

1 **Quality attributes of cultivated white crowberries (*Corema album* (L.)**

2 **D. Don) from a multi-origin clonal field**

3 João Jacinto¹, Manuela Giovanetti^{1,2}, Pedro Brás Oliveira³, Teresa Valdiviesso³, Cristina

4 Máguas¹, Carla Alegria^{1*}

5 ¹cE3c - Centre for Ecology, Evolution and Environmental Changes, Faculdade de Ciências,
6 Universidade de Lisboa, 1749-016 Lisboa, Portugal.

7 ²CREA - Research Centre for Agriculture and Environment, Via di Saliceto 80, 40128 Bologna,
8 Italy.

9 ³INIAV - Instituto Nacional de Investigação Agrária e Veterinária, I.P., Unidade Estratégica de
10 Sistemas Agrários e Florestais e Sanidade Vegetal, 2780-157 Oeiras, Portugal.

11
12 ***Corresponding author**

13 Carla Alegria (csmalegria@gmail.com; csalegria@fc.ul.pt)

14 Current address: cE3c - Centre for Ecology, Evolution and Environmental Changes, Faculdade
15 de Ciências, Universidade de Lisboa. Campo Grande, Ed. C2, room 2.5.37, 1749-016 Lisboa,
16 Portugal; Tel.: + 351 217 500 000 ext. 22556.

17
18 **Author's ORCID:**

19 João Jacinto (0000-0002-7808-4238); Manuela Giovanetti (0000-0001-9442-0062); Teresa
20 Valdiviesso; (0000-0003-2832-0658); Cristina Máguas (0000-0002-4396-707); Carla Alegria
21 (0000-0002-9461-4569)

22
23 **Declarations**

24 **Funding**

25 Authors JJ, MG, CM and CA acknowledge financial support from Fundação para a Ciência e
26 Tecnologia (FCT), through the strategic project UIDB/00329/2020 granted to the Centre for

27 Ecology, Evolution and Environmental Changes, cE3c, Faculdade de Ciências, Universidade de
28 Lisboa. Author CA also acknowledges the financial support from FCT, through a postdoctoral
29 fellowship (SFRH/BPD/126703/2016). The white crowberry clonal field established at "Herdade
30 Experimental da Fataca" was funded by the Operational Group "CompetitiveSouthBerries"
31 (Partnership no. 21/Initiative no. 29/PDR2020-101-031721) which was co-financed by the
32 PDR2020, Portugal 2020 and the European Commission.

33 **Conflicts of interest**

34 The authors have no conflict of interest to report. The funders had no role in the design of the
35 study; in the collection, analyses, or interpretation of data; in the writing of the manuscript, or in
36 the decision to publish the results.

37 **Availability of data and material**

38 The data that support the findings of this study are available from the corresponding author, CA,
39 upon reasonable request.

40 **Acknowledgements**

41 Authors JJ, MG, CM and CA acknowledge financial support from Fundação para a Ciência e
42 Tecnologia (FCT), through the strategic project UIDB/00329/2020 granted to the Centre for
43 Ecology, Evolution and Environmental Changes, cE3c, Faculdade de Ciências, Universidade de
44 Lisboa. Author CA also acknowledges the financial support from FCT, through a postdoctoral
45 fellowship (SFRH/BPD/126703/2016). The white crowberry clonal field established at "Herdade
46 Experimental da Fataca" was funded by the Operational Group "CompetitiveSouthBerries"
47 (Partnership no. 21/Initiative no. 29/PDR2020-101-031721) which was co-financed by the
48 PDR2020, Portugal 2020 and the European Commission.

49 **Abstract**

50 There is a growing interest in *Corema album* (L.) D. Don fruits due to the unique white colour,
51 mildly acidic lemony flavour and health-promoting properties associated with its bioactive
52 composition. This study performs a physical-chemical characterisation of cultivated *C. album*
53 fruits from a multi-origin clonal field. The field comprises ten wild populations with distinct
54 geographical origins, grown under the same edaphoclimatic conditions. We analysed fruits
55 CIELab colour parameters, texture profile (TPA), pH, acidity (TA, g.100 mL⁻¹), soluble solids
56 content (SSC, %) and total phenolic content (TPC, mg CAE.100 g⁻¹). Our results showed
57 differences between fruits physical-chemical attributes. Variation patterns in fruits SSC and
58 hardness suggest that the differences might be related to the original geographical location of the
59 populations. The determined TPC levels in all samples were very encouraging at a bioactive level,
60 ranging from 185.3 to 355.6 mg CAE.100 g⁻¹. Fruits from Mira and Pego populations stood out
61 from the ten geographical provenances. Mira fruit samples had higher sweetness and lower
62 acidity, while the Pego ones had firmer fruits and higher phenolic content. The multi-origin clonal
63 field allowed us to offer an interesting scientific comparative background, highlighting the large
64 potential of these berries for introduction in the commercial market. Not only our results support
65 the potential of white crowberry as a new crop; the detected differences also indicate a hidden
66 capacity for small fruit market diversification.

67

68 **Keywords:** Cultivated plant populations; geographic provenance; fruit quality; hardness;
69 soluble solids content; phenolic content.

70

71 **1. Introduction**

72 *Corema album* (L.) D. Don, known as white crowberry, is an Iberian Peninsula endemic
73 species, from the Ericaceae family. The genus *Corema* has an amphi-Atlantic distribution with
74 only two known species: *Corema conradii* (Torr.) Torr. Ex Loud, in the eastern coast of North
75 America and *C. album*, with two subspecies, *C. album* spp. *azoricum* Pinto da Silva in Azores
76 and *C. album* spp. *album* in the Portuguese mainland and Spanish Atlantic coasts (Castroviejo et
77 al. 1993; Li et al. 2002). This evergreen shrub inhabits the coastal dune systems of the Atlantic
78 coast, or even in pine tree understory near the ocean and, in the Iberian Peninsula, it is distributed
79 from the North of Galicia to Gibraltar, in the south (Valdés et al. 1987; Álvarez-Cansino et al.
80 2012). An isolated population can also be found in Alicante, in the Mediterranean coast of Spain
81 (Martínez-Varea et al. 2019). The species develops blueberry-like fruits shaped in a drupe, with
82 a 5-8 mm diameter, usually with three seeds (Simmonds 1979). Fruit production ranges from July
83 to September, depending on geographical origin. When fully ripe, fruits develop a white or
84 pinkish-white colouration and, depending on genotypes, turn to translucent as maturation
85 progresses (Oliveira and Dale 2012). Still, reports on a winter fructification are known (Alegria
86 et al. 2020) and describe fruit maturation progression from white to black fruits, a newly reported
87 stage.

