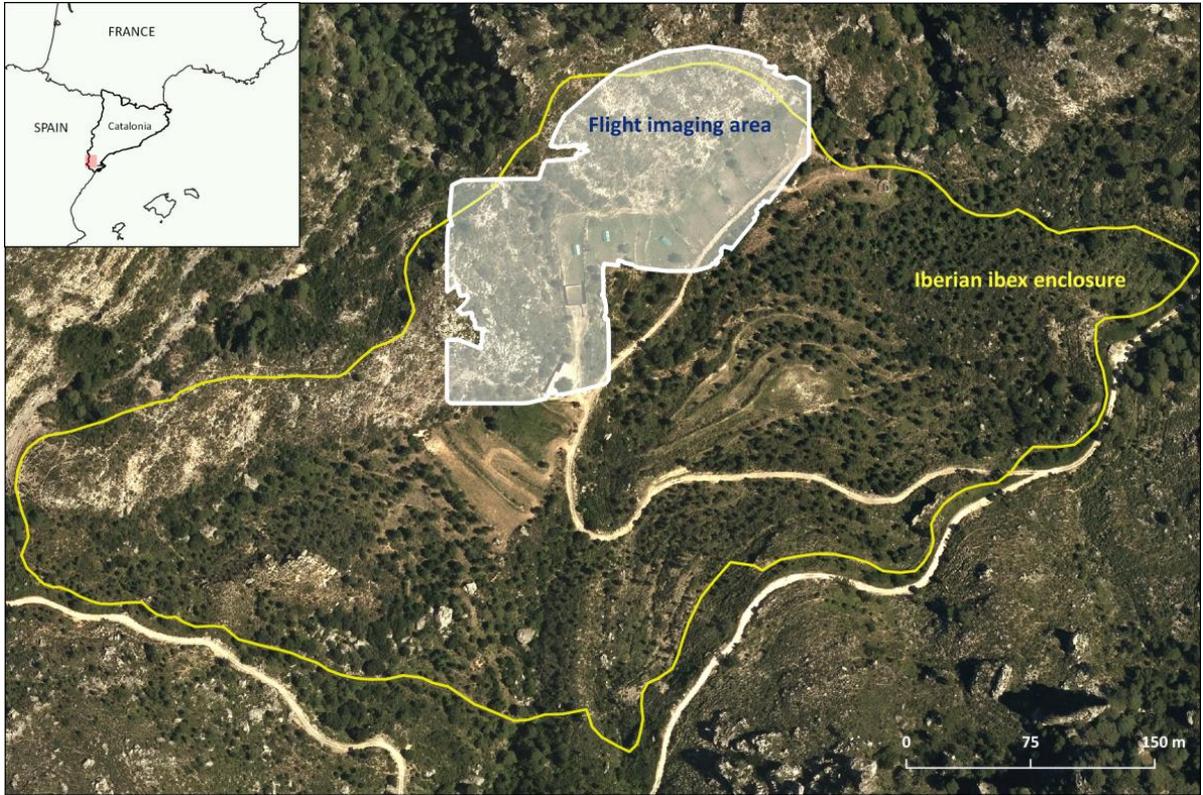
877 **Figure 7.** Foodscape map overlaid with the infrared false-colour orthomosaic of the study area.
878 Predicted categories are based on RFMI with an OOB error of 18.4%. A pie chart illustrates the
879 proportion of the 6 dietary categories: ‘F’: *Quercus* spp. as the family Fagaceae; ‘L’: *Rosmarinus*
880 *officinalis* and *Thymus vulgaris* as the family Labiatae; ‘E’: *Erica multiflora*; ‘C’: *Cistus albidus*; ‘G’:
881 *Brachypodium phoenicoides*, *B. retusum*, and other grass-like plants as Graminoids. ‘Others’ category
882 includes plants without dietary interest, bare soil and rock.

