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1 **Life story of Tunisian durum wheat landraces revealed by their genetic and phenotypic**
2 **diversity**

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13

14 **Abstract:**

15 Durum wheat (*Triticum turgidum* L. subsp. *durum*) landraces represent a prominent genetic resource
16 for Mediterranean farming systems and breeding programs. Fourteen landraces sampled in Tunisia
17 were genotyped with 9 microsatellite markers and characterized with 15 morphological descriptors,
18 including resistance to the fungal disease *Septoria tritici* blotch (STB). The genetic diversity, nearly was
19 as important within landraces populations (45%) than between populations (54%). It was structured in
20 seven genetic groups and was only partly explained by the variety name or the locality of origin.
21 Populations were also greatly diversified phenotypically (Shannon-Weaver $H' = 0.54$) with traits related
22 to spike and awn colours being the most diversified. Resistance to STB was either qualitative in two
23 populations or with varying degrees of quantitative resistance in the others. A P_{st} - F_{st} comparison
24 indicate a local adaptation of the populations. Overall, the genetic structure of Tunisian durum wheat
25 landraces revealed a complex selection trajectory and seed exchanges between farmers.

26 **Keywords:** genetic resources, population genetics, phenotypic divergence, P_{st} - F_{st} comparison,
27 septoria leaf blotch.

28 INTRODUCTION

29 The importance of diversity in plant genetic resources used in agriculture and the need for biodiversity
30 conservation is now widely recognized (Maxted & al., 2010). Intra- and interspecific diversity represent
31 a heritage value that is important to preserve because it is the basis for breeders *sensu lato* – from
32 traditional farmers to global breeding companies – to adapt crops to abiotic stresses (*i.e.*
33 heterogeneous and changing environments) and provide them with resistance to biotic stresses (*i.e.*
34 pests and diseases) (Bellon, 1996). The Green Revolution, which occurred between 1950 and the late
35 1960s, led to a loss of this diversity. Traditional varieties, thereafter called landraces, were threatened
36 by genetic extinction primarily due to their replacement by modern genetically uniform varieties (Villa
37 & al., 2007). Although humans have historically domesticated and cultivated more than 7,000 species,
38 few high-yielding modern varieties from a limited number of these species constitute nowadays most
39 of the world's food resources (Perrings, 2018). This evolution to modern production systems promotes
40 the cultivation of some varieties at the expense of traditional and local crops (De luca & al., 2018) and
41 consequently is responsible of a huge intraspecific genetic erosion (Perrings, 2018; Wallace & al, 2019).
42 Moreover, this steady decline of genetic diversity canalizes the evolution of pests and forces to adopt
43 management practices, *e.g.* use of pesticides that damage the agro-ecosystem (Conversa & al, 2020).
44 Wild relatives, weedy forms and traditional varieties are especially important for conservation
45 purposes. One of the most threatened components of agricultural plant genetic resources are
46 traditional varieties commonly referred as landraces (Brush, 1997), which constitute the bulk of
47 genetic diversity in domesticated species (Conversa & al., 2020; Poudel & Johnsen, 2009; Villa & al.,
48 2007). Villa & al. (2007) define a landrace as being « a dynamic population of a cultivated plant that
49 has historical origin, distinct identity and lacks formal crop improvement, as well as often being
50 genetically diverse, locally adapted and associated with traditional farming systems ». Landraces are
51 essential heritage from farmer generations at the local and regional scale, as they are associated with
52 traditional farming systems and food trends (Negri, 2003). As a result, they are related to the biological,
53 historical, cultural and socio-economic contexts where they have been grown over generations
54 (Conversa & al, 2020), and are specifically well adapted to the environmental conditions of their
55 cultivation area (*i.e.* tolerance to biotic and abiotic stresses) (De Ron & al., 2018; A.C. Zeven, 1998). In
56 conclusion, there is a strong paradox between the loss of landraces being replaced by high yielding
57 modern varieties and the necessity for breeding programs to preserve intraspecific diversity to develop
58 innovative varieties and hybrids (De Ron & al., 2018; Govindaraj & al., 2015; Hammer & Diederichsen,
59 2009). This statement is particularly relevant to durum wheat.

60 Durum wheat (*Triticum turgidum* L. ssp. *durum*) is one of the most crucial crops in the Mediterranean
61 countries. It's a selfing tetraploid species ($2n = 4x = 28$, AABB) that originated and diversified in the
62 Mediterranean basin (Martínez-Moreno & al., 2020), which is the largest durum wheat producing
63 region worldwide, accounting for about 60% of the total growing area (Royo & al., 2017). This
64 traditional crop is the raw material for the fabrication of local dishes and products including couscous,
65 pasta, several kinds of bread and other semolina products such as bulgur and frike (Belaid 2000; Nazco
66 & al., 2014; Hammami & Sissons, 2020). Durum wheat originated in the Fertile Crescent around 10,000
67 years B.P. It spread to the western coast of the Mediterranean basin (MacKey, 2005) reaching North
68 Africa around 7,000 years B.P. (Feldman, 2001). During the migration, a combination of natural and
69 farmer's selection resulted in the development of local durum wheat landraces well adapted to their
70 region of origin and environment. But these durum wheat landraces were recently replaced by
71 improved, genetically uniform and more productive modern varieties also called 'elite cultivars'
72 (Soriano & al., 2016). In this context, the National Gene Bank of Tunisia (NGBT) has implemented an
73 ambitious program for the conservation of Tunisian durum wheat landraces both *ex situ* in gene banks
74 and *in situ* on farms. Prospecting activities carried out since 2012 by the NGBT revealed that durum

75 wheat landraces are still cultivated by some farmers in mountainous areas from the North and the
76 Center of Tunisia, under traditional farming systems. These landraces, transmitted by farmers from
77 one generation to the next, are designed by a variety name linked to a historical origin and specific
78 phenotypic characteristics. Previous studies have demonstrated that Tunisian durum wheat landraces
79 are genetically diversified (Medini & al., 2005; Robbana & al., 2019; Slim & al., 2019). Robbana & al
80 (2019) showed a variation of genetic diversity between six durum landraces using Diversity Arrays
81 Technology sequencing (DArTseq). They reported as well a higher level of genetic diversity between
82 landraces than within landraces. Slim & al. (2019) demonstrated for instance a great diversity of 41
83 Tunisian landraces using 16 molecular markers based on simple sequence repeats (SSRs), also called
84 microsatellites, with clear differentiation between landraces and elite cultivars. They detected also five
85 genetic clusters structuring landraces with a strong North-South stratification. SSRs have been
86 reported to be the most widely used markers to study the genetic diversity in wheat germplasm due
87 to their large distribution in the genome, codominant nature, high polymorphism, good reproducibility
88 and ease of application (Russell & al., 1997; Medini & al., 2005). These multiallelic markers allow to
89 capture higher variability than biallelic markers like Single Nucleotide Polymorphism (SNP), Amplified
90 Fragment Length Polymorphism (AFLP) or DArT markers (Semagn & al., 2014; Targońska & al., 2016;
91 Hurtado & al., 2008).

92 Several studies have shown that Tunisian durum wheat landraces are agro-morphologically diversified
93 and can be exploited for their large panel of technological properties (Ayed & al., 2010; Nazco & al.,
94 2012; Ayadi & al., 2012; Chamekh & al. 2015; Babay & al. 2019; Bouacha & Rezgui, 2017; Yacoubi & al.,
95 2020). Ayed & al. (2010) showed a huge phenotypic diversity of Tunisian durum wheat landraces for
96 six qualitative traits: seed colour, seed size, glume colour, glume pubescence, spike density and beak
97 length (Ayed & al., 2010). Nazco & al. (2012) highlighted that landraces from the western
98 Mediterranean countries such as Tunisia have heavier grains and higher grain-filling rates than those
99 from the eastern Mediterranean countries. Previous studies established that landraces overstep
100 improved genotypes for agronomic traits such as plant height, biomass, straw yield and also have high
101 grain yield, high nitrogen utilization efficiency, high nitrogen use efficiency, high NADH-dependent
102 glutamate synthase activity and high NADH-dependent glutamate dehydrogenase activity (Ayadi & al.,
103 2012; Chamekh & al. 2015). Additional studies shown that landraces have a high protein content and
104 consequently high physico-chemical technological propriety of semolina and pasta (Babay & al., 2019,
105 Bouacha & Rezgui, 2017). Landraces are known to be sources of increased biomass and thousand
106 kernel weight, both being important traits for adaptation to drought and heat stresses (Yacoubi & al.,
107 2019), but also of genetic resistances to pests, including fungal pathogens. Recently, Huhn & al. (2012)
108 screened a wheat collection including Tunisian durum wheat landraces and revealed that five lines are
109 moderately resistant to *Fusarium* head blight (Huhn & al., 2012). Ferjaoui & al. (2011, 2015) and Ouaja
110 & al. (2020) identified other lines with resistance to *Septoria tritici* blotch (STB). This foliar disease,
111 caused by the fungal pathogen *Zymoseptoria tritici*, can cause important yield losses in Tunisia
112 (Ferjaoui & al., 2015). Most of the Tunisian durum wheat landraces remain to be genetically and
113 phenotypically characterized, including for their level of resistance to STB disease.

114 The large majority of studies on durum wheat landraces examine collections of lines belonging to
115 different populations, with only a few or sometimes only one individual per population. It is advisable
116 to extend the population-study to more individuals per population in order to better characterize
117 genetic and phenotypic diversity both between and within populations. To this end, we studied 14
118 Tunisian durum wheat landraces collected by the NGBT between 2015 and 2017. Some landraces had
119 the same name but were grown in different localities while others were grown by the same farmer but
120 had different names. We thus decided to investigate whether these landraces are different (at the
121 molecular and phenotypic levels) or not and what are their relationships. Concretely, these landraces

122 consisted here and in the whole manuscript as a ‘population’, *i.e.* a group of individuals collected at a
123 specific farmer’s field and reported by each farmer to be a landrace. 16 to 51 individuals were randomly
124 selected from each population, which were characterized for: (i) their neutral genetic diversity and
125 structure with SSR markers; (ii) their phenotypic diversity based on agro-morphological characters and
126 their response to STB; and (iii) their phenotypic differentiation (P_{st}) comparatively with neutral genetic
127 differentiation (F_{st}) in order to determine whether phenotypic differences between populations were
128 due to selection.