88 In both Portuguese and Spanish coastal areas, these berries are part of the traditional folk
89 culture, accounting for their consumption as fresh fruits at beaches and their commercial
90 exploitation in local markets, sold as fresh fruits, made into jams and liquors or even as traditional
91 medicine (Font-Quer and Davit 1993; Gil-López 2011; González 2006). Due to the recently up-
92 raised interest as a novel "fresh beach" fruit, driven by their colour and mildly acidic lemony
93 flavour, efforts are being made to convert this wild species into a new crop for future integration
94 in the berry market (Oliveira and Dale 2012). Moreover, a factor driving agronomic and market
95 possibilities are the *C. album* potential health benefits from its recognised antioxidant properties
96 (Pimpão et al. 2013), a trait evermore demanded by health-conscious consumers.

97 Existing reports on *C. album* biochemical properties mainly focus on its phenolic profile
98 and antioxidant capacity (Andrade et al. 2017a; León-González et al. 2012; León-González et al.
99 2013; Pimpão et al. 2013). Andrade et al. (2017b) and Alegria et al. (2020) also characterised the
100 physical-chemical properties of the white crowberry fruits and defined the maturation progression
101 of the berry in natural conditions. However, all these studies refer to *C. album* fruits collected
102 from wild specimens and, therefore, provide information regarding a single population. *C. album*
103 populations hold distinct genetic backgrounds which could significantly influence fruits physical-
104 chemical attributes (Jacinto et al. 2020), together with local edaphoclimatic conditions
105 (Åkerström et al. 2010; Rohloff et al. 2015). Moreover, Oliveira et al. (2020b) concluded that
106 within the same wild population, different genotypes gather distinct traits of interest, which
107 supports the establishment of a breeding program for the species. Considering the interest for
108 future cropping practices, there is an augmenting need to comparatively test different populations
109 grown in controlled conditions.

110 This study was designed to compare the physical-chemical properties and the total phenolic
111 content of ten (10) cultivated *C. album* populations established by rooted cuttings of wild plants
112 from different geographical origins, grown under the same edaphoclimatic conditions.

113 **2. Materials and Methods**

114 **2.1 Sampling**

115 Fruits from female plant individuals were collected from several genotypes, from ten (10)
116 different locations of the Portuguese coast (Figure 1), grown under the same conditions in
117 Herdade Experimental da Fataca, INIAV, I.P (37°34'56.8"N 8°44'23.6"W), on September 4,
118 2019. The experimental station is located in Southwest Alentejo and is characterised by an
119 average annual temperature of 17.1 °C and annual precipitation of 516 mm. The white crowberry
120 field was established in 2015, with plants obtained by vegetative propagation (rooted cuttings)
121 from wild plants collected in ten distinct geographical locations (Oliveira et al. 2020a). Plant
122 density is one meter along the line and three meters between lines, with one male plant separating
123 12 female plants along the line. Irrigation is achieved with drippers separated by 40 cm, watering

124 2 L.h⁻¹ twice a month for 30 min. Plants were able to produce significant amount of fruits after
125 three years in the field and its average volume is around 2 m³. Onwards, we adopt the term
126 "population" preceded by the original geographic location to identify the cultivated plants present
127 in Fataca.

128 Only white fruits (the mature stage) were randomly harvested from plants of each
129 provenance, collected in an average of 186±55 g (per replicate, n=3). Fruits were packed in
130 commercial vented clamshell containers (with snap-on lids) placed in a 38 L refrigerated
131 incubator and then transported to the laboratory. At the laboratory, fruits were screened and
132 defective fruits (crushed, cracked, or immature) eliminated. The selected fruits were placed in
133 vented clamshell containers (n=3 per population) for further analysis.

134 **2.2. Biometric measurements**

135 For the assessment of biometric characteristics, weight and calibre, all selected fruits from
136 each population were used. The calibre of each fruit sample (based on berries diameter) was sorted
137 with the aid of calibration sieves (Ø 10.25, 8.25, 7.5 mm) and fruits with <7.5 mm in diameter
138 discarded. Calibrated fruits were then counted on an automated seed counter and weighted on a
139 precision scale.

140 **2.3 Colour**

141 Berries superficial colour was evaluated with a CR 300 Minolta colourimeter (Osaka,
142 Japan) by measuring the CIELab parameters (C illuminant, 2nd observer). The instrument was
143 calibrated using a white tile standard (L* = 97.10; a* = 0.08; b* = 1.80). A total of 45
144 measurements were made per sample type (one measurement per fruit).

145 **2.4 Texture**

146 Uniform size fruit samples (n=15 fruits) were used for textural measurements. Prior to
147 analysis, samples were kept for 2 h at room temperature (20 °C) to prevent temperature influence
148 on fruits firmness (Chiabrande et al. 2009). Instrumental texture profile analysis (TPA) was
149 carried out on a TA-XT2i texture analyser (Stable Micro Systems, Godalming, UK) equipped
150 with a 30 kg load cell and HDP/90 platform. Samples were compressed to 30% of the original

151 height using a crosshead speed of $0.8 \text{ mm}\cdot\text{s}^{-1}$ and a 60 mm diameter cylinder stainless flat probe.
152 Each sample was subjected to a two-cycle compression with 5 s between cycles. Data was
153 collected using Exponent Version 6.1.4.0 software. The following parameters were calculated
154 from the resulting force–time curve: hardness (N); cohesiveness (adimensional); gumminess (N);
155 springiness (mm); chewiness (mJ) and resilience (adimensional).

156 **2.5 pH, Soluble solids content and Titrable acidity**

157 The pH and soluble solids content (SSC, %) of freshly prepared juice were determined
158 using a pH meter (Crison Micro pH 2001, Crison Instruments, Spain) and a digital refractometer
159 (DR-A1, ATAGO Co Ltd., Japan), respectively. Titrable acidity (TA) was determined by titrating
160 the freshly prepared juice with 0.1 N NaOH to an endpoint of pH 8.2 using a Mettler Toledo DL21
161 automatic titrator. Results were expressed as the mass equivalent (g) of citric acid per 100 mL of
162 juice ($\text{g}\cdot 100 \text{ mL}^{-1}$). The pH, SSC and TA determinations were carried out in 15 mL juice triplicates
163 for each sample type and the average values considered.

164 **2.6 Total phenolic content**

165 Samples ($n=3$ per population) were extracted with methanol (1:4, w:v) and the clear
166 supernatant used for the determination of the total phenolic content (TPC) using the Folin-
167 Ciocalteu reagent according to Heredia and Cisneros-Zevallos (2009). Results were expressed as
168 mg chlorogenic acid equivalents per 100 g of fresh tissue ($\text{mg CAE}\cdot 100 \text{ g}^{-1}$).

169 **2.7 Statistical Analyses**

170 R Studio was used to perform all statistical analyses (R Core Team 2013). To test the
171 differences in physical-chemical properties and total phenolic content among the 10 populations,
172 Kruskal-Wallis tests, at a significance level of $\alpha=0.05$, were performed, followed by Warden's
173 post hoc test ($\alpha=0.05$), for mean separation, with *agricolae* R package (De Mendiburu 2019).
174 Spearman's correlation ($\alpha=0.05$) was performed (Supporting Information Table S1), using the
175 *Hmisc* R package (Harrell 2014), to seek relations between studied variables and non auto-
176 correlated variables used to perform a Principal Component Analysis (PCA). The PCA was built
177 on eight of the studied variables, using *factorextra* R package (Kassambara and Mundt 2017).