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885 **FIGURES**

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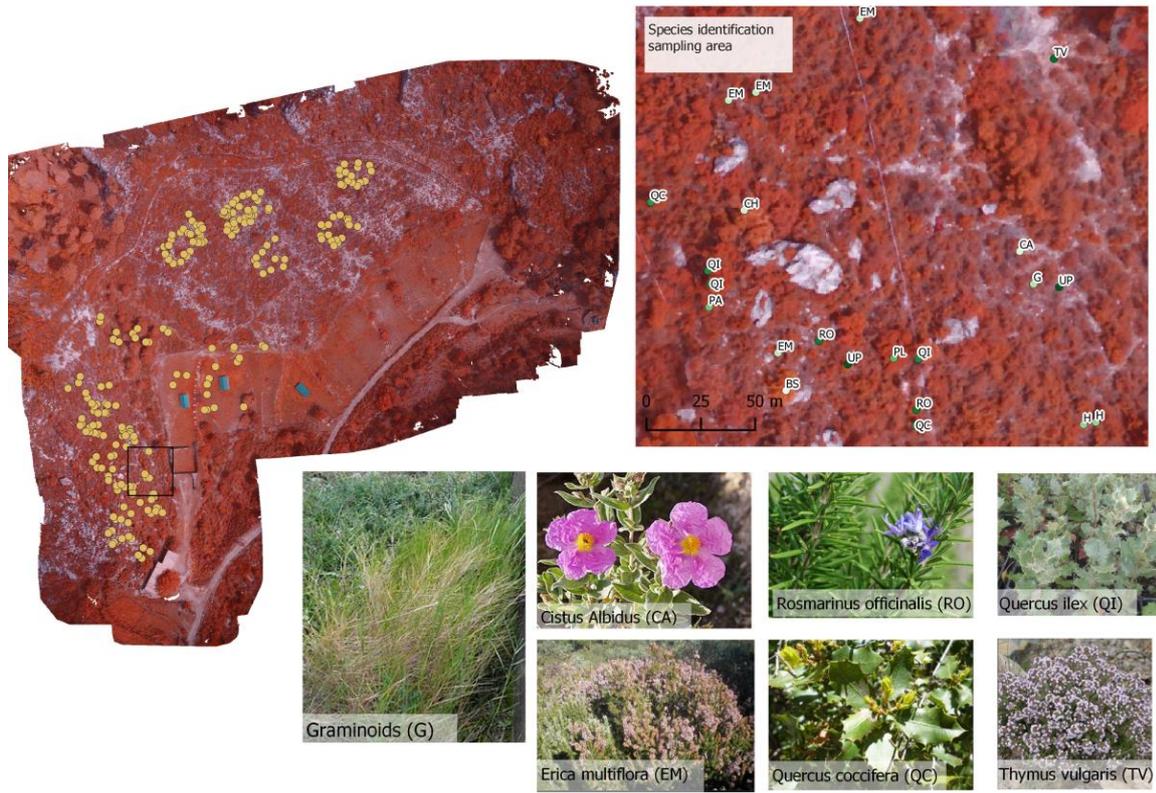
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888 **FIGURE 1**