129

130 **RESULTS**

131 **Identification of hexaploid lineages, historical contaminant of Tunisian durum wheat fields**

132 A first genetic structure analysis of the durum wheat germplasm showed that the lineages from the 14
133 populations can be divided into 10 genetically distinct groups (strongest ΔK of 14.93) (data not shown).
134 Most genetic groups were composed of lineages coming from different “variety-locality” populations,
135 designed by a “variety” name and its “locality” of origin. At $k=10$, 33 lineages corresponding to 23
136 unique multilocus genotypes (MLGs) belong exclusively to three genetic groups genetically close from
137 each other. These lineages stood out for having a characteristic of spikes different from all the others,
138 *i.e.* long cylindrical white spikes. These lineages were found in eight out of the 14 populations collected
139 in the North or the Center of Tunisia, *i.e.* Roussia Joumine, Mahmoudi Amdoun, Mahmoudi Oued
140 Sbaihia, Mahmoudi El Jouf, Chili El Jouf, Chili Lansarine, Aouija Msaken and Mahmoudi Msaken. They
141 correspond to what is called ‘mule’s tail’ or ‘mare’s tail’ by Tunisian farmers. ‘Mule’s tails’ are
142 undesirable contaminants growing in Tunisian durum wheat fields, whose grains are too “soft” and
143 become flour rather than semolina when milled. The karyotype analysis showed that all mule tails’
144 lineages suspected to be hexaploid wheat species rather than tetraploid durum wheat carried indeed
145 42 chromosomes vs. 28 for ‘Karim’ and ‘Mahmoudi-101’ varieties (Figure 1). The highest proportion of
146 ‘mule’s tail’ was found on the two populations Mahmoudi Oued Sbaihia and Mahmoudi El Jouf, both
147 cultivated in the North East of Tunisia (Governorate Zaghuan). Therefore, 33 hexaploid lineages were
148 eliminated from the dataset and further analyses were performed with tetraploid wheat lineages only.

149 **Characterization using microsatellite markers**

150 From the collection of 335 lineages of durum wheat, the polymorphic microsatellite (SSR) markers
151 amplified 61 different alleles (Table 1). Size ranges of alleles did not overlap between markers tagged
152 with the same dye. The number of alleles per locus ranged from three for the marker Xgpw2239 to 12
153 for the marker Xgwm372. In total, 23 private alleles were identified in eight out of the nine loci. The
154 low mean- H_o (0.003-0.014) and $-H_s$ (0.147-0.317) values revealed a low level of heterozygosity. All
155 markers were highly informative and polymorphic as evidenced from their high PIC value (ranging from
156 0.503 for Xgpw2239 to 0,966 for Xgwm413), and were characterized by a high Fixation index (0.961-
157 0.986). An analysis of the SSR markers with the BayeScan program revealed no outlier loci.
158 Furthermore, genotype accumulation curves indicated that eight loci only are required to discriminate
159 between individuals in the studied populations (Figure-ESM1). Hence, this multiplex of nine SSR
160 markers is a valuable tool for population genetics studies of durum wheat and genotyping results can
161 be further used for characterizing the diversity of the 14 durum wheat populations.

162 **Distribution of genetic diversity**

163 The AMOVA test determined that there is almost as much inter and intra-population genetic variation
164 (54% of genetic variability was explained by inter-populations variability vs. 45% by diversity within

165 populations; Table 2). As the level of heterozygosity is very low (intra-individual variation around 1%),
166 the studied lineages can be considered fixed after one or two generations of selfing.

167 The genetic diversity indices (genotypic richness; Shannon, Stoddart-Taylor, Simpson and Evenness
168 indexes) for nine SSR markers calculated for each population are given in Table 3. The genotypic
169 richness ranges from 0.107 in the durum wheat population Chili El Jouv to 0.551 in Roussia Joumine.
170 The Shannon index ranges from 0.545 in Chili El Jouv to 2.625 in Roussia Joumine. The Stoddart and
171 Taylor's index ranges from 1.33 for Chili El Jouv to 11.25 in Roussia Joumine. The Shannon Wiener index
172 is sensitive to genotypic richness in samples of uneven sizes (Grünwald & al., 2003). However, the high
173 positive correlation between the Shannon and Stoddart-Taylor indexes ($r=0.86$) reinforces the
174 conclusions that can be drawn. The Simpson's index, measuring the probability that two randomly
175 selected genotypes are different, varies from 0.25 in Chili El Jouv to 0.911 in Roussia Joumine and was
176 also highly correlated with the Shannon index ($r=0.96$). The Evenness index ranges also from 0.25 in Chili
177 El Jouv to 0.801 in Roussia Joumine, indicating that the MLGs observed in Roussia Joumine are closer
178 to equal abundance than for the other populations. As estimates of genotypic diversity include
179 genotype richness and evenness of distribution of genotypes within the sample (Grünwald & al., 2003),
180 combining results of different evaluated parameters lead us to state that Roussia Joumine stands out
181 as being the most diversified population. Indeed, it has the highest genotypic richness, highest
182 genotypic diversity index and highest Evenness index. On the other hand, Chili El Jouv is the least
183 genetically diverse population.

184 Overall, 118 MLGs were identified from all populations, meaning that 64.8% of lineages are clonal. At
185 equal population sizes, the eMLGs range from 2.6 for Chili El Jouv to 9.6 for Roussia Joumine. Roussia
186 Joumine stands out by its high number of MLGs.

187 **Genetic structure of Tunisian durum wheat populations**

188 For a number of genetic groups (k) varying from $k=3$ to $k=7$, the analysis of the genetic structure of the
189 14 durum wheat populations highlighted that the strongest ΔK was obtained for $k=3$ (with $\Delta K=193.84$),
190 $k=5$ (with $\Delta K=77.46$) and $k=7$ (with $\Delta K=44.19$) (Figure 2). The number of genetic groups $k=7$ was chosen
191 because it allows a better description of the genetic structuration of the 14 populations and it
192 maximises the genetic differentiation, *i.e.* F_{st} , compared to the other partitioning of groups. The seven
193 populations Bidi Kasserine, Ajimi Kasserine, Chili Lansarine, Chili El Jouv, Mahmoudi Msaken,
194 Mahmoudi Sejnane and Roussia Joumine are made of individuals belonging mainly to one genetic
195 group, showing the homogeneity of individuals within these populations. The two populations from
196 Kasserine (collected from the same farmer), Bidi Kasserine and Ajimi Kasserine, are identical,
197 suggesting that they have a similar origin despite their different names. Similarly, individuals from the
198 populations Chili Lansarine and Chili El Jouv also belong to the same genetic group, implying that
199 populations with the same variety name but from different localities aroused from a common origin.
200 The structure of these populations is therefore explained by a combination of effects, including the
201 locality of origin and the variety name.

202 The minimum spanning network (MSN) visualizes relationships among MLGs and indicates the
203 existence of one MLG of high frequency in nine different populations (Figure 3). The populations
204 sharing this common MLG (MLG.66) are Mahmoudi Oued Sbaihia, Mahmoudi Joumine, Mahmoudi
205 Amdoun, Mahmoudi El Jouv, Richi El Jouv, Aouija Msaken, Beskri Msaken, Chili Lansarine and Chili El
206 Jouv. Other high frequency MLGs were found in two or three populations. Several MLGs detected a
207 single time were highly distant from other MLGs belonging to the same population. MLGs from the
208 population Roussia Joumine were close to each other, and close to MLGs from the two populations
209 Mahmoudi Msaken and Mahmoudi Sejnane. These three populations were distant from the other 11

210 populations. MLGs belonging to Ajimi Kasserine and Bidi Kasserine were exceptionally close to each
211 other.

212 Genetic differentiation between populations was estimated with pairwise F_{st} (F_{st} values between pairs
213 of populations). The matrix of pairwise F_{st} shows values ranging from 0.016 (between Mahmoudi El
214 Jouf and Mahmoudi Oued Sbaihia) to 0.887 (between Ajimi Kasserine and Mahmoudi Msaken), with
215 significant genetic differentiation between the majority of populations (Table 4). In some cases, the
216 genetic differentiation is not significant (5% threshold), for example between the two populations from
217 Kasserine (Bidi and Ajimi varieties), the two Mahmoudi populations from El Jouf and Oued Sbaihia, and
218 the two Mahmoudi populations from Oued Sbaihia and Joumine; while the two Mahmoudi populations
219 from El Jouf and Joumine were significantly different. Similarly, even if significant, the F_{st} value
220 between the two Chili populations from El Jouf and Lansarine is low and very close to the significance
221 threshold. Similarities and differences between populations testify of the history of seed exchanges
222 that took place between farmers. Genetic differentiation between the seven genetic groups
223 determined by STRUCTURE was also calculated using the F_{st} index (Table-ESM1). An individual was
224 considered to belong to a specific genetic group when it has at least 50% affiliation to this group. The
225 matrix of pairwise F_{st} shows values ranging from 0.36 to 0.64 between the seven genetic groups (Table-
226 ESM1) indicating that they are significantly different from each other.

227 **Phylogenetic relationship between the 14 populations and selection trajectory of Tunisian** 228 **landraces**

229 Tunisian durum wheat landraces were collected and stored in genebanks since the studies realized by
230 Félicien Boeuf (1875-1961) in the beginning of the 20th century. We studied the relationship between
231 the 14 durum wheat populations collected by the NGBT in 2015 and 2017 and 40 landraces collected
232 between 1911 and 1976 carrying the same variety name (seed samples received from NPGS and NGBT
233 genebanks) (Table-ESM2). The 14 populations, the 40 landraces and individuals from three seed lots
234 of the modern variety 'Karim' were genotyped with the nine microsatellite markers described earlier.
235 A neighbour-joining phylogenetic tree was built with microsatellite genotypes (Figure 4). The tree
236 revealed a complex relationship between historical landraces, as samples with the same variety name
237 did not necessarily clustered together. First, the historical landraces called Mahmoudi were mostly
238 divided into two clusters: 14 from the 24 Mahmoudi landraces clustered at the bottom of the tree and
239 were only admixed with the two Bidi landraces (PI534469, 1976) and (PI576736, 1976); 6 of the
240 remaining clustered at the top of the tree and were admixed with one Bidi (Cltr3811, 1912) and one
241 Chili (PI534336, 1976). In this last cluster, the Mahmoudi Msaken population was strongly related to
242 Chili (PI534336, 1976) (bootstrap support of 98.9), and the population Roussia Joumine remains
243 relatively distant from other historical landraces and studied populations. This suggests that
244 Mahmoudi was submitted to at least two different environments, and that in each environment it was
245 confronted to different landraces with which some exchanges could have occurred through open
246 pollination. Second, most of the landraces called Chili were grouped in the same cluster in the middle
247 of the tree, together with few Mahmoudi and Bidi landraces. Among them, landraces Mahmoudi
248 (Cltr15501, 1972) and Bidi (PI157961, 1947) were identical to three Chili landraces. This identity could
249 not have arisen by chance or mutation and this strongly suggests that these two landraces belong to
250 the Chili group and that their names have been badly assigned. This raises the question about the
251 collection and maintenance of accessions in gene banks. Nine of the 14 studied populations also
252 grouped in this cluster with Chili landraces: including Chili Lansarine and Chili El jouf, but also
253 Mahmoudi El Jouf, Mahmoudi Oued Sbaihia, Beskri Msaken, Mahmoudi Joumine, Aouija Msaken, Richi
254 El Jouf and Mahmoudi Amdoun; indicating a strong contribution of Chili historical landraces to the
255 genetic constitution of the studied populations. Third, the populations Ajimi Kasserine and Bidi

256 Kasserine grouped apart from other populations and landraces, and not surprisingly were strongly
257 related to each other (bootstrap support of 100). Finally, in the NGBT and INAT seed lots of 'Karim',
258 the same MLG was observed for all individuals with eight of the nine markers, leading us to conclude
259 that these two seed lots are genetically homogeneous; variability of the ninth marker was attributed
260 to a technical bias. For the CRP seed lot, three different MLGs were observed among which the MLG
261 detected in the two other seed lots of 'Karim'. Thus, this third seed lot corresponds well to 'Karim' but
262 is not genetically pure. More surprisingly, Mahmoudi Sejnane was strongly related to Karim (bootstrap
263 support of 98.4), which may indicate that the farmer from Sejnane erroneously considered his seed lot
264 as a landrace.