178 **3. Results and discussion**

179 Among the studied populations, differences emerged on all physical-chemical properties
180 we addressed. These differences however grouped populations according to given parameters, as
181 explained in more details in the following paragraphs.

182 For biometrics, calibre showed that fruits with a diameter between 8.25 to 10.25 mm were
183 the most common among all samples (Supporting Information Figure S1). 47% to 71% of fruits
184 were within this calibre. In wild populations, similar fruit calibre (in the range of 8.25-10.25 mm)
185 has been reported (Andrade et al. 2017b; Jacinto et al. 2020; Larrinaga and Guitián 2016; Oliveira
186 and Dale 2012); however, in our study, we also found fruits with a diameter >10.25 mm in
187 samples collected from the Meco, Comporta and Cabo Sardão populations (from 36% up to 47%
188 of the total fruits). On the other hand, also fruits with smaller calibre (7.5-8.25 mm) were frequent,
189 especially in fruits collected from Mira and Quiaios populations (\approx 30% of the total fruits).
190 Considering a potential future use for fresh fruit production, fruits with higher calibres (>10.25
191 mm) potentially represent higher production yields and a more appealing marketability option.
192 Saftner et al. (2008) demonstrated that consumer preference on choosing blueberries from
193 different cultivars was mainly driven by fruit size perception, with larger fruits being preferred
194 over smaller ones, and related to high sensory textural scores (eating quality).

195 Regarding fruit weight, fruits from the highest calibre (>10.25 mm) ranged from 0.41 g in
196 VRSTAntónio to 0.71 g in Comporta. Average fruit weight from the most representative calibre
197 (8.25-10.25 mm) was between 0.32 g and 0.41 g, and similar fruit weights have been reported in
198 fruits collected from wild plants (Andrade et al. 2017b; Oliveira and Dale 2012; Oliveira et al.
199 2020b). Also, a study conducted in wild plants from Doñana, Spain, showed that plants with an
200 average canopy size of 0.96 m produce around 2200 fruits with an average weight of \approx 0.4 g
201 (Zunzunegui et al. 2006). Since the calibre range of 8.25-10.25-mm was the most common among
202 all evaluated samples, fruits from this calibre were selected for colour and texture assessments.

203 Fruits CIELab colour parameters are reported in table 1 and indicate significant differences
204 between fruit samples. *Corema album* is known for its white coloured berries. Thus, the

205 luminosity parameter L^* , ranging from 0 (pure black) to 100 (pure white), is well suited to
206 differentiate *C. album* fruits colour. Regarding samples L^* colour parameter (Table 1), we found
207 differences ($p < 0.05$) among fruits from different plant origins but all related to a white colour
208 perception ($L^* > 65$). We found most evident differences between fruit samples from Santo André
209 and Mira populations (both with L^* of ≈ 69 , $p > 0.05$), and VRSTAntónio, Cabo Sardão and
210 Quiaios populations (L^* ranging from 74.6 to 76.1, $p > 0.05$). In these latter samples, with higher
211 ($p < 0.05$) L^* values, fruit surface lightness was less influenced by variations in red (positive a^*)
212 chroma, being perceived as whiter fruits. Reports on the presence of low amounts of anthocyanins
213 are found in *C. album* fruits (León-González et al. 2013), which influences the white/pinkish-
214 white berry perception. Indeed, regarding the a^* parameter (Table 1) (sample redness), we found
215 significant differences, with fruit samples from Cabo Carvoeiro, Quiaios, Meco and Comporta
216 populations ($p > 0.05$) representing the lower a^* values and of Santo André, Moledo and Pego
217 ($p > 0.05$) populations representing the highest a^* values. Notwithstanding the found differences
218 between fruit samples, all samples had positive a^* values suggesting that all fruits tend to be, to
219 some extent, more pinkish/reddish than greenish (negative a^* values). The higher a^* values of
220 fruit samples from the Santo André population supports the lower numerical L^* values in regard
221 to those of, e.g., Quiaios, leading to a decreased white perception, probably associated with higher
222 amounts of anthocyanins (León-González et al. 2013). As for the b^* values (table 1), relating to
223 blue (negative values) and yellow (positive values) chromas, despite the found differences
224 ($p < 0.05$), all fruit samples had positive values ranging from 8.07 ± 2.02 (Quiaios) to 11.37 ± 1.98
225 (Pego). These variations in yellow chromas were more substantial than the ones found for red
226 chromas, leading to the assumption that these variations can also contribute to the overall white
227 colour perception of the fruits.

228 Among all fruits, those from the Mira population had the highest variability regarding
229 colour parameters: 68.80 ± 8.25 , 1.68 ± 3.02 , 11.23 ± 3.48 for L^* , a^* , b^* , respectively. Andrade et al.
230 (2017b) assessed wild plants from Mira and reported higher values for L^* (79.82 ± 2.82) and
231 lower values for a^* (1.27 ± 2.05) and b^* (5.88 ± 2.1). These differences could possibly be related
232 to the distinct edaphic-climatic conditions of growing sites, influencing pigment synthesis, as

233 documented in blueberry (Howell et al. 2001; Routray et al. 2011). Nevertheless, other
234 mechanisms apart from climatic context might influence fruit colour seeing as we also found
235 colour differences of similar range between fruit samples from the Fataca collection. Díaz-
236 Barradas et al. (2016) assessed *C. album* wild fruits reflectance spectra from plants of Donñana,
237 Spain, finding that berries reflectance is related mainly to two pentacyclic triterpenes, ursolic and
238 oleanolic acid. Thus, the found differences in colour parameters, particularly in L*values, might
239 be due to different amounts of triterpenes present in the berries.

240 The results of white crowberries' texture profile (TPA) are shown in Figure 2 and
241 Supporting Information Table S2. From the evaluated texture parameters, hardness was the
242 parameter that best represented textural differences in white crowberry fruits (Figure 2). Fruit's
243 hardness varied from 3.9 ± 0.8 N (VRSTAntónio) to 7.7 ± 2.2 N (Pego). Moreover, from Figure 2,
244 it is possible to observe an interesting pattern regarding fruit samples hardness, describing a
245 visible bell-shaped pattern related to the geographical origin of the Fataca populations. Fruit
246 samples from Comporta, Pego and Santo André populations have significantly higher hardness
247 values than remaining populations. This pattern suggests that fruits from these populations,
248 originally located in the northern shores of the Alentejo Litoral region (Figure 1), have a
249 significantly different textural imprint regarding the other populations, originally located to the
250 north and south of this cluster. This behaviour might be linked to specific functional traits
251 contingent on a "memory effect", most likely genetic, related to the particular "in natura"
252 geographical origins/environmental conditions.