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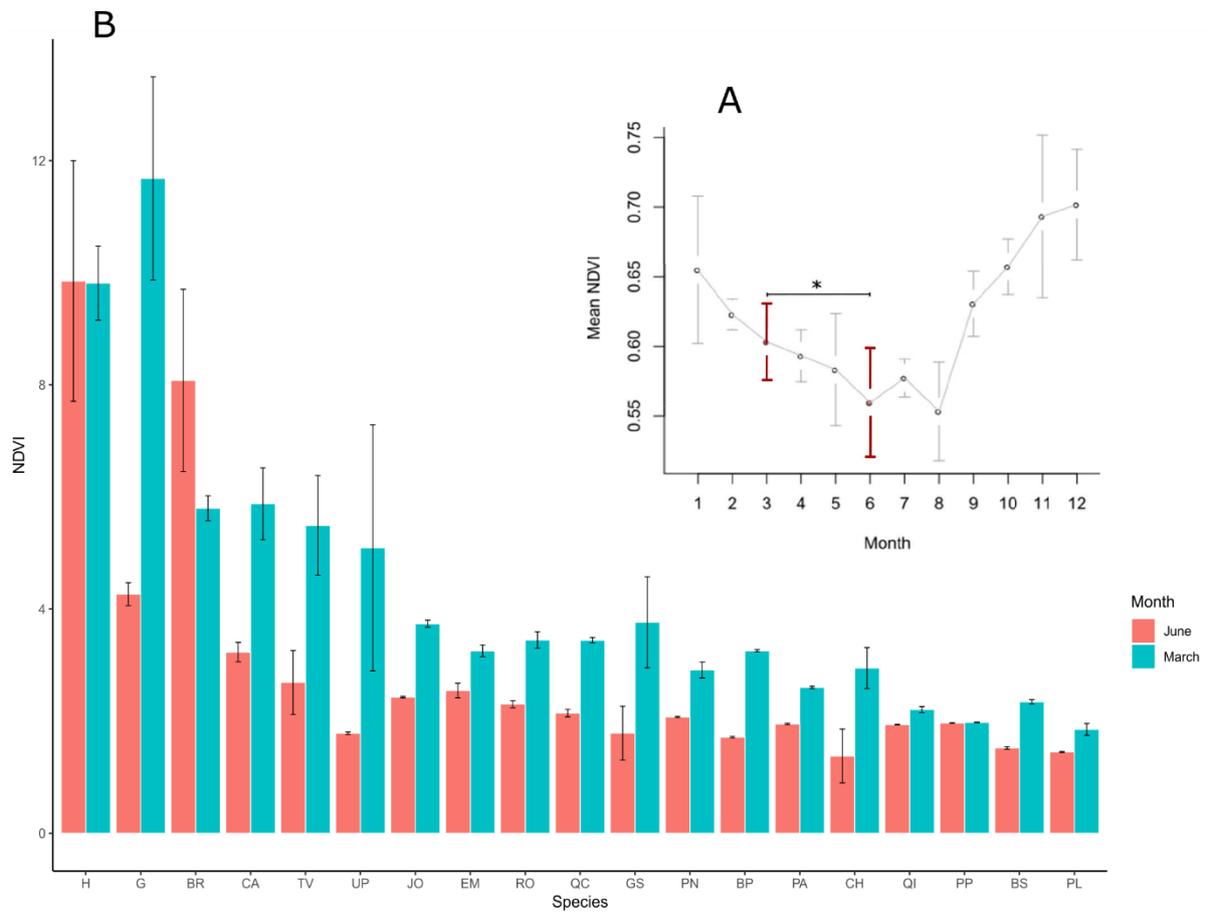
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FIGURE 2