265 **Phenotypic diversity of the 14 populations**

266 **-Shannon Index**

267 The phenotypic diversity was assessed on 273 from the 335 genotyped lineages for the 15 evaluated
268 phenotypic traits (Table ESM3) using the Shannon diversity index (H') at the population and at the
269 genetic group levels (Table 5). At the population level, the mean H' for all characterized traits ranged
270 from $H'=0.41$ in Mahmoudi Msaken to $H'=0.66$ in Richi El Jouf, making Richi El Jouf the most
271 phenotypically diversified population. The spike colour (SC) was the most polymorphic trait over the
272 14 populations (mean $H'=1.06$), with spikes being mostly slightly coloured. The seed shape (SS) was
273 the least diversified trait over all populations (mean $H'=0.08$), with the majority of populations having
274 semi-elongated seeds.

275 At the genetic group level, the greatest diversity ($H'=0.85$) was found in genetic group 4 composed by
276 individuals belonging to Richi El Jouf, Mahmoudi Msaken and Aouija Msaken. Six from the 15 evaluated
277 traits were highly polymorphic in this genetic group compared to others: SS ($H'=0.87$), spike density
278 (SD) ($H'=0.93$), length of spike without awns (LSWA) ($H'=0.95$), awn colour (AC) ($H'=1.12$), number of
279 spikelets per spike (NSS) ($H'=1.28$) and thousand grains weight (TGW) ($H'=1.03$). The genetic group 7
280 was the least phenotypically diversified ($H'=0.28$) but this group consists of only two lineages and is
281 therefore not representative. The phenotypic diversity observed within the genetic groups indicate
282 that they are not phenotypically homogeneous. Nevertheless, some phenotypic traits are clearly
283 distinct between them. For example, SS with parallel-edge spikes being dominant in genetic groups 2,
284 3 and 7 (monomorphic $H'=0$) while less frequent in groups 1, 4 and 5. Genetic group 6 was
285 characterized by a particular stunted-spike shape, which is not listed in the official classification from
286 UPOV. Few individuals from genetic group 4 also had this particular stunted-spike shape. Moreover,
287 SC was the most diverse phenotypic trait among all the genetic groups (mean $H' = 1.00$).

288 **-Statistics on quantitative traits**

289 Nine phenotypic traits were quantitatively evaluated and subjected to statistical analyses. Significant
290 positive correlations (at $p<0.01$) were detected between heading date (HS) and plant height (HI) (0.71),
291 and not surprisingly between length of awns (LA) and length of awns in relation to spike (LAS) (0.82)
292 (Figure-ESM2). Concerning *Septoria tritici* blotch (STB), a significant negative correlation (at $p<0.01$)
293 was detected between area under disease progress curve (AUDPC) and HI (-0.43), and between AUDPC
294 and HS (-0.57), corroborating that HI and HS can influence disease development. Early heading and
295 short lineages tend to be more infected by STB. Finally, LSWA and SD were also significantly negatively
296 correlated (-0.50).

297 Phenotypes of all quantitative variables were not normally distributed. Only the trait NSS could be
298 normalized following a Box-Cox transformation (Box & Cox, 1964). A significant difference between
299 population means was found for 8 quantitative traits (at $p<0.001$); LAS was not significantly different

300 between population means. We observed a maximum of pairwise significant differences between
301 populations for the trait NSS (35 pairs) and a minimum for the trait LA (five pairs) (Table 6).

302 The population Mahmoudi Sejnane cumulated the highest number of differences with other
303 populations for quantitative phenotypic traits, except with the population Mahmoudi Msaken (Table
304 6). In total, 31 pairs of populations were not significantly different for any of the phenotypic traits
305 studied, including the phylogenetically related (Figure 4) Richi El Jouf and Aouija Msaken, Mahmoudi
306 Joumine and Beskri Msaken, Mahmoudi El Jouf and Mahmoudi Oued Sbaihia, Bidi Kasserine and Ajimi
307 Kasserine, and even Mahmoudi Sejnane and Mahmoudi Msaken (Table 6).

308 **-Response of the 14 populations to STB infection**

309 The 14 durum wheat populations were contrasted in their response to STB infection (Figure 5). The
310 two populations Ajimi Kasserine and Roussia Joumine show a nearly complete and qualitative
311 resistance level to the *Z. tritici* strain IPO91009, while Mahmoudi Sejnane appeared highly susceptible.
312 The other 11 populations show varying degrees of resistance or susceptibility to STB. Within each
313 population, a variation in response to STB was also observed between lineages.

314 **-Factor Analysis of Mixed Data (FAMD)**

315 A Factor Analysis of Mixed Data (FAMD) was performed on both quantitative and qualitative variables
316 (Figure 6). The first two principal components in the FAMD accounted for 16.5% and 9.8 % of the total
317 variation, respectively, and together explained 26.3% of the total variation (Figure 6A). SS, HI, HS, NSS,
318 TGW, AUDPC and LA were the most important traits contributing to the first principal component
319 (Figure ESM3). Shape of grain (SG), SS, SD, AUDPC, TGW and LSWA contributed significantly to the
320 second principal component (Figure ESM3). FAMD allowed to discriminate between three groups of
321 populations. The first group includes Mahmoudi Sejnane and Mahmoudi Msaken (top left quadrant of
322 Figure 6A), which are differentiated from the others populations by their stunted spike shape (Figure
323 6B), earlier heading date, shorter plant height, higher susceptibility to STB, and lower number of
324 spikelets per spike ($p < 0.05$). The second group includes Aouija Msaken and Richi El Jouf (top right
325 quadrant of Figure 6A), which are distinguishable by their fusiform spike shape (Figure 6C), elongated
326 shape of grains (Figure 6D), and the highest TGW compared to other populations ($p < 0.05$). Finally, a
327 third group includes the other 10 populations, which are characterized by spikes with parallel edges
328 (Figure 6F) and semi-elongated grains (Figure 6E).

329 **- P_{st} - F_{st} comparison and sensitive analysis**

330 Comparisons of P_{st} and F_{st} show that most phenotypic traits are highly divergent reflecting therefore a
331 local adaptation rather than a genetic drift, because lower 95% CI for P_{st} were higher than the $F_{st}=0.57$
332 (Figure 7). Only for LAS, P_{st} didn't differ from F_{st} since the lower 95% CI= $0.5066 < F_{st}$. However, the
333 observed degree of differentiation is unlikely due to genetic drift for this trait (Leinonen & al., 2013).
334 Evidence for the robustness of the sensitive P_{st} - F_{st} analysis varied among the eight highly divergent
335 phenotypic traits but was exceptionally strong for some traits such as HI, HS and AUDPC, which had
336 critical c/h^2 around 0.1 (Table 7). Similarly, LSWA, NSS and TGW have a critical c/h^2 ranging from 0.16
337 to 0.21. Therefore, assuming that these traits are under selection and not under genetic drift is a very
338 robust inference since phenotypic variance across populations that is explained by additive genetic
339 effects would need to be at least six times lower than the phenotypic variation within populations to
340 be explained by genetic drift. For SD and LA, the higher critical c/h^2 of 0.30 and 0.61, indicates that
341 phenotypic differentiation is lower and the conclusion of selection is clearly not conservative.

342

343 **DISCUSSION**

344 Durum wheat landraces traditionally used by Tunisian farmers have been progressively replaced by
345 modern cultivars provided by international (i.e. CIMMYT) and Tunisian (i.e. INRAT) breeding programs.
346 This led to a rapid increase in productivity of wheat and was accompanied by the multilateral trade
347 liberalization on its value chain as well as staple food affordability in both rural and urban Tunisian
348 areas (La Rovere & al., 2010). However, few farmers are still until now cultivating landraces in some
349 mountainous or dry areas of Tunisia characterized by low input farming systems, for their own
350 consumption or to be sold on local markets. Understanding the genetic and phenotypic diversity of
351 landraces can be used to legitimate the preservation of these important genetic resources in order to
352 improve the adaptability of durum wheat crop to marginal environments and threatening pathogens
353 (Pietrusińska & al., 2018).

354 **1- Durum wheat landraces are genetically diverse and dynamic populations**

355 **-Microsatellite markers**

356 PIC values are a good indication of the informativeness of the multiplexed panel of the microsatellite
357 (SSR) markers used in this study. According to the classification by Botstein & al. (1980), our nine SSRs
358 were moderately ($0.25 < PIC < 0.5$) to highly ($PIC > 0.5$) informative, therefore sufficient to discriminate
359 between populations and useful for further genetic diversity studies and seed bank management.

360 The genetic diversity of the 14 populations assessed with these SSRs ($H_e = 0.57$) was almost equivalent
361 to the genetic diversity of several collections of durum wheat landraces from different Mediterranean
362 countries such as Tunisia, Morocco or Italy with average gene diversity values between 0.44 and 0.69
363 (Nefzaoui & al., 2014; Slim & al., 2019; Sahri & al., 2014; Soriano & al., 2016). This high genetic diversity
364 can explain why Tunisia has previously been considered as a centre of diversity for durum wheat (Ayed
365 & al., 2010).

366 **-Repartition of genetic diversity**

367 The Analysis of MOlecular Variance (AMOVA) evidenced the great genetic diversity present within the
368 durum wheat populations since variation was only slightly greater between populations (54%) than
369 within population (45%) (Table 2). It can be the result of outcrossing and fitness-relevant mutations.
370 The global F_{st} value of 0.5 also supports a high differentiation between the 14 populations. Similar
371 results were obtained for Ethiopian durum wheat landraces with a slightly higher genetic variance
372 between populations (52%) than between individuals within genetic groups (48%) (Alemu & al., 2020).
373 However, several other studies rather report a greater genetic variance within than between
374 populations (Kabbaj & al., 2017; Soriano & al., 2016; Kyratzis & al., 2019; Asmamaw & al., 2019),
375 potentially explained by seed exchange or selection by farmers for similar traits.