253 In literature, textural properties of *C. album* fruits are only described in reference to wild
254 fruits from Mira (Andrade et al. 2017b), reporting values of ca. 1.9 N for hardness. Despite
255 differences in texture determination methodology, the reported values are much lower than the
256 ones determined in the cultivated fruits collected from the Fataca population (Mira; 5.0 ± 1.9 N).
257 As previously mentioned, we should not rule out the differences in environmental conditions from
258 each location (Mira and Fataca's). For instance, Lobos et al. (2018) assessed different irrigation
259 conditions in blueberry (cv. Brigitta) plants and demonstrated that plants under deficit irrigation
260 had firmer fruits. Although plants in Fataca were sparsely irrigated, the drier environmental

261 conditions in the site might have influenced fruits textural attributes, leading to firmer fruits.
262 Moreover, in blueberries, Ochmian et al. (2009) reported that soil composition also has a
263 significant effect on fruit quality, including firmness, which can similarly contribute to explain
264 the found differences between studies.

265 Mean values (\pm SD) of fruit samples soluble solids content (SSC), pH, titrable acidity (TA)
266 are shown in Table 2. The distinctive taste found in *C. album*, sweet-sour or acidic taste, makes
267 sugar concentration and pH important parameters for assessing fruits quality. SSC expresses an
268 approximate measure of the amount of sucrose (g) per 100 g of solution. We found significant
269 differences ($p < 0.05$) regarding fruit samples SSC, ranging from $8.2 \pm 0.2\%$ (VRSTAntônio) to
270 $10.6 \pm 0.1\%$ (Mira), higher than the ones reported in other works in wild *C. album* fruits (Alegria
271 et al. 2020; Andrade et al. 2017b; Pimpão et al 2013). We found a decreasing trend regarding fruit
272 samples SSC, which can possibly be related to the original geographical location of the
273 populations. The decrease tendency is compliant to the north-south positioning of the wild *C.*
274 *album* populations from which Fataca's clonal field was established. Again, since the edaphic-
275 climatic context is the same for all sampled plants/fruits, the found differences might be related
276 to adaptation strategies of the wild populations to local factors and specific climate conditions
277 which are "passed on" through a form of "genetic memory". This "memory effect" could,
278 therefore, influence plants functional traits and, consequently, fruit quality (in this case, sugar
279 content).

280 All fruit samples had low pH values (Table 2), ranging from 2.6 to 3.2 pH units, which is
281 similar to the reported by Alegria et al. (2020) and to the ranges reported for different blueberry
282 cultivars (Chiabrande et al. 2009; Giovanelli and Buratti 2009; Liu et al. 2019). Even though we
283 found statistical differences between fruit samples, differences were, at most, of 0.5 pH units,
284 which does not relate to any expressive physiological outcome. Nevertheless, the low pH values
285 label *C. album* fruits as acidic and promote microbial development inhibition, therefore
286 contributing to fruit preservation. Titrable acidity (TA, Table 2) showed to be coincident with
287 sample pH, with low TA corresponding to high pH and vice-versa. White crowberries are
288 described as high acidity fruits (Andrade et al. 2017b; Pimpão et al. 2013). Pimpão et al. (2013)

289 alluded that such high acidity might be a concerning issue for fresh consumption. However, it
290 also creates an opportunity window for other commercial valorisation strategies, namely as a food
291 additive, as suggested by Alegria et al. (2020).

292 Fruit sample total phenolic content (TPC) was determined, and results shown in Figure 3.
293 Reports on high contents of phenolic compounds in wild white crowberry fruits are closely related
294 to the fruits antioxidant properties (Andrade et al. 2017a; León-González et al. 2012; León-
295 González et al. 2013; Pimpão et al. 2013). In our study, fruit sample TPC levels ranged from
296 185.3 to 355.6 mg CAE.100 g⁻¹. Among evaluated fruit samples, the fruits from the Pego
297 population stand out with the highest TPC levels, 1.5 times higher than the average TPC values
298 for all samples. Nonetheless, the determined TPC levels ascribe to *C. album* fruits a high
299 antioxidant potential, irrespective of populations geographical origin.

300 As previously mentioned, several studies reported high contents of phenolics in *C. album*
301 fruits: 12 mg GAE/g (dw) (Pimpão et al. 2013); 1214.4±122 mg GAE/kg (fw) and 7316.6±740
302 mg GAE/kg (dw) (León-González et al. 2013); 1997±75 mg GAE/100 g (Andrade et al. 2017a)
303 and 1393.91±0.06 mg/100 g characterised in a water extract (León-González et al. 2012). These
304 studies agree on the high antioxidant potential of the *C. album* fruits, attributed to the phenolic
305 composition which supports our results. The high phenolic content has been related particularly
306 to the high amounts of phenolic acids, with benzoic and hydrocinnamic acids, especially
307 chlorogenic acid, reported as the most abundant phenolic. Considering that phenolic acids are the
308 main group of phenolics found in *C. album* fruits, it is also possible that this prevalent composition
309 influences the acidic taste perception (Tomás-Barberán and Espín 2001).

310 We used a Principal Component Analysis (PCA) to explore which physical-chemical traits
311 best describe the differences among fruits of cultivated *C. album* populations. We used eight non
312 auto-correlated variables (Supporting Information Table S1) for the PCA: the L*, a*, b* colour
313 parameters, hardness and the pH, SSC, TA and TPC parameters. The obtained PCA accounted
314 for 79.5% of the total variance on the first two axes (53.4% and 26.2%, in PC1 and PC2,
315 respectively). The original data variability explained in the first two dimensions is considered
316 suitable to define a good qualitative model as a significant percentage of the original information

317 (>70%) accumulates within the first two PC's (Larrigaudière et al. 2004). The first axis (PC1)
318 was most heavily loaded by pH, SSC, TA and two colour parameters (L^* and a^*), while the
319 second axis (PC2) was heavily loaded by hardness and total phenolic content (Supporting
320 Information Table S3). The PCA confirms a major cluster, grouping eight of the ten fruit samples,
321 with samples from the Pego and Mira populations independently segregated. The segregation of
322 the Mira population, established by PC1, relates to the pH and the SSC ($r>0.80$) and with L^* , a^*
323 and TA ($r<-0.8$), distinguishing fruits with sweeter traits and pinkish-white colour perception. On
324 the other hand, the segregation of the Pego sample relied mostly on the TPC and hardness vectors,
325 both positively correlated with PC2 ($r>0.85$), indicating that this sample is distinguished by its
326 high phenolic levels and firmer fruits.