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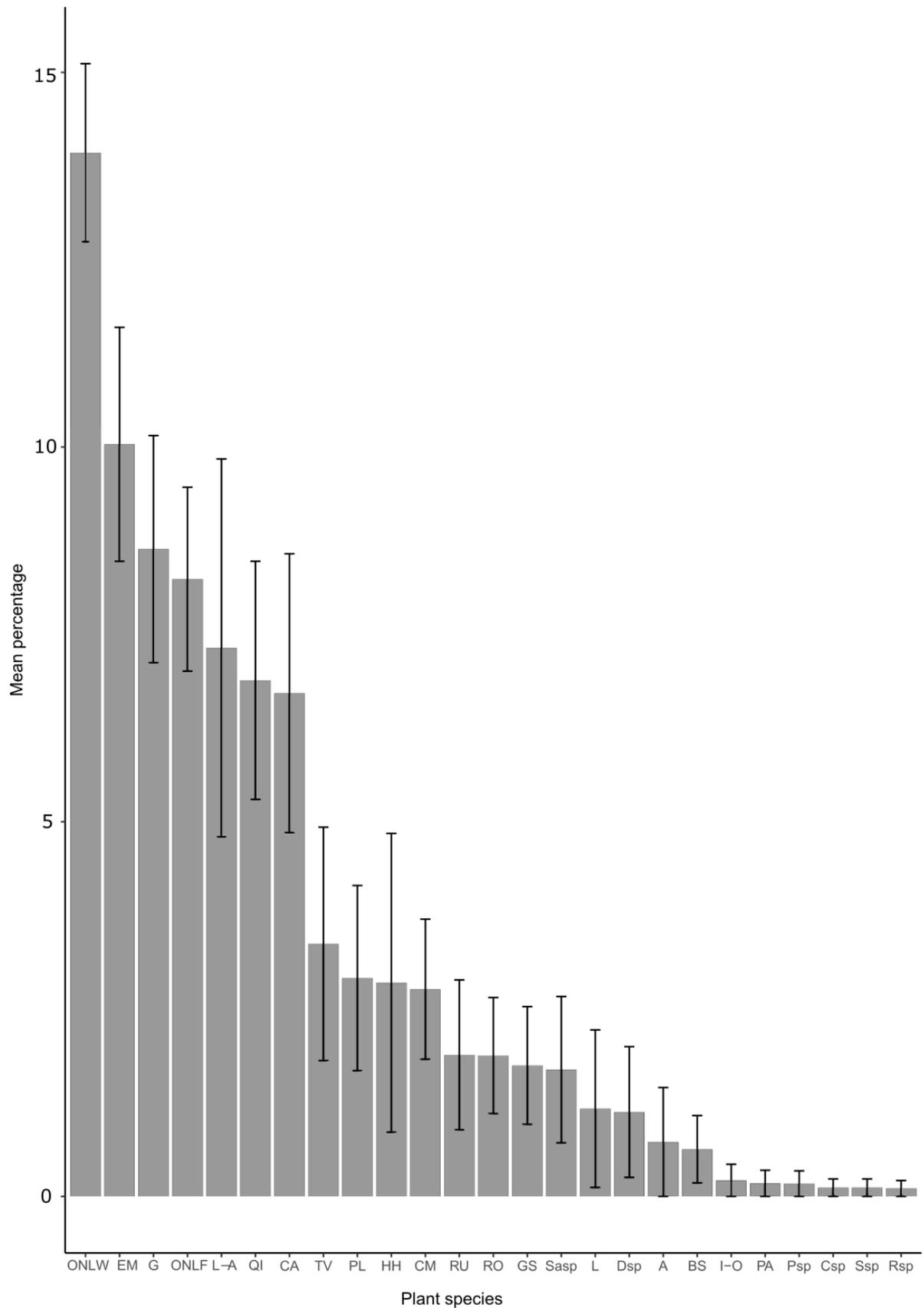
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FIGURE 3

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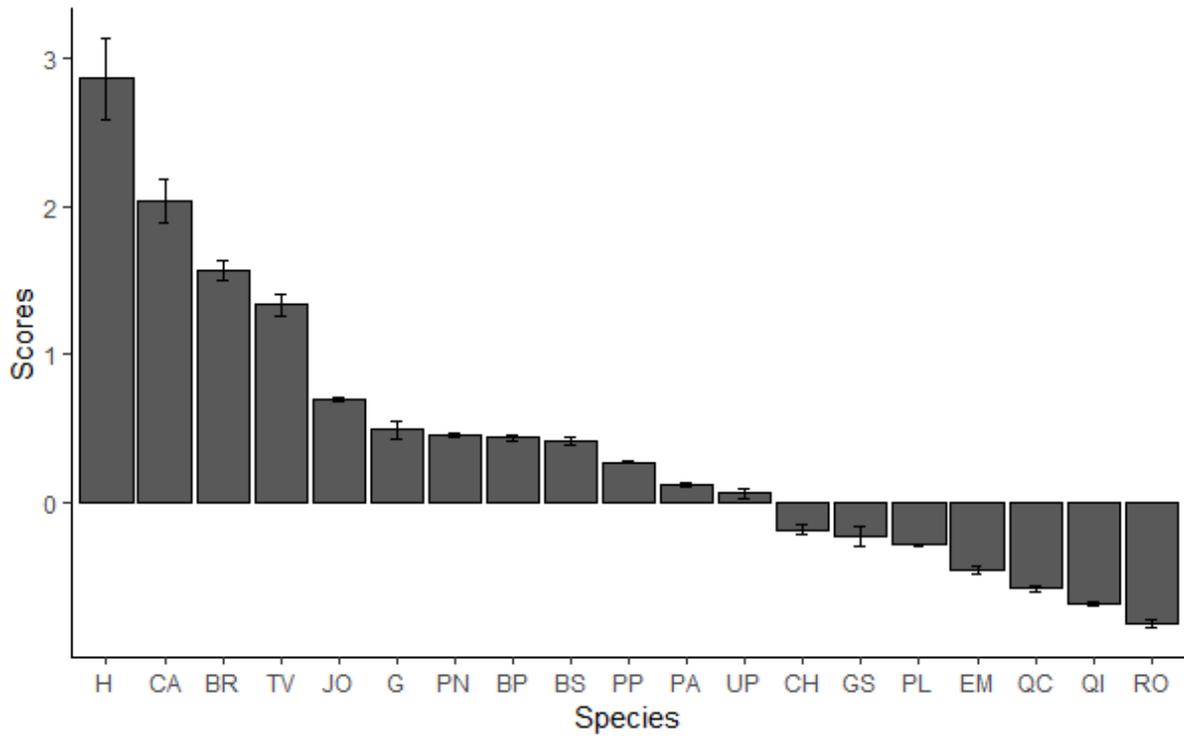