376 **-Structuration of lineages**

377 Based on a bayesian method, the 14 populations were stratified into seven genetic groups (Figure 2).
378 Only few lineages showed a low membership coefficient q -value suggesting a weak level of admixture
379 among the populations. This is probably due to the fact that wheat is mostly autogamous and to the
380 selection pressure exerted by farmers. Some genetic groups are composed of lineages belonging to a
381 single population while other genetic groups include lineages from different populations. Two genetic
382 groups (G2 and G6) are exclusive to a single population (Roussia Joumine and Mahmoudi Sejnane,
383 respectively), as supported by the distant position of each population on the minimum spanning
384 network (Figure 3). Roussia Joumine is the only population with variety name 'Roussia', and this
385 landrace is likely to be genetically distinct from the others. Mahmoudi Sejnane seems unrelated to

386 other 'Mahmoudi' populations. This is also true for Mahmoudi Msaken composed only from lineages
387 belonging to a group (G4) which also includes individuals from the Richi El jouf and Aouija Msaken
388 populations. Another group (G5) is composed of lineages coming from eight populations, including
389 nearly all individuals from Chili El Jouf and Chili Lansarine. This suggest that both populations probably
390 have a common origin and thus highlights the pronounced impact of the variety name in the genetic
391 structuration. Furthermore, a group (G3) is composed of lineages coming from four populations,
392 including nearly all individuals from Ajimi Kasserine and Bidi Kasserine. This strong evidence of the
393 impact of locality on the genetic of Tunisian durum wheat landraces structuration, in addition to the
394 expected effect of the variety name, has been demonstrated in previous studies (Slim & al., 2019;
395 Soriano & al., 2016; Rufo & al., 2019). Finally, a group (G1) is composed of lineages coming from seven
396 populations having different origins and different names of variety, underlining the more complex
397 genetic structure of several populations. Genetic structuration of these 14 durum wheat populations
398 is therefore explained by a combination of effects, including the locality of origin and the variety name,
399 that likely results from the farmer-mediated selection and exchange trajectory of each population.

400 **-Agro-morphological diversity**

401 Characterizing the phenotypic diversity of durum wheat landraces, known to be often agro-
402 morphologically heterogeneous (Royo & al., 2017), is prominent towards an understanding of their life
403 history, in other words the selection trajectory and seed exchanges between farmers. Indeed, the 14
404 studied populations were agro-morphologically diversified with a mean Shannon-Wiener index of
405 $H' = 0.54$, estimated with 15 phenotypic traits. In previous studies, even greater agro-morphological
406 diversities were observed with $H' = 0.77$ within 30 Tunisian durum wheat landraces characterized for
407 11 phenotypic traits (Ayed & al., 2010) and $H' = 0.63$ within a collection of 368 Tunisian durum wheat
408 accessions characterized for 9 phenotypic traits (Slim & al. 2011). The least diversified traits among the
409 14 populations were SG (shape of grain) ($H' = 0.08$), SS (spike shape) ($H' = 0.14$) and LAS (length of awns
410 in relation to spike) ($H' = 0.14$). The most frequent shape of grains is semi-elongated, which is associated
411 with a spike shape with parallel edges. Thus, it suggests a selection pressure exerted by Tunisian
412 farmers on their landraces for the same spike shapes. However, grain and spike shapes appeared highly
413 diversified in several other studies (Belhadj & al., 2015; Bechere & al., 1996; Al Khanjari & al., 2008).
414 The most diversified traits among the 14 populations were SC (spike colour) ($H' = 1.06$), PgA
415 (anthocyanin colouration of awns) ($H' = 0.82$) and AC (awn colour) ($H' = 0.75$) (Table 6). Spike and awn
416 colours are important for Tunisian farmers. For example, farmers associate the black colour of awns
417 with the Mahmoudi landrace distinguished for its higher quality for semolina and bread bakery. These
418 findings are consistent with previous studies showing that awn colour was often among the most
419 diversified traits in Tunisian and Omani landraces (Belhadj & al., 2015; Al Khanjari & al., 2008).

420 Population genetic groups did not appear congruent with the structure of agro-morphological diversity
421 while another study highlighted a reliable relationship between genetic and phenotypic population
422 structures, and the connection of both with the geographic origin of the landraces (Soriano & al., 2016).
423 The overall agro-morphological diversity measured with Shannon index does not differ significantly
424 between the 14 populations. A better discrimination between the populations could be obtained with
425 more informative traits and tighter descriptors. Traits related to spike and awn colours could be
426 described using a colour palette for better discrimination between shades, such as Roussia Joumine's
427 russet colour of awns. A broader list of spike shapes could be established including for example the
428 stunted form of spikes found in Mahmoudi Msaken and Mahmoudi Sejnane populations.

429

430

431 **2- Assessing the origin of durum wheat landraces**

432 Characterization of the 14 populations revealed the complexity of the history of durum wheat
433 cultivation in Tunisia. Behind each population hides a particular life story that can only be understood
434 in light of converging evidences from genetic and agro-morphological descriptions of the populations,
435 and from their local context.

436 **-Roussia Joumine, standing-out redhead**

437 The population Roussia Joumine, the only one with variety name 'Roussia', stood-out for the
438 homogeneous russet colour of spikes from all its characterized lineages, although this criterion is not
439 in the UPOV classification. It was also strongly genetically distinguishable from all the other studied
440 populations. In literature, Roussia 875 is a local landrace from the north of Tunisia identified in 1927
441 and registered from 1953 to 1973 (Ammar & al., 2011). Gharbi & El Felah (2013) reported that
442 'Roussia', keep its name from an original population that was cultivated extensively in field crops. Slim
443 & al. (2019) previously reported a dendrogram for the durum wheat collection of the NGBT on which
444 Roussia Joumine and Mahmoudi Amdoun clustered together, while in our study both populations were
445 highly differentiated ($F_{st}=0.6$). Indeed, Roussia Joumine showed higher values for all genetic indices
446 evaluated, and it was the most genetically diversified population while being phenotypically less
447 divergent. This landrace was as well remarkably resistant to STB and could be an interesting source of
448 resistances to this disease.

449 **-Chili El Jouf and Chili Lansarine, a common origin**

450 The two populations Chili El Jouf and Chili Lansarine, were significantly weakly differentiated with SSR
451 markers ($F_{st}=0.03$) and almost exclusively composed of lineages classified in the same genetic group
452 (Figure 2) and sharing an MLG (Figure 3). Both populations were grouped with 'Chili' landraces
453 collected by USDA in Tunisia in 1979 (Figure 4). In literature, the origin of 'Chili' landrace is not very
454 clear. According to Saade & al (1996) 'Chili' was selected from a commercial shipment imported from
455 Chile in the early 1960s and rapidly adopted by Tunisian farmers. Deghais & al. (2007) reported that,
456 in 1932, sir Racine, a French industrialist, sent from Marseille (France) 100 kg of Chili seeds to sir
457 Charles Fabre, a farmer in Bou Salem. In 1938, the Botanical Service of Tunis recovered a part of these
458 Chili seeds and incorporated them in regional field trials to evaluate its adaptative potential to regional
459 conditions. Propagation of Chili then took place as early as 1941/1942 and the Botanical Service of
460 Tunis selected the variety 'Chili-931', which was registered in the official Tunisian catalogue of varieties
461 in 1953. Since 1985, the cultivation of this variety has been largely abandoned with the exception of a
462 few niches and it was removed from the catalogue in 1993.

463 Chili El Jouf and Chili Lansarine were identical for most phenotypic traits studied, except for the NSS,
464 being higher in Chili El Jouf than in Chili Lansarine (Table 6). The difference between the two
465 populations could result from a selection by the farmers from El Jouf for a better yield, whose NSS is a
466 component. Moreover, as NSS is influenced by environmental conditions, such as temperature and
467 day length (Knezevic & al., 2007; Rawson, 1970), the difference between the two Chili populations
468 could also have resulted from a local adaptation to environmental conditions. Genetic analysis
469 demonstrated that Chili El Jouf had lower genetic diversity than all studied populations (Table 3).

470 **-Aouija Msaken and Richi El Jouf, distant cousins**

471 The populations Aouija Msaken and Richi El Jouf stood out for their similar fusiform spike shape and
472 elongated grain shape (Figure 6). These populations are genetically heterogeneous and share lineages
473 belonging to the same genetic groups (Figure 2) but remain genetically differentiated from each other

474 ($F_{st}=0.27$). Based on Neighbour joining clustering, Richi El Jouf and Aouija Msaken were genetically
475 closely related (bootstrap support of 85.2), with an F_{st} index around 0.2 being the lowest between Richi
476 El Jouf and any other population. In addition, both populations share a common MLG (Figure 3)
477 suggesting that they have a common ancestor. Richi is a very old variety related to the species *Triticum*
478 *polonicum* (personal communication from NGBT). According to some studies, Aouija is also called
479 "Aouej", "Swabaa Aljia" or "Aouiji". Several forms were introduced in 1913 and 1916 by Boeuf from
480 Morocco where they were called "Zréa Laouaja" in relation to the hunchback shape of the grain
481 reminiscent of that of *Triticum polonicum*. Richi El Jouf was the most diversified population based on
482 all measured traits ($H'=0.66$), while Aouija Msaken had a lower phenotypic diversity ($H'=0.50$).

483 **-Ajimi Kasserine and Bidi Kasserine, from the same bag**

484 The two populations collected in Kasserine, Bidi Kasserine and Ajimi Kasserine, were undifferentiated
485 from each other, genetically ($F_{st}=0.039$) and phenotypically for quantitative traits. Both of them shared
486 three MLGs, had exclusively the same heading dates, same parallel edged spikes and same spike
487 shapes. Some lineages from Ajimi Kasserine were shorter, with shorter and more compact spikes,
488 longer awns and lower number of spikelets per spike, albeit statistically not significant. In addition,
489 some lineages of Ajimi Kasserine had white colour of seeds and spikes and high anthocyanin
490 pigmentation of awns that was not observed in Bidi Kasserine. The populations Ajimi Kasserine and
491 Bidi-Kasserine were genetically close based on neighbour joining clustering (bootstrap support of 100,
492 Figure 4) and on their F_{st} index around 0.04. These results support the hypothesis of a common origin
493 for these two populations, considering that the NGBT collected seed samples from these two
494 populations from the same farmer in the region of Kasserine. Our findings are consistent with an
495 interview with the farmer suggesting a mishandling of seed bags that led to name two bags with
496 different variety names while they originated from the same landrace.

497 **-Mahmoudi Sejnane, a fake identity**

498 Mahmoudi Sejnane was highly differentiated from all the other 'Mahmoudi' populations (F_{st} : 0.53-0.8),
499 but was phenotypically identical to Mahmoudi Msaken. Both, Mahmoudi Sejnane and Mahmoudi
500 Msaken, were genetically homogeneous with lineages belonging to only one genetic group each,
501 respectively G4 and G6. These two populations were characterized by their stunted spike shape and
502 higher susceptibility to STB. The history of Mahmoudi landraces is unclear and several landraces may
503 have been wrongly attributed the variety name 'Mahmoudi' by farmers. In fact, it appears that the
504 population Mahmoudi Sejnane is genetically very close from the modern variety 'Karim' (bootstrap
505 88.4, Figure 4). A recent study by Slim & al. (2019), studying the same population Mahmoudi Sejnane
506 and the modern variety 'Karim' by neighbour joining cluster analysis, reached the same conclusion.
507 Thus, the population Mahmoudi Sejnane is suspected to be derived from the modern variety 'Karim'.
508 The short plant height of Mahmoudi Sejnane (and Mahmoudi Msaken) compared to other populations
509 and its high level of susceptibility to STB strongly looks alike the characteristics of the variety 'Karim',
510 which has been introduced in Tunisia more than 40 years ago by CYMMIT. It is by far the most
511 cultivated variety in Tunisia. Its resistance to STB has undergone years of genetic erosion making it
512 today very susceptible to *Z. tritici*.