327 **4. Conclusions**

328 This study contributes to the valorisation of *Corema album* and is the first focused on the quality
329 evaluation of cultivated white crowberry fruits from multiple origins. On the base of specific fruit
330 quality attributes from which this species might be desirable (e.g. acidity, SSC), and despite minor
331 differences, most fruits were similar. From the ten different geographical origins studied, grown
332 under the same conditions in Fataca, only fruits from Mira and Pego populations were clearly
333 segregated. The dissociation was based on fruit sweetness (SSC), firmness, and phenolic content.
334 These specific quality attributes might be linked to specific functional traits conditional to a
335 "memory effect", most likely genetic, related to adaptation strategies of the wild populations to
336 local factors and specific climate conditions. In this context, it will be important to understand
337 how different cultivation conditions (emulating the natural habitat) affect fruit quality and to
338 select appropriate genotypes for the viability of a crop with the desirable quality characteristics.
339 The white crowberry has an interesting physical-chemical profile and high phenolic content,
340 supporting its evaluation as a new crop for a potential small fruit market expansion.

341

342 **References**

343 Åkerström A, Jaakola L, Bång U, Jaderlund A (2010) Effects of latitude-related factors and
344 geographical origin on anthocyanidin concentrations in fruits of *Vaccinium myrtillus* L.
345 (bilberries). J. Agric. Food Chem. 58(22):11939-11945. Doi: 10.1021/jf102407n

346 Alegria C, Abreu M, Máguas C, Giovanetti M (2020) Winter Collection of the Underutilized
347 Berry *Corema album* (L.): New Insights on its Maturation Progression. Agri Res & Tech:
348 Open Access J. 24(4):556274. Doi: 10.19080/ARTOAJ.2020.24.556274

349 Álvarez-Cansino L, Díaz-Barradas MC, Zunzunegui M, Esquivias MP, Dawson TE (2012)
350 Gender-specific variation in physiology in the dioecious shrub *Corema album* throughout its
351 distributional range. Funct. Plant Biol. 39(12):968-978. Doi: 10.1071/FP12131

352 Andrade SC, Guiné RP, Gonçalves FJ (2017a) Evaluation of phenolic compounds, antioxidant
353 activity and bioaccessibility in white crowberry (*Corema album*). J Food Meas. Charact.
354 2017a; 11(4):1936-1946. Doi: 10.1007/s11694-017-9576-4

355 Andrade SC, Gonçalves F, Guiné R (2017b) Contribution for the physical-chemical
356 characterization of Portuguese Crowberry (*Corema album*). Int J Food Sci Nutr. 2(4):9-14.

357 Castroviejo S, Aedo C, Gómez Campo C, Laínz M, Montserrat P, Morales R, Muñoz Garmendia
358 F, Nieto Felinier G, Rico E, Talavera S, Villar L (1993). Flora Ibérica. Plantas vasculares de
359 la Península Ibérica e Islas Baleares.- Cruciferae-Monotropaceae Vol. IV. Real Jardim
360 Botánico, C.S.I.C. Madrid.

361 Chiabrando V, Giacalone G, Rolle L (2009) Mechanical behaviour and quality traits of highbush
362 blueberry during postharvest storage. J Sci Food Agric. 89:989–992. Doi: 10.1002/jsfa.3544

363 De Mendiburu F (2019) Agricolae: statistical procedures for agricultural research. R package
364 version. 1.3-1.

365 Diaz-Barradas MC, Costa C, Correia O, León-González AJ, Navarro-Zafra I, Zunzunegui M,
366 Álvarez-Cansino L, Martín-Cordero C (2016). Pentacyclic triterpenes responsible for
367 photoprotection of *Corema album* (L.) D. Don white berries. *Biochemical Systematics and*
368 *Ecology*, 67, 103-109. Doi: 10.1016/j.bse.2016.05.009

369 Font-Quer P, Davit S (1993) Plantas medicinales: el Dioscórides renovado. Labor.

370 Gil-López MJ (2011) Etnobotánica de la camarina (*Corema album*, Empetraceae) en Cádiz. Acta
371 Botanica Malacitana. 36:137-144. Doi: 10.24310/abm.v36i1.2784

372 Giovanelli G, Buratti S (2009) Comparison of polyphenolic composition and antioxidant activity
373 of wild Italian blueberries and some cultivated varieties. Food Chem. 112(4):903-908. Doi:
374 10.1016/j.foodchem.2008.06.066

375 Gonzáles G (2006) Los árboles y arbustos de la Península Ibérica e Islas Baleares. Madrid. Esp,
376 editorial SA, 2ed; p. 1727.

377 Harrell Jr FE (2014) Hmisc package version 4.1-1; 2014.

378 Heredia JB, Cisneros-Zevallos L (2009) The effect of exogenous ethylene and methyl jasmonate
379 on pal activity, phenolic profiles and antioxidant capacity of carrots (*Daucus carota*) under
380 different wounding intensities. Postharvest Biol. Technol. 51(2):242–249. Doi:
381 10.1016/j.postharvbio.2008.07.001

382 Howell A, Kalt W, Duy JC, Forney CF, McDonald JE (2001) Horticultural factors affecting
383 antioxidant capacity of blueberries and other small fruit. Horttechnology. 11(4):523-528. Doi:
384 10.21273/HORTTECH.11.4.523

385 Jacinto J, Oliveira PB, Valdiviesso T, Capelo J, Arsénio P, Nóbrega F (2020) Genetic diversity
386 assessment among *Corema album* (L.) D. Don (Ericaceae) genotypes based on ISSR markers
387 and agro-morphological traits. Genet. Resour. Crop Evol. 67(3):715-726. Doi:
388 10.1007/s10722-019-00849-8

389 Kassambara A, Mundt F (2017) Factoextra: extract and visualize the results of multivariate data
390 analyses. R package version 1.0. 4.

391 Larrigaudière C, Lentheric I, Puy J, Pintó E (2004) Biochemical characterisation of core browning
392 and brown heart disorder in pear by multivariate analysis. Postharvest Biol. Technol. 31:29–
393 39. Doi: 10.1016/S0925-5214(03)00132-7

394 Larrinaga AR, Guitián P (2016) Intraspecific variation in fruit size and shape in *Corema album*
395 (Ericaceae) along a latitudinal gradient: from fruits to populations. Biol. J. Linn. Soc. Lond.
396 118(4):940-950. Doi: 10.1111/bij.12794

397 León-González AJ, Mateos R, Ramos S, Martín MÁ, Sarriá B, Martín-Cordero C, et al. (2012)
398 Chemo-protective activity and characterization of phenolic extracts from *Corema album*. Food
399 Res. Int. 49(2):728-738. Doi: 10.1016/j.foodres.2012.09.016

400 León-González AJ, Truchado P, Tomás-Barberán FA, López-Lázaro M, Díaz-Barradas MC,
401 Martín-Cordero C (2013) Phenolic acids, flavonols and anthocyanins in *Corema album* (L.)
402 D. Don berries. J Food Compost Anal. 29(1):58-63. Doi: 10.1016/j.jfca.2012.10.003

403 Li J, Alexander Iii J, Ward T, Del Tredici P, Nicholson R (2002) Phylogenetic relationships of
404 Empetraceae inferred from sequences of chloroplast gene matK and nuclear ribosomal DNA
405 ITS region. Mol. Phylogenet. Evol. 25(2):306-315. Doi: 10.1016/S1055-7903(02)00241-5