FIGURE 4

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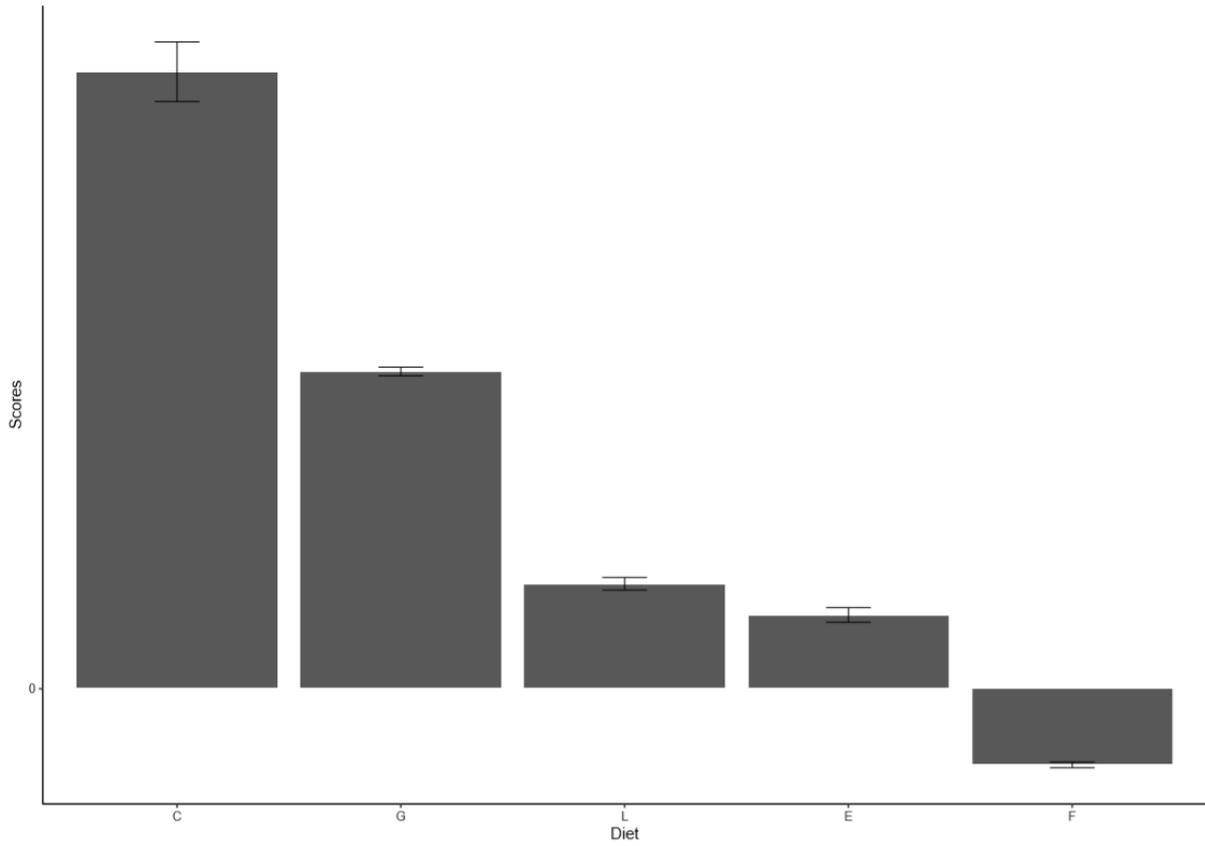
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FIGURE 5