513 **-Mahmoudi Msaken, a probable 'Chili' origin**

514 Mahmoudi Msaken was highly differentiated from all the other populations with variety name
515 'Mahmoudi'. The phylogenetic tree revealed that Mahmoudi Msaken is rather strongly related to a
516 landrace carrying the name 'Chili' (bootstrap support of 98.9), which do not group with other 'Chili'
517 landraces (Figure 4). However, the short length, stunted spike shape (Figure 6) and higher susceptibility
518 to STB looked alike the characteristics of Mahmoudi Sejnane, while these two populations were

519 genetically highly differentiated ($F_{st}=0.8$). This suggests that Mahmoudi Msaken is inherited from the
520 landrace Chili (PI534336, 1976) but might have evolved through mutations and crosses with other
521 varieties, such as the widespread variety Karim. This hypothesis could be investigated further through
522 the phenotypic characterization of the historical landrace.

523 **-Beskri Msaken, a 'Mahmoudi' on disguise**

524 Beskri Msaken was composed of lineages belonging to the same genetic groups (i.e. G1 and G5) as
525 other populations with the variety name 'Mahmoudi' (Figure 2). Moreover, Beskri Msaken was weakly
526 differentiated from the populations Mahmoudi El Jouv and Mahmoudi Joumine, these populations
527 sharing two MLGs. The close relationship between 'Beskri' and 'Mahmoudi' landraces was previously
528 established (Robbana & al., 2019; Slim & al., 2019). Based on Neighbour joining clustering, the
529 populations Beskri Msaken and Mahmoudi Joumine also appeared genetically close, even when the F_{st}
530 index was around 0.5. It suggests a common ancestor between Beskri Msaken and other Mahmoudi
531 populations.

532 **-'Mahmoudi', a popular variety with a complex history**

533 The four populations Mahmoudi El Jouv, Mahmoudi Oued Sbaihia, Mahmoudi Joumine and Mahmoudi
534 Amdoun had a heterogeneous genetic structure but all have lineages belonging to G1 and G5 (Figure
535 2). In addition, Mahmoudi Oued Sbaihia and Mahmoudi Joumine had lineages belonging to G3.
536 Mahmoudi Amdoun was genetically differentiated from the three other populations with variety name
537 'Mahmoudi' (F_{st} 0.247-0.346) but remained identical phenotypically. The genetic differentiation
538 between 'Mahmoudi' populations from El Jouv and Oued Sbaihia, and from El Jouv and Joumine, was
539 weak while they did differ phenotypically for quantitative traits. The complexity of the genetic
540 structure of 'Mahmoudi' populations emphasises the complexity to decipher and identify the
541 determinants of the life history, *i.e.* selection trajectory and seed exchanges between farmers, of those
542 landraces. In the literature, 'Mahmoudi' landraces were reported from Tunisia, Algeria, Libya or
543 Palestine (Deghais & al., 2007; Erroux, 1991; Boeuf, 1926; Gharbi & El Felah, 2013; Ammar & al., 2011).
544 In Tunisia, Boeuf (1926) reported seven indigenous botanical Mahmoudi varieties that were selected
545 and propagated by the Botanical Service of Tunis in the early 1900s. In the middle of the twentieth
546 century 'Mahmoudi' landraces became the most commonly cultivated landraces by traditional farmers
547 in Tunisia, even exclusively in certain locality such as Joumine. Farmers questioned during a survey
548 appeared very attached to this landrace for its yield stability, and high quality for making couscous and
549 traditional bread. Based on pairwise F_{st} analysis, Chili (PI534351, 1976) was almost genetically identical
550 to Mahmoudi Joumine ($F_{st}=0$), Mahmoudi El Jouv ($F_{st}=0$) and Mahmoudi Oued Sbaihia ($F_{st}=0.04$). These
551 three varieties named 'Mahmoudi' were also weakly differentiated from Chili Lansarine and Chili El
552 Jouv, suggesting that the landrace 'Chili' has contributed to the complex structure of 'Mahmoudi'
553 landraces in Tunisia. On the phylogenetic tree with historical collections of Tunisian landraces, these
554 three varieties named 'Mahmoudi' populations and Mahmoudi Amdoun, clustered with landraces
555 carrying the variety name 'Chili' rather than with landraces carrying the variety name 'Mahmoudi'
556 (Figure 4). It strengthens the hypothesis that seeds from 'Chili' have been introduced into 'Mahmoudi'
557 landraces. Also, the success of 'Mahmoudi' in Tunisia may explain why many seed lots (or populations)
558 have been called by traditional farmers with the name 'Mahmoudi' while having different origins,
559 resulting therefore in the complex genetic and phenotypic structure that we observed. Overall, our
560 findings emphasise the fact that the name of varieties is not sufficient to explain the origin and history
561 of Tunisian durum wheat landraces.

562

563

564 **3- Challenges and opportunities in the conservation of durum wheat landraces**

565 **-Contamination of durum wheat landraces with a hexaploid wheat**

566 Our study revealed the presence of hexaploid lineages in eight out of the 14 studied populations, easily
567 distinguishable in the farmers' fields because of their long cylindrical white spikes. They are considered
568 as impurities and result in a semolina with a flour-like texture with poorer quality. The origin of this
569 species in Tunisian durum wheat fields is not well understood. Some authors describe it as indigenous
570 and others as introduced by Europeans (Bœuf, 1926; Portères, 1958), while more recent studies
571 established that durum wheat fields grown with landraces are frequently contaminated with hexaploid
572 wheats (Zeven & Waninge, 1989; Figliuolo & al., 2007). In the middle of the twentieth century the
573 proportion of bread wheat in Tunisian durum wheat fields approached 50% (Bœuf, 1926; Portères,
574 1958), but this proportion is lower nowadays and was only 9% in our study. These bread wheats or
575 'mule's tail' existing as a mixture in the durum wheat fields are locally called « Baabous Bghal »
576 = « *Babous el brel* », « Dhil Bghal » or « Boujlida » = « *Bou jelida* » (in English: mule's tail), and Bœuf
577 and collaborators were unsuccessful in domesticating them through breeding (Bœuf, 1926; Gharbi &
578 Elfalah, 2013). Portères (1958) suggested that this form already existed before the arrival of Europeans,
579 and that durum wheat brought by Arabs in North Africa competed bread wheat due to its resistance
580 to drought and its better use for semolina. He further mentioned that the Romans, perhaps the
581 Phoenicians and Greeks, brought bread wheat to North Africa. Ducellier described in 1923 an Algerian
582 wheat named 'Hachadi' (white, dense, very rough ear; strongly curved, inflated glumes; strongly
583 spreading ears; strong divergent beards; reddish or reddish-brown seeds), which might be synonymous
584 to Tunisian 'mule's tail' (Laumont & Erroux, 1962). 'Hachadi' is probably an old form of cultivated
585 wheat, which is nowadays never found in its pure state but always mixed with traditional durum or
586 bread wheats. Nevertheless, these ancient forms have no cultural value anymore and are not
587 interesting for breeders. Similarly, Laumont & Erroux (1962) reported the presence of a hexaploid
588 wheat form in Algerian durum wheat fields, known as 'Guelia' (root akli: to cook, because of the
589 reddish colour of the grain which would have been scorched in the heat). This form of wheat has spread
590 in Algeria since 1950 as a result of the importation of wheat (Florence x Aurore) by Tunisian flour mills.
591 It looks like the « Baabous Bghal » or « Boujlida » (Laumont & Erroux, 1962), is morphologically
592 different from 'Hachadi' and characterised by rough, white ears, with a keeled husk similar to durum
593 wheat, and with a red grain often confused with durum wheat grains.

594 The presence of these undesirable hexaploid wheats in durum wheat fields obliges Tunisian farmers
595 to conduct a dynamic selection in order to eradicate 'mule's tail', unsuccessfully so far. Farmers who
596 are not undergoing this selection process are gradually losing the purity of their landrace, which can
597 lead to a loss of its diversity and sometimes to its complete loss due to resignation.

598 **-Divergence among landraces is explained by selection rather than genetic drift**

599 The majority of evaluated quantitative traits were highly divergent ($P_{st} > F_{st}$), with a robust
600 approximation of P_{st} . Especially, HI, HS, AUDPC, NSS and LSWA were distinguishable with low critical
601 $c/h^2 < 0.20$. Indeed, we can suppose that divergence in genes coding for these five traits exceeds what
602 is expected on the basis of genetic drift. It indicates that selection has resulted in different phenotypes
603 for these five traits between the studied populations, which is the footprint of local adaptation.
604 Selection made by farmers on their landraces should have contributed to this local adaptation. Indeed,
605 a local survey informed us that some farmers perform irregular selection based especially on spike
606 length, spike density and awn colour. Thus, the genetic variation present in the landraces for these
607 traits constitute an important resource to breed for improved varieties adapted to local constraints
608 and meeting Tunisian farmers' expectations.

609 -Use of most resistant lineages to STB for conservation and breeding purposes

610 Landraces are a well-known source of resistance to diseases (Akem & al. 2000; Nazco & al., 2012; Xu
611 & al., 2018; Agnoun & al., 2019; Yao & al., 2019), such as durum wheat landraces and their resistance
612 to *Septoria tritici* blotch (STB) caused by *Z. tritici*. Previous studies by Ferjaoui & al. (2011, 2015)
613 identified for instance new resistance genes to STB in 'Azizi27', 'Agili37', 'Agili39' and 'Derbessi12'
614 landraces. Ouaja & al. (2020) also identified valuable sources of resistance among a collection of 304
615 Tunisian durum wheat landraces. In our study, the 14 populations responded differently to STB (Figure
616 5). Roussia Joumine and Ajimi Kasserine were the most resistant populations to the *Z. tritici* strain used
617 for inoculations, IPO91009. The resistance is qualitative (total resistance controlled by major genes) in
618 both populations suggesting the presence of highly effective resistance genes that could be useful for
619 breeding. Bidi Kasserine appeared slightly less resistant to STB than Ajimi Kasserine, although the two
620 populations are genetically and phenotypically identical. On the contrary Mahmoudi Sejnane was
621 highly susceptible to STB, suggesting that it does not carry any source of resistance to this disease or
622 that the resistance genes it may contain have been overcome by *Z. tritici* populations. Intermediary,
623 the remaining 10 populations expressed varying levels of resistance, probably quantitative (partial
624 resistance controlled by polygenes with moderate-to-small effects).

625 Intra-population variation of the level of resistance between lineages was observed as well (Figure 5),
626 reflecting the genetic diversity previously described within the populations. Few lineages from Roussia
627 Joumine and Ajimi Kasserine had some level of susceptibility. Inversely, several populations expressing
628 quantitative resistance contained few completely resistant lineages, such as Mahmoudi Amdoun and
629 Mahmoudi Oued Sbaihia. Moreover, all populations expressing quantitative resistance as a whole were
630 composed of lineages with varying levels of resistance, which was especially pronounced for
631 Mahmoudi Msaken. This intra-population diversity represents as well an important resource of genetic
632 variation, which should not be overseen.