406 Liu B, Wang K, Shu X, Liang J, Fan X, Sun L (2019) Changes in fruit firmness, quality traits and
407 cell wall constituents of two highbush blueberries (*Vaccinium corymbosum* L.) during
408 postharvest cold storage. Sci. Hortic. 246:557-562. Doi: 10.1016/j.scienta.2018.11.042

409 Lobos TE, Retamales JB, Ortega-Farías S, Hanson EJ, López-Olivari R, Mora ML (2018)
410 Regulated deficit irrigation effects on physiological parameters, yield, fruit quality and
411 antioxidants of *Vaccinium corymbosum* plants cv. Brigitta. Irrig Sci. 36(1):49-60. Doi:
412 10.1007/s00271-017-0564-6

413 Martínez-Varea CM, Ferrer-Gallego PP, Raigón MD, Badal E, Ferrando-Pardo I, Laguna E,
414 Villaverde V (2019) *Corema album* archaeobotanical remains in western Mediterranean basin.
415 Assessing fruit consumption during Upper Palaeolithic in Cova de les Cendres (Alicante,
416 Spain). Quaternary Science Reviews, 207, 1-12. Doi: 10.1016/j.quascirev.2019.01.004

417 Ochmian I, Grajkowski J, Skupień K (2009) Influence of substrate on yield and chemical
418 composition of highbush blueberry fruit cv. 'Sierra'. J Fruit Ornam Plant Res. 17(1):89-100.

419 Oliveira PB, Dale A (2012) *Corema album* (L.) D. Don, the white crowberry - a new crop. J Berry
420 Res. 2(3):123-133. Doi: 10.3233/JBR-2012-033

421 Oliveira PB, Luz FR, Magalhães T, Lisboa A, Oliveira CM, Valdivieso T (2020a) Propagação
422 vegetativa e seminal em *Corema album* (L.) D. Don. Actas Portuguesas de Horticultura.
423 30:347-356.

424 Oliveira PB, Valdivieso T, Luz FR (2020b) Melhoramento Genético da camarinha; Seleção e
425 Avaliação de plantas. Actas Portuguesas de Horticultura. 30:338-346.

426 Pimpão RC, Dew T, Oliveira PB, Williamson G, Ferreira RB, Santos CN (2013) Analysis of
427 phenolic compounds in Portuguese wild and commercial berries after multienzyme
428 hydrolysis. J. Agric. Food Chem. 61(17):4053-4062. Doi: 10.1021/jf305498j

429 R Core Team. R: A language and environment for statistical computing (2013) R foundation for
430 statistical computing, Vienna, Austria. ISBN 3-900051-07-0. <http://www.Rproject.org/>

431 Rohloff J, Uleberg E, Nes A, Krogstad T, Nestby R, Martinussen I (2015) Nutritional composition
432 of bilberries (*Vaccinium myrtillus* L.) from forest fields in Norway—Effects of geographic
433 origin, climate, fertilization and soil properties. J Appl Bot Food Qual. 88:274-287. doi:
434 10.5073/JABFQ.2015.088.040

435 Routray W, Orsat V (2011) Blueberries and their anthocyanins: factors affecting biosynthesis and
436 properties. Compr. Rev. Food Sci. Food Saf. 10(6):303-320. Doi: 10.1111/j.1541-
437 4337.2011.00164.x

438 Saftner R, Polashock J, Ehlenfeldt M, Vinyard B (2008) Instrumental and sensory quality
439 characteristics of blueberry fruit from twelve cultivars. Postharvest Biol. Technol. 49(1):19-
440 26. Doi: 10.1016/j.postharvbio.2008.01.008

441 Simmonds NW (1979) Principles of crop improvement, Longman. p. 408.

442 Tomás-Barberán FA, Espín JC (2001) Phenolic compounds and related enzymes as determinants
443 of quality in fruits and vegetables. J Sci Food Agric. 81(9):853–876. Doi: 10.1002/jsfa.885

- 444 Valdés B, Talavera S, Fernández Galiano E (1987) Flora vascular de Andalucía Occidental.
445 Barcelona: Ketres.
- 446 Zunzunegui M, Díaz-Barradas MC, Clavijo A, Álvarez-Cansino L, Lhout FA, Novo FG (2006).
447 Ecophysiology, growth timing and reproductive effort of three sexual forms of *Corema album*
448 (Empetraceae). Plant Ecology, 183(1), 35-46. Doi: 10.1007/s11258-005-9004-4

449 **Tables**

450 **Table 1.** CIELab colour parameters of white crowberry fruit samples from Fataca's clonal field,
 451 established with plants from ten distinct geographical origins.

Sample ID	L*	a*	b*
Moledo	71.75 ^{ef} ± 3.00	0.99 ^{ab} ± 0.82	8.51 ^{bc} ± 1.72
Mira	68.80 ^{fg} ± 8.25	1.68 ^{bcd} ± 3.02	11.23 ^a ± 3.48
Quiaios	75.62 ^{ab} ± 4.38	0.24 ^{ef} ± 0.67	8.07 ^c ± 2.02
Cabo Carvoeiro	72.50 ^{de} ± 3.89	0.33 ^f ± 0.64	8.99 ^b ± 1.66
Meco	74.20 ^{bcd} ± 4.57	0.17 ^{def} ± 0.50	8.65 ^{bc} ± 1.49
Comporta	73.00 ^{cde} ± 4.48	0.37 ^{cdef} ± 0.74	8.36 ^{bc} ± 1.84
Pego	72.77 ^{de} ± 4.35	0.66 ^{abc} ± 0.70	11.37 ^a ± 1.98
Santo André	68.57 ^g ± 4.53	1.12 ^a ± 1.08	8.27 ^{bc} ± 1.66
Cabo Sardão	74.66 ^{abc} ± 6.11	0.48 ^{cdef} ± 0.89	8.23 ^c ± 2.11
VRSTAntónio	76.13 ^a ± 4.56	0.48 ^{cde} ± 0.56	8.61 ^{bc} ± 1.85

452 Within a column, different letters represent significant differences at p=0.05 (Warden's post hoc test). L*
 453 values represent the luminosity of samples (0-black to 100-white), a* and b* values indicate the variation
 454 of greenness to redness (-60 to + 60) and blueness to yellowness (-60 to + 60), respectively.

455 **Table 2.** Quality parameters of pH, soluble solids content (SSC) and titrable acidity (TA) of white
 456 crowsberry fruit samples from Fataca's clonal field, established plants from ten distinct
 457 geographical origins.