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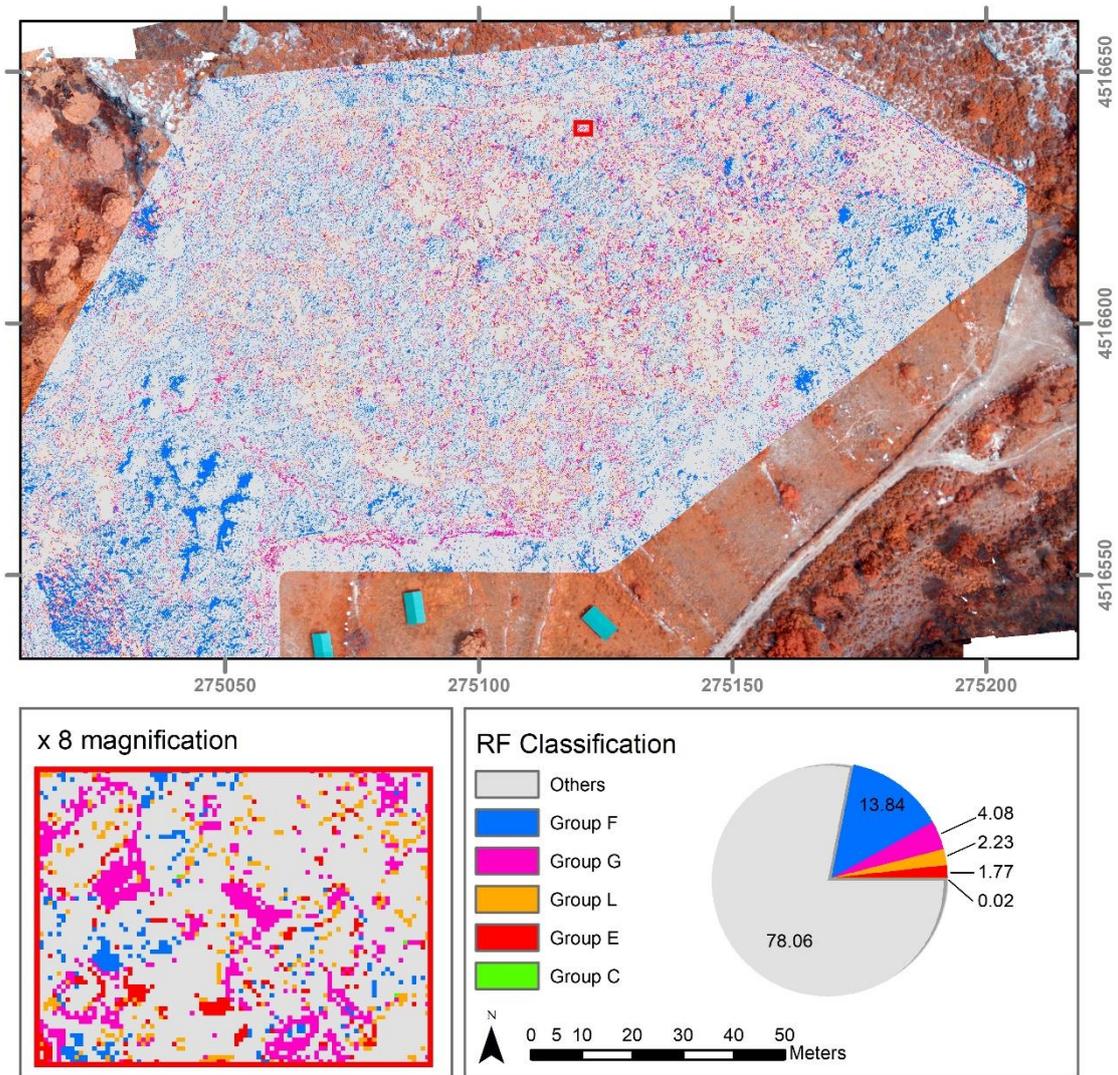


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FIGURE 6

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FIGURE 7