633

634 Conclusion

635 Our study showed a complex structuration of 14 durum wheat populations, only partly explained by
636 their geographic origin and name of variety. Among them, landraces called 'Chili' contributed a lot to
637 the history of Tunisian durum wheat landraces. These results highlight that a landrace is the outcome
638 of a complex selection trajectory, driven not only by the heritage of a biological material (seed lots)
639 but also by traditional breeding practices (selection) deeply-rooted in a territory and subject to several
640 kinds of disruptions (exchange, mixing, loss, ingression of exogeneous material, misidentification, etc.).
641 These two components are an integral part of the "life history" of landraces. The dynamic nature of
642 this life story must be considered and characterized, even more thoroughly than we have done in this
643 study, for instance with sociologists, if one wants to better exploit the resources that landraces
644 represent. It is also crucial to enlarge prospection and collection efforts of actual cultivated landraces
645 to retrace the history of these genetic resources and, particularly, distinguishing the "real" landraces
646 from the "false" by combining genetic and phenotypic approaches. The conservation of durum wheat
647 landraces is important for future breeding programs. And the role of gene banks is prominent to
648 develop appropriate and relevant *in situ* and *ex situ* conservation plans.

649

650

651

652 **Materials and Methods**

653 **Plant material**

654 Since 2012 the National Gene Bank of Tunisia (NGBT) conducts a program aiming at the conservation
655 of landraces from traditional Tunisian crops, such as durum wheat. Farmers still cultivating those
656 landraces were identified and seed samples were collected for conservation purposes. Within this
657 framework, we focused on 14 populations, 10 being collected in 2015 from five localities and four
658 being collected in 2017 from three additional localities (Table 8). Each population was considered as
659 representative of a given durum wheat landrace and was designated by a variety name combined with
660 its locality of origin. Therefore, several of these “variety-locality” populations collected in different
661 “localities” have the same “variety” name, and vice-versa. Seeds of each population were randomly
662 selected from the seed samples and sown under open field conditions in micro-plots (2 m x 1 m) at the
663 experimental station of Thiverval-Grignon, France (48°50'21.3" N, 1°57'07.2" E). For each population,
664 32 plants were randomly selected and labelled from 1 to 32, except for the four populations collected
665 in 2017 for which 16, 23, 38 and 51 plants were selected, leading to a total of 448 plants. At the
666 inflorescence emergence stage (Zadoks' 50-59), the wheat spikes were bagged in permeable
667 cellophane wraps in cellophane to promote self-fertilization and prevent the expression of
668 heterozygosity (< 5% in cultivated wheat). When matured (Zadoks' 90-99), spikes from each plant were
669 collected and stored in separate bags before being threshed, spike by spike. This procedure was
670 repeated twice for the 10 populations collected in 2015 and only once for the four populations
671 collected in 2017. As each cycle of self-fertilization is expected to reduce natural intercrossing by 50%,
672 the residual heterozygosity was estimated less than 1.25%. Each lineage obtained from the 32 plants
673 originally selected and represented by seeds coming from a single spike were sown as a single head-
674 row during the growing season 2017-2018 on the CRP-Wheat Septoria Phenotyping Platform from
675 CIMMYT-IRESA (Kodia Bou Salem, Tunisia). This trial was conducted as randomized incomplete block
676 design (12 blocks of 16 m x 1 m separated by the modern variety 'Karim'). Eight reference varieties
677 were included in each block: 'Karim', 'Khiar', 'Maali', 'Nasr', 'Salim', 'Agili-39' and 'Mahmoudi-101'. At
678 the end of the trial, six spikes per wheat lineage (i.e. head-row) were harvested, threshed, and stored
679 for further characterizations.

680 **Molecular analyses**

681 *DNA isolation and genotyping

682 For each wheat lineage, one seed was sown and grown under controlled conditions (growth chamber)
683 with a 16h/24h light (300 photons $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$) and a temperature of 18°C night/20°C day. Two weeks
684 after sowing, a 2-3 cm segment from the youngest leaf of each plant was sampled for DNA extraction
685 performed using the DNA plant MiniKit protocole (www.qiagen.com). Purity and concentration of the
686 extracted DNAs were estimated using a Nanodrop spectrophotometer (ND-1000). The DNA samples
687 were normalized to a concentration of 20 ng. μL^{-1} and genotyped using nine SSR markers (Table 9).
688 Genotyping was outsourced at Eurofins (www.eurofins.fr). These SSR markers were assembled in a
689 single multiplex following the methodology described by Gautier & al. (2014). Briefly, the forward
690 primers were labelled with four fluorochromes (i.e. Ned, Fam, Pet, Vic) in a way that the same colour
691 was given only to markers with non-overlapping range of allele sizes. Candidate markers were
692 individually tested on eight reference durum wheat DNAs, before constitution and validation of the
693 multiplex. Chromatograms were visually inspected for all markers and for all individuals using Peak
694 scanner software version 1.0, before final assignment of SSR alleles. Individuals with missing data for
695 at least one marker were removed from our dataset not to influence the detection of unique multilocus
696 genotypes (MLGs) per population. Finally, genotypes of 335 lineages were obtained (13 to 32 lineages

697 per population), after the exclusion of hexaploid lineages (see below, section “Determination of the
698 ploidy level in lineages”).

699 *Marker characterization

700 Neutrality of the SSR markers was preliminary confirmed using BayeScan v2.1 software, which allows
701 identifying candidate loci under natural selection from genetic data using differences in allele
702 frequencies between populations. The default settings were used (Foll & Gaggiotti, 2008). The plotting
703 and identification of outliers was performed with R software using the output of the MCMC algorithm.

704 A genotype accumulation curve was drawn with the package poppr in Rstudio v3.5.2 to determine the
705 minimum number of loci necessary to discriminate between individuals in a population. This function
706 randomly sampled the loci without replacement and counted the number of multilocus genotypes
707 observed.

708 Estimates of number of alleles (N_a), number of private alleles (N_{ap}), mean observed heterozygosity (H_o),
709 mean genetic diversity (H_s) and estimate of F_{is} following Nei (1987) were obtained using the package
710 hierfstat of Rstudio v.5.2.

711 The Polymorphism Information Content (PIC) of each microsatellite locus was evaluated using the
712 formula of Botstein & al. (1980), with n corresponding to the number of alleles, p_i to the frequency of
713 the i^{th} allele, and p_j to the frequency of the j^{th} allele:

$$714 \quad PIC = 1 - \sum_{i=1}^n p_i^2 - \sum_{i=1}^{n-1} \sum_{j=i+1}^n 2p_i^2 p_j^2$$

715 Markers with PIC value greater than 0.5 were considered highly informative (Botstein & al., 1980).

716 *Population structure, molecular variance and diversity analyses

717 Population structure was analysed on MLGs with the Bayesian model-based clustering software
718 STRUCTURE v.2.3.4 (Pritchard & al., 2000). Simulations were performed under an admixture model
719 with independent allele frequencies. The analysis was carried out for a number of tested clusters (K)
720 ranging from 1 to 20, 10 runs, 100,000 burn-in period and 100,000 Markov Chain Monte Carlo (MCMC)
721 repetitions after burning. The most likely number of populations (K) was identified using the Delta K
722 method implemented in Structure Harvester. Structure results were summarized using Excel to obtain
723 the probability of each lineage to belong to each cluster. Lineages of the 14 populations were assigned
724 to one genetic cluster, or several genetic clusters when genotypes were admixed. A lineage was
725 assigned to a genetic group if > 50% of its genome fraction value was derived from that group.

726 A Minimum Spanning Network on MLGs based on Nei distances with 1000 bootstraps and neighbor
727 joining clustering method was performed using the package poppr (kamvar & al., 2014) of R Studio
728 V3.5.2.

729 To assess the partitioning of the total genetic variance within and among populations, an analysis of
730 molecular variance (AMOVA) based on F -statistics with 999 permutations was performed using
731 GenAlEx 6.5 (Peakall & Smouse, 2012).

732 The genetic distance between populations was assessed by calculating pairwise F_{st} using the software
733 Arlequin v3.5.2.2 (Excoffier & Lischer, 2010).

734 Diversity measures incorporate both genotypic richness R (number of genotypes in a population;
735 Dorken & Eckert, 2001) and the Evenness index E_5 (distribution of genotypes within a population;

736 Grünwald & al., 2003). Common diversity indices were calculated, i.e. Shannon Weaver (H), Stoddart
737 and Taylor's index (G) and Simpson's index (λ). Finally, several genetic diversity parameters,
738 including the number of alleles per population (N_a), the observed heterozygosity (H_o), the expected
739 heterozygosity (H_e), the number of multilocus genotypes (MLGs), the number of expected MLG at the
740 smallest sample size ≥ 10 based on rarefaction (eMLGs), and Weir and Cockerham estimates of F
741 statistics over all loci (F_{st}), were estimated using the software Genclone v 2.0 (Arnaud-Haond & Belkhir,
742 2007), the packages poppr (Kamvar & al., 2014) and Hierfstat (Goudet, 2005) of Rstudio v3.5.2.

743 *Phylogenetic analysis

744 Several international gene banks have accessions from durum wheat landraces collected through the
745 20th century in their collections. We ordered from the U.S. National Plant Germplasm System (NPGS)
746 of the USDA (<https://npgsweb.ars-grin.gov/>) seed samples corresponding to 39 accessions from durum
747 wheat landraces carrying the names 'Bidi', 'Chili' and 'Mahmoudi' from Tunisia, Algeria, Morocco and
748 Italy (Table-ESM2). A 40th accession, called 'Mahmoudi-101', was provided by the NGBT. Four
749 individuals of each of the 40 accessions were genotyped with the nine SSR markers as described
750 previously. In addition, three different seed lots were collected from the modern variety 'Karim'
751 provided by the NGBT, the CRP-Wheat Septoria Phenotyping Platform, and the National Agronomic
752 Institute of Tunisia (INAT). 30, 30 and 12 individuals from each respective lot were genotyped. Then, a
753 phylogenetic analysis was conducted integrating genotypes of the 14 studied populations, the 40
754 accessions from landraces and the three seed lots from 'Karim'. A phylogenetic tree based on
755 Neighbour Joining clustering method on Edward's distances, with 1000 bootstraps, was generated
756 using the package poppr of Rstudio V3.5.2.

757 **Evaluation of phenotypic characters:**

758 *Determination of the ploidy level in lineages

759 Preliminary molecular and phenotypic analyses revealed lineages within eight populations belonging
760 to the same genetic groups and being phenotypically more similar to hexaploid wheat species rather
761 than tetraploid durum wheat. The ploidy level of 14 lineages, of the landrace Mahmoudi-101 and of
762 variety Karim was verified by establishing their karyotype. The seeds were sterilized 10 min with
763 sodium hypochlorite 5% and then rinsed with water. They were placed for 48 hours at 4°C in the dark
764 for germination on a petri dish containing a water-moistened filter paper and then moved at 22°C in
765 an oven for 48 hours. Roots of 2 cm were cut and placed in a glass tube containing a solution of ultra-
766 pure water previously cooled for 24 hours on ice to stop mitosis. The roots were then fixed in an
767 ethanol/acetic acid solution (3:1) at 4°C for 24 hours. The roots were placed in 1 mL of acetocarmine
768 (carmin 10 g/L with 45% acetic acid) in a watch glass for 1 hour. The tip of a meristem is placed on a
769 slide with a drop of 45% acetic acid, covered with a coverslip and then taped gently with the tip of a
770 pencil to release the cells. The slide is lightly heated for a few seconds at 90 °C to remove the
771 cytoplasmic veil. The quality of the spreads chromosomes is observed under a conventional phase
772 contrast microscope. The photos are taken in bright-field at X40 magnification. Chromosome numbers
773 are counted for five cells to estimate the hexaploid/tetraploid nature of the plants.