Sample ID	pH	SSC (%)	TA (g.100 ml⁻¹)
Moledo	2.98 ^b ± 0.02	9.17 ^c ± 0.12	10.85 ^g ± 0.03
Mira	3.24 ^a ± 0.03	10.57 ^a ± 0.06	6.77 ⁱ ± 0.07
Quiaios	2.89 ^{bc} ± 0.04	9.10 ^c ± 0.10	14.19 ^c ± 0.11
Cabo Carvoeiro	2.69 ^f ± 0.02	9.37 ^b ± 0.06	11.75 ^e ± 0.10
Meco	2.62 ^g ± 0.02	8.83 ^d ± 0.15	14.49 ^b ± 0.12
Comporta	2.79 ^{de} ± 0.01	9.13 ^c ± 0.15	9.95 ^h ± 0.07
Pego	2.80 ^{de} ± 0.05	9.03 ^{cd} ± 0.06	12.62 ^d ± 0.07
Santo André	2.84 ^{cd} ± 0.01	8.43 ^e ± 0.06	11.93 ^e ± 0.13
Cabo Sardão	2.76 ^e ± 0.02	8.47 ^e ± 0.15	11.02 ^f ± 0.07
VRSTAntónio	2.67 ^f ± 0.02	8.17 ^f ± 0.15	16.53 ^a ± 0.13

458 Within a column, different letters represent significant differences at p=0.05 (Warden's post hoc test).

459

460 **Figure captions**

461 **Figure 1.** Location of Herdade Experimental da Fataca (INIAV), and identification of the plant's
462 collection sites for the clonal field establishment.

463

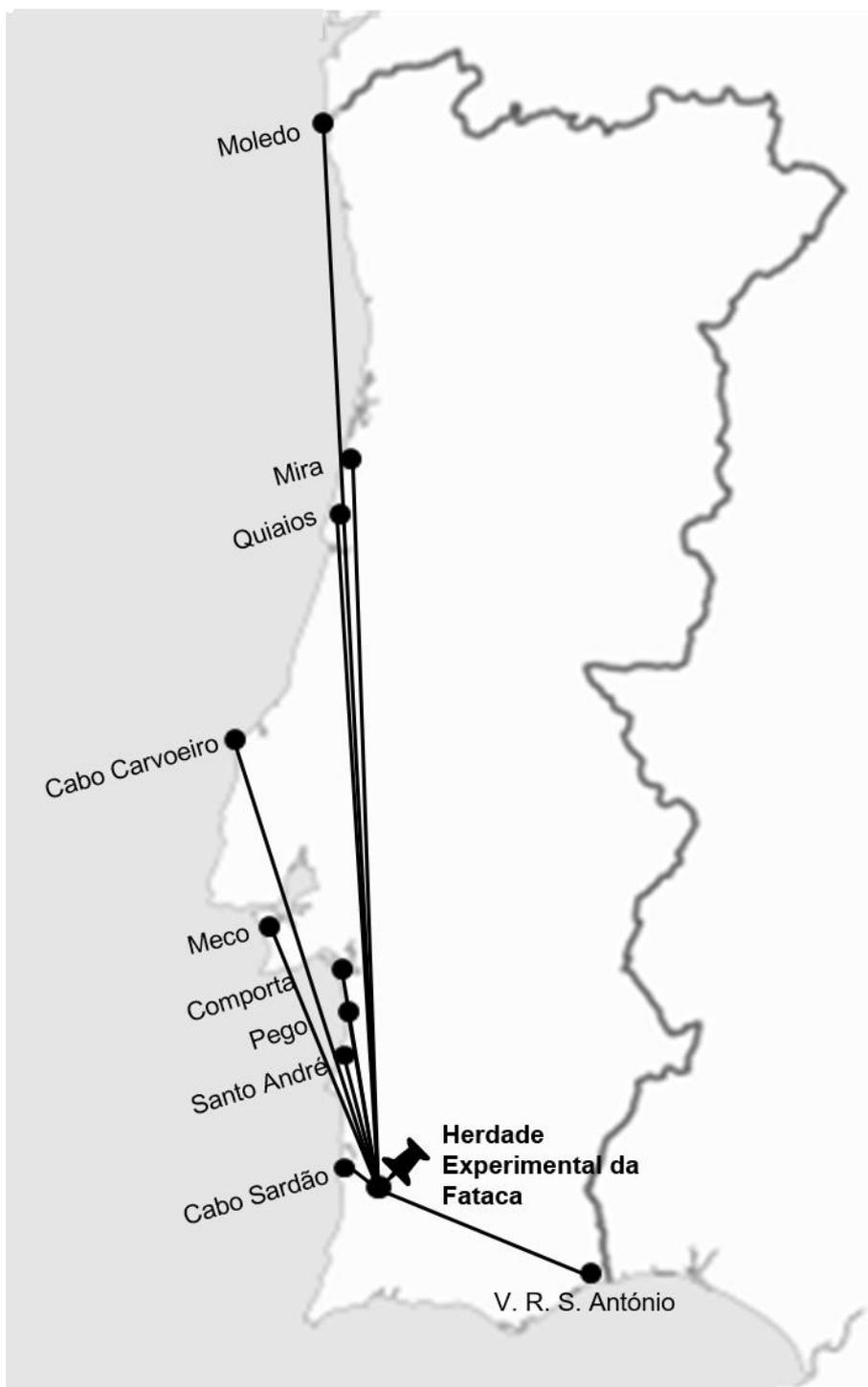
464 **Figure 2.** Hardness of white crowberry fruit samples from Fataca's clonal field, established with
465 plants from ten distinct geographical origins. Significant differences between fruit samples are
466 denoted with a different letter. n=15 per fruit sample.

467

468 **Figure 3.** Total Phenolic Content (TPC) of white crowberry fruit samples from Fataca's clonal
469 field, established with plants from ten distinct geographical origins. Significant differences
470 between fruit samples are denoted with a different letter. n=9 per fruit sample.