774 All hexaploid lineages were removed from further analyses on genetic and phenotypic diversity in the
775 14 populations.

776 *Agro-morphological characterization of landraces

777 Four representative spikes were randomly sampled at maturity from each lineage sown as a headrow
778 in the Kodia Bou Salem's field trial. Fifteen qualitative and quantitative agro-morphological traits were

779 measured on the plants in the field or on the sampled spikes: plant height (HI), heading date (HS), spike
780 shape (SS), spike colour (SC), spike density (SD), length of awns in relation to spike (LAS), length of spike
781 without awns (LSWA), length of awns (LA), awn colour (AC), anthocyanin pigmentation of awns (PgA),
782 number of spikelets per spike (NSS), colour of grains (CG), shape of grains (SG), and thousand grain
783 weight (TGW). Scoring was done according to the recommendations of the International Plant Genetic
784 Resource Institute (IPGRI) and the International Union for the Protection of New Varieties of Plants
785 (UPOV) wheat descriptor lists (Table ESM3). Some lineages were lost during the field trial leaving us
786 with 273 lineages for which a complete genotypic and phenotypic data set was available.

787 *Evaluation of landraces for resistance to *Septoria tritici* blotch

788 During the trial conducted on the CRP-Wheat Septoria Phenotyping Platform at Kodia Bou Salem,
789 lineages from landraces and reference varieties were evaluated for their resistance to *Septoria tritici*
790 blotch (STB). In the field, spreader rows of the modern variety 'Karim', highly deployed in Tunisia and
791 considered susceptible (Berraies & al., 2014), were sown in all blocks perpendicularly to head-rows.
792 Spreader rows and head-rows were spray-inoculated with the *Zymoseptoria tritici* strain IPO91009,
793 collected in Tunisia (Béja) in 1991 (Kema & al., 1996). The inoculation was performed twice on 15 and
794 29 January 2018, between Zadoks' 13 and (three fully unfolded leaves) and Zadoks' 26 (tillering) stages
795 with a conidial suspension adjusted to the concentration 10^6 spores.mL⁻¹. STB severity was assessed at
796 two different dates following the "double digit" scoring method (Saari & Prescott, 1975). The area
797 under the disease progression curve (AUDPC) was calculated from the percentage of disease severity
798 (DS) at both observation dates (Sharma & Duveiller, 2007; Das & al., 1992) according to the formula:

799
$$DS = \left(\frac{D1}{9}\right) \times \left(\frac{D2}{9}\right) \times 100$$

800 where, D1 is the first digit (vertical disease progress) and D2 is the second digit (severity of infection),
801 and

802
$$AUDPC = \sum_{I=1}^{n-1} ((DS_{i+1} + DS_i) \times (T_{i+1} - T_i))/2$$

803 where, DS_i is disease severity at the ith assessment, T_i is the time (number of days) on which the ith
804 assessment was performed, and n is the total number of assessments.

805 Statistical analyses were performed to assess differences between populations for their
806 resistance/susceptibility to STB at p-value=0.05. Kruskal-Wallis tests were applied and Mann-Whitney
807 tests were performed using Rstudio v 3.5.2.

808 *Statistical analysis of quantitative traits

809 A matrix of correlations between the nine quantitative traits was calculated at confidence interval of
810 0.99 using the package corrplot of Rstudio v 3.5.2. One-way Anova and Kruskal-Wallis tests -
811 parametric and non-parametric tests, respectively - were applied depending on the trait's distribution
812 to test for phenotypic differences between populations. For the non-normally distributed traits, a first
813 normalization was performed using the boxcox function (Box & Cox, 1964) from the package MASS
814 (Venables & Ripley, 2002). In case a significant difference was detected for a trait, post-hoc tests were
815 performed using Rstudio v 3.5.2: a Tukey's Honest Significant Difference test (*i.e.* parametric) or a
816 Mann-Whitney test (*i.e.* non-parametric), at p-value=0.001.

817 *Phenotypic diversity between populations and genetic groups

818 For qualitative phenotypic traits, we considered the most frequent phenotypic class among the four
819 spikes of each lineage. For quantitative traits, the mean values of the four spikes were categorized into
820 classes (Table ESM3). All qualitative and quantitative values grouped into classes were used to
821 calculate the Shannon–Weaver diversity index (H) (Shannon & Weaver 1949; Jain & al. 1975) according
822 to the formula:

$$823 \quad H = - \sum_{i=1}^n p_i \log_2 p_i$$

824 where p_i is the frequency of individuals from the i^{th} class and n is the number of classes for the
825 designated phenotypic trait.

826 A relative phenotypic diversity index (H') was calculated according to the formula:

$$827 \quad H' = H/H_{\max} \quad \text{with} \quad H_{\max} = \log_e(n)$$

828

829 *Factor Analysis of Mixed Data (FAMD)

830 A Factor Analysis of Mixed Data (FAMD) was performed using the FactoMineR and factoextra packages
831 in Rstudio v3.5.2, using both quantitative and qualitative variables to ensure a balance of their
832 influence in the analysis. This method allows to reveal similarities between lineages and to explore
833 associations between all variables.

834 **Relationship between genotypic and phenotypic diversity parameters**

835 A P_{st} - F_{st} analysis was performed using the 273 lineages for which both phenotypic and genetic data
836 were available to test whether phenotypic differences in nine quantitative traits between populations
837 were due to selection or genetic drift. P_{st} was used as an approximation of Q_{st} , whose exact calculation
838 requires common garden experiments to measure the additive genetic variances (Leinonen & al., 2008;
839 Pujol & al., 2008; Brommer, 2011) and was not possible here.

840 The genetic differentiation F_{st} and the corresponding 95% confidence interval (CI) were calculated
841 using the package hierfstat. P_{st} was estimated according to Brommer (2011) according to the formula:
842

$$843 \quad P_{st} = \frac{\frac{c}{h^2} \sigma_b^2}{\frac{c}{h^2} \sigma_b^2 + 2 \sigma_w^2}$$

844 where σ_b^2 and σ_w^2 are the phenotypic variances between and within populations, respectively, c is an
845 estimate of the proportion of the total variance due to additive genetic effects across populations, and
846 h^2 is the heritability (the proportion of phenotypic variance due to additive genetic effects). When
847 $P_{st}=F_{st}$, the differentiation of quantitative traits may be the result of genetic drift, even if a contribution
848 of natural selection cannot be discarded nor estimated. If $P_{st}>F_{st}$, quantitative traits have a higher level
849 of differentiation, which could be an evidence for directional selection (or heterogeneous selection).
850 Finally, if $P_{st}<F_{st}$, quantitative traits are less diversified than neutral differentiation, suggesting that
851 these traits have been under the influence of stabilizing selection (or homogeneous selection).

852 As the determination of c is difficult, a sensitive analysis was performed using the package Pstat of R
853 studio v3.5.2 to infer the robustness of the estimation of c/h^2 which is critical for P_{st} to correctly
854 approximate Q_{st} . The lower critical c/h^2 ratio is obtained when P_{st} exceeds F_{st} and at null assumption
855 ($c/h^2 = 1$) the proportion of phenotypic variance due to additive genetic effects is the same within and
856 across populations. The trait is considered divergent at the point where P_{st} exceeds F_{st} ($c<h^2$). P_{st} values

857 were extracted at null assumption and P_{st} with a 95% CI were estimated using Pstat package with 1000
858 permutations. If the lower 95% CI of P_{st} is higher than F_{st} then the quantitative trait is considered highly
859 divergent. For each trait with $P_{st} > F_{st}$, the robustness of the P_{st} as an approximation of F_{st} , indicating
860 local adaptation, was estimated by the critical c/h^2 value. This ratio is referring to the value when the
861 lower 95% CI of P_{st} equal the upper 95% CI of F_{st} as described in Brommer & al. (2011). As the critical
862 c/h^2 was here lower than 0.20 for some traits, comparing P_{st} and F_{st} allowed us to make robust
863 conclusions on the selection regime (Brommer, 2011). The critical c/h^2 ratio was calculated according
864 to the formula:

$$865 \quad \sigma_w^2 (lower) = \frac{[1 - P_{st}(lower) c/h^2]}{2 \times P_{st}(lower)},$$

867 Then :

$$868 \quad c/h^2 (critical) = \frac{[2 \times \sigma_w^2 (lower) F_{st}(upper)]}{1 - F_{st}(upper)}$$