471

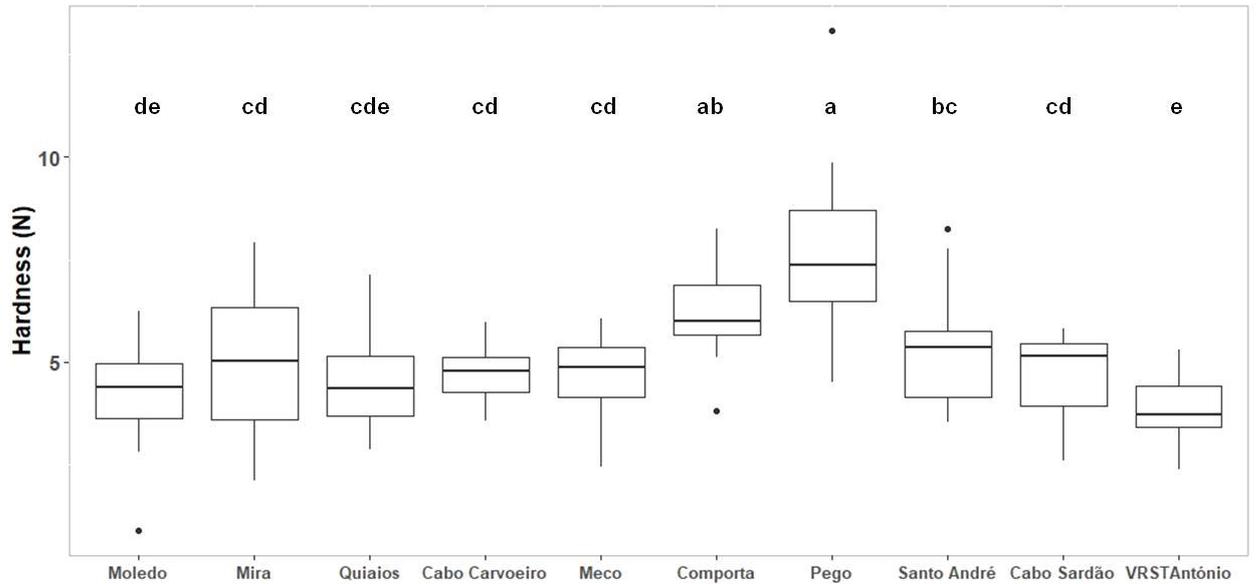
472 **Figure 4.** Principal component analysis (PCA) of white crowberry fruit samples from Fataca's
473 clonal field, established with plants from ten distinct geographical origins. For traits considered
474 see methods section. PCA abbreviations are the following: Total Phenolic Content (TPC); Soluble
475 Solids Content (SSC); Titrable Acidity (TA); L. (L*colour parameter); a. (a* colour parameter);
476 b. (b* colour parameter).



478

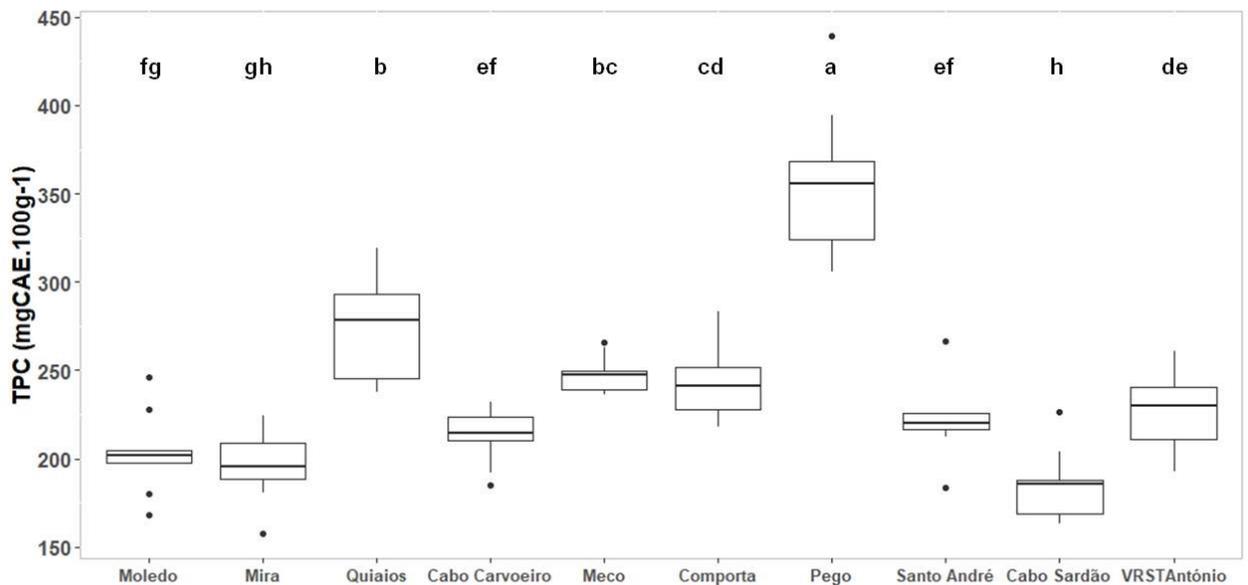
479 **Figure 1.** Location of Herdade Experimental da Fataca (INIAV), and identification of the plant's
480 collection sites for the clonal field establishment.

481



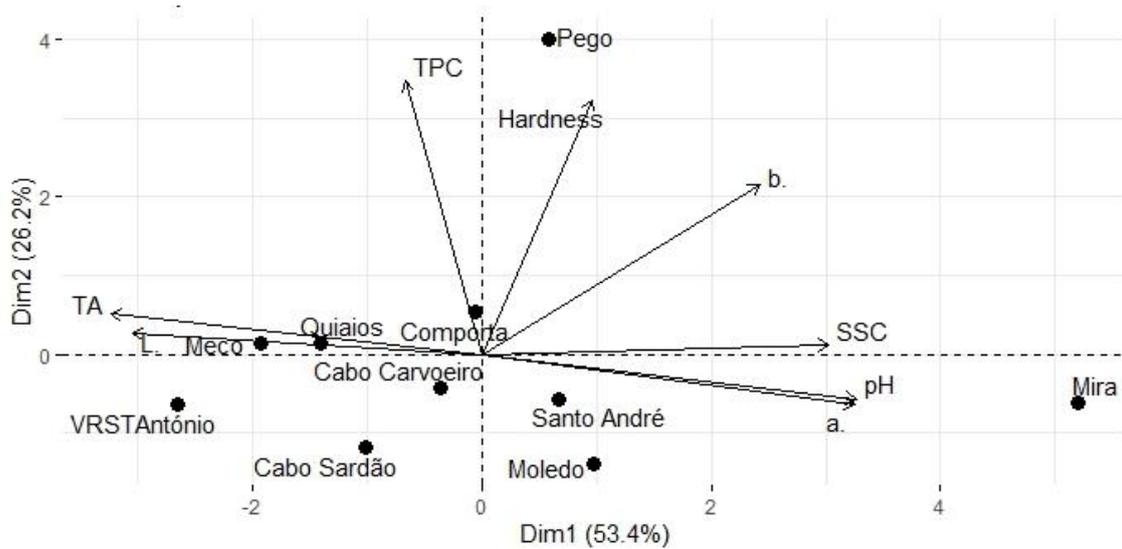
482

483 **Figure 2.** Hardness of white crowberry fruit samples from Fataca's clonal field, established with
 484 plants from ten distinct geographical origins. Significant differences between fruit samples are
 485 denoted with a different letter. n=15 per fruit sample.



486

487 **Figure 3.** Total Phenolic Content (TPC) of white crowberry fruit samples from Fataca's clonal
 488 field, established with plants from ten distinct geographical origins. Significant differences
 489 between fruit samples are denoted with a different letter. n=9 per fruit sample.



490

491 **Figure 4.** Principal component analysis (PCA) of white crowberry fruit samples from Fataca's
 492 clonal field, established with plants from ten distinct geographical origins. For traits considered
 493 see methods section. PCA abbreviations are the following: Total Phenolic Content (TPC); Soluble
 494 Solids Content (SSC); Titrable Acidity (TA); L. (L*colour parameter); a. (a* colour parameter);
 495 b. (b* colour parameter).

496