869

870

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881

882 Competing interests

883 The authors declare that they have no competing interests.

884

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- 1193

1194 **Figures**

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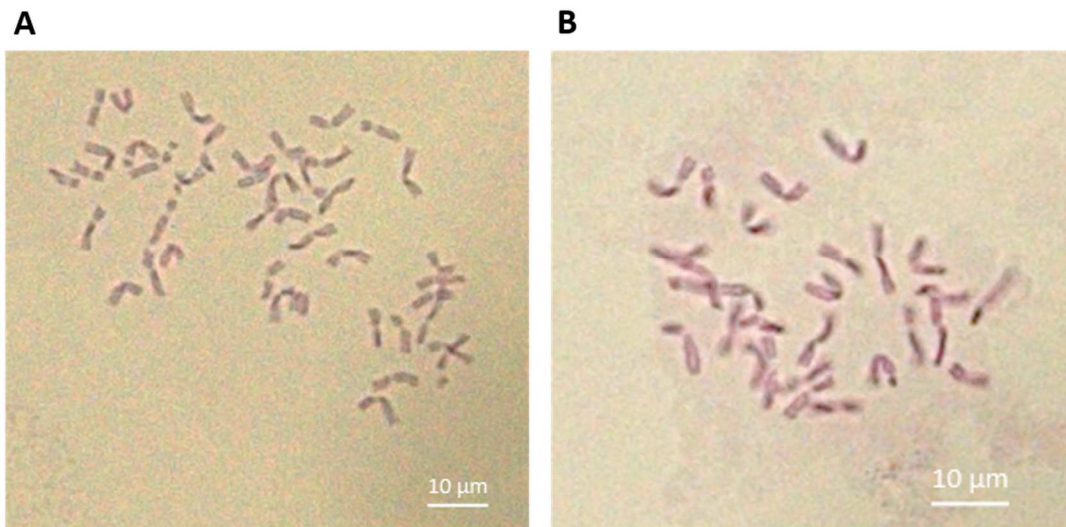
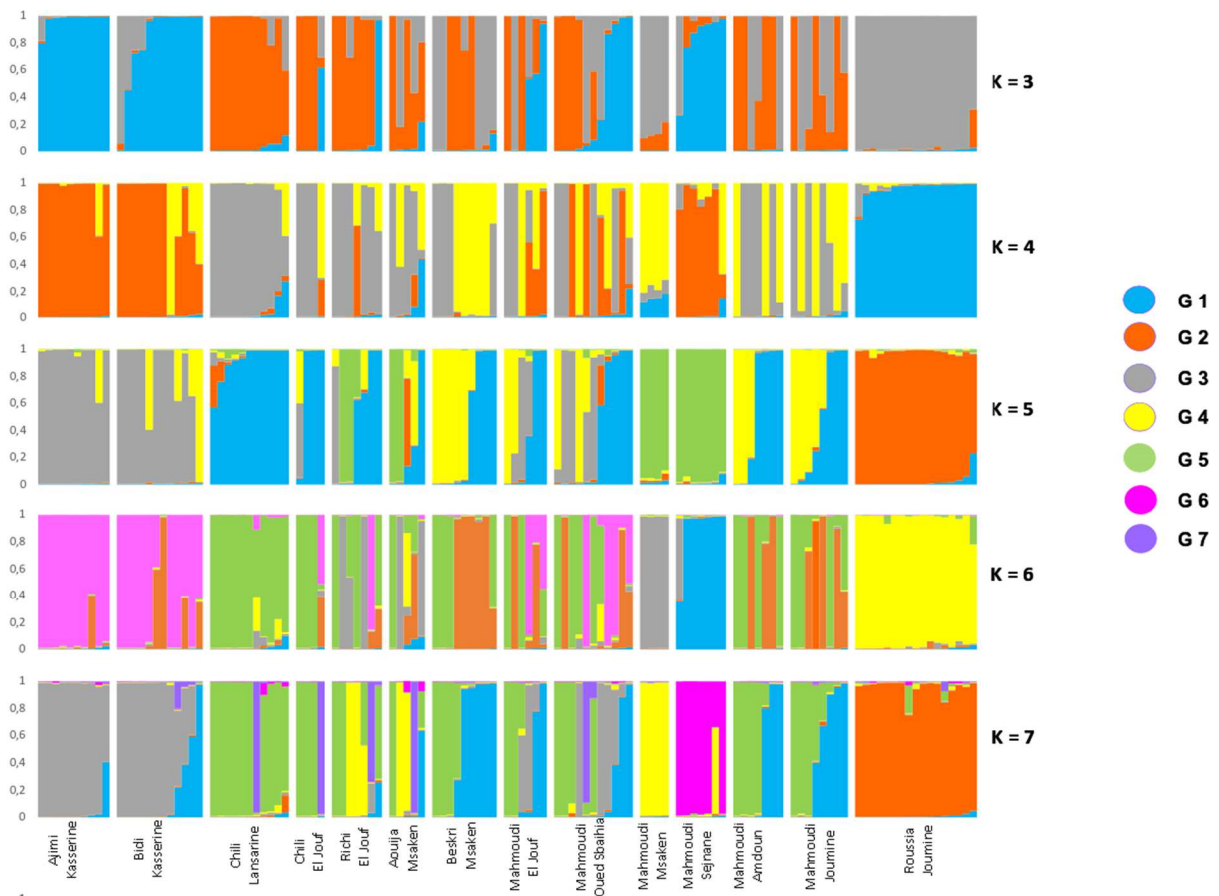


Figure 1. Karyotypes of A. a 'mule's tail' lineage carrying 42 chromosomes (hexaploid), and B. the durum wheat cultivar Karim carrying 28 chromosomes (tetraploid).

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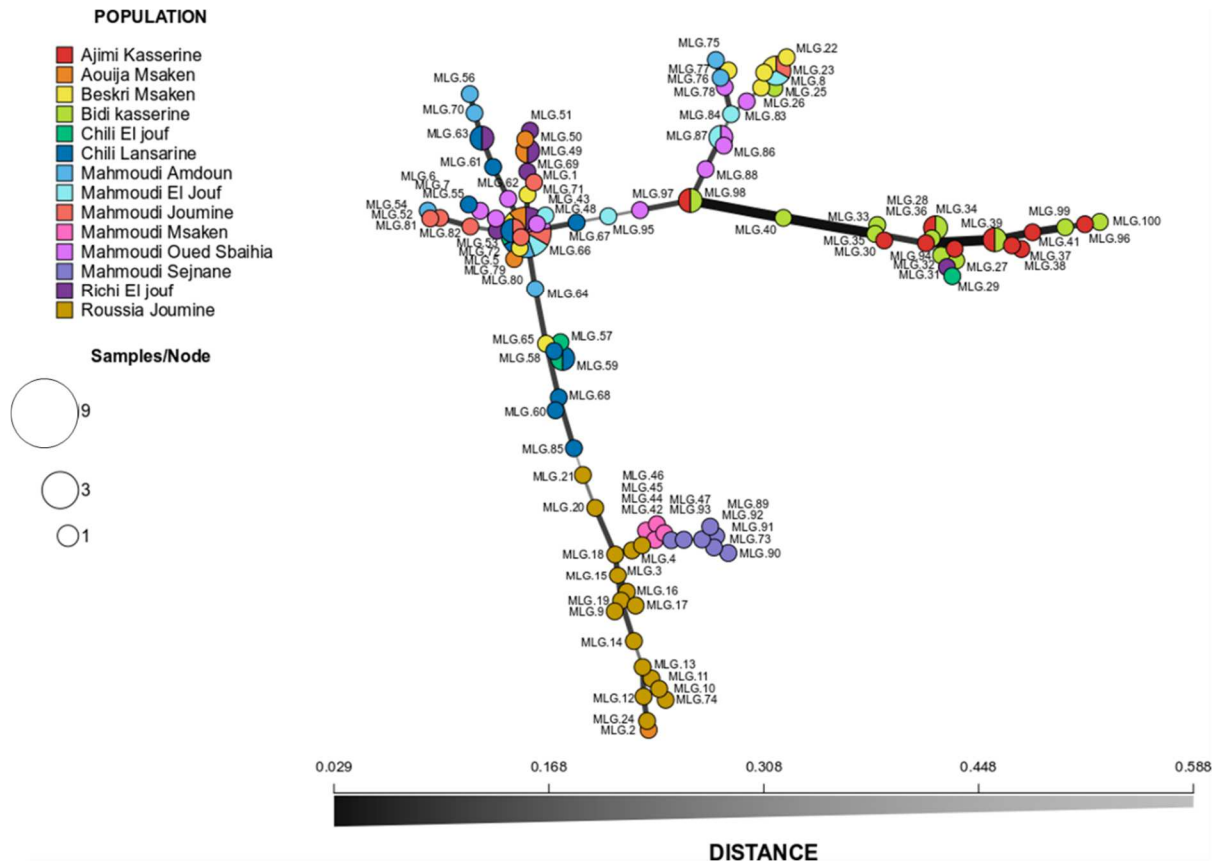
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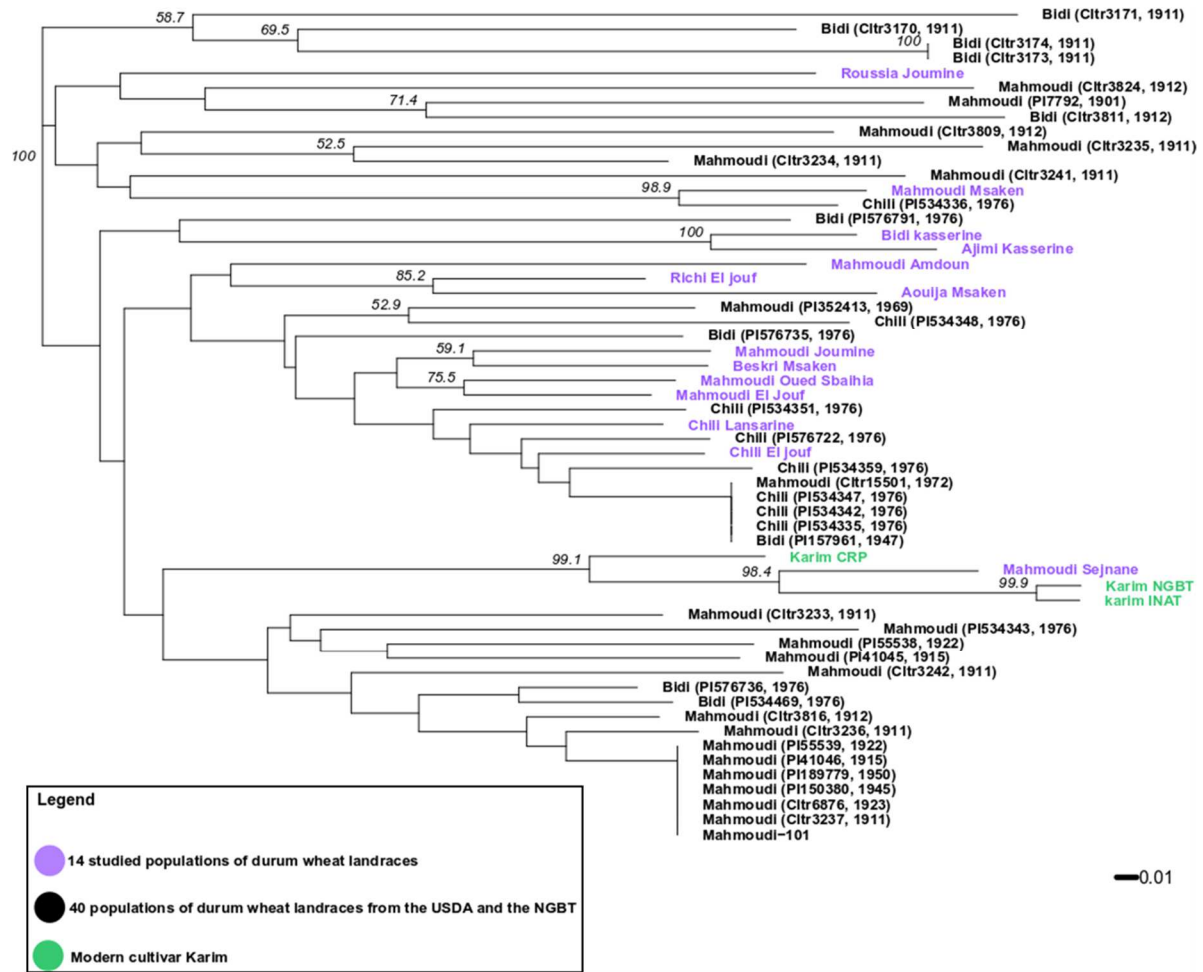
Figure 2. Admixture proportions of the 14 durum wheat populations estimated with STRUCTURE (K=3 to K=7) leading to the identification of different genetic groups (G1 to G7 on the left of the figure). Each vertical bar represents an individual. The colour proportion within each bar represents the posterior probability of assignment of each individual to one of the groups of genetic similarity. The range of assignment probability varied from 0 to 100%.



1205

1206 **Figure 3. Minimum spanning network between multilocus genotypes (MLGs) of the 14 durum wheat**
1207 **populations.** Each node represents a different MLG with the size proportional to the frequency of the
1208 MLG. Colours represent the population of origin of each MLG. Edges represent minimum genetic
1209 distances between MLGs based on Nei distances (1000 bootstraps) and Neighbor-Joining clustering
1210 method. Nodes that are more closely related have thicker and darker edges, whereas nodes that are
1211 more distantly related have lighter and thinner edges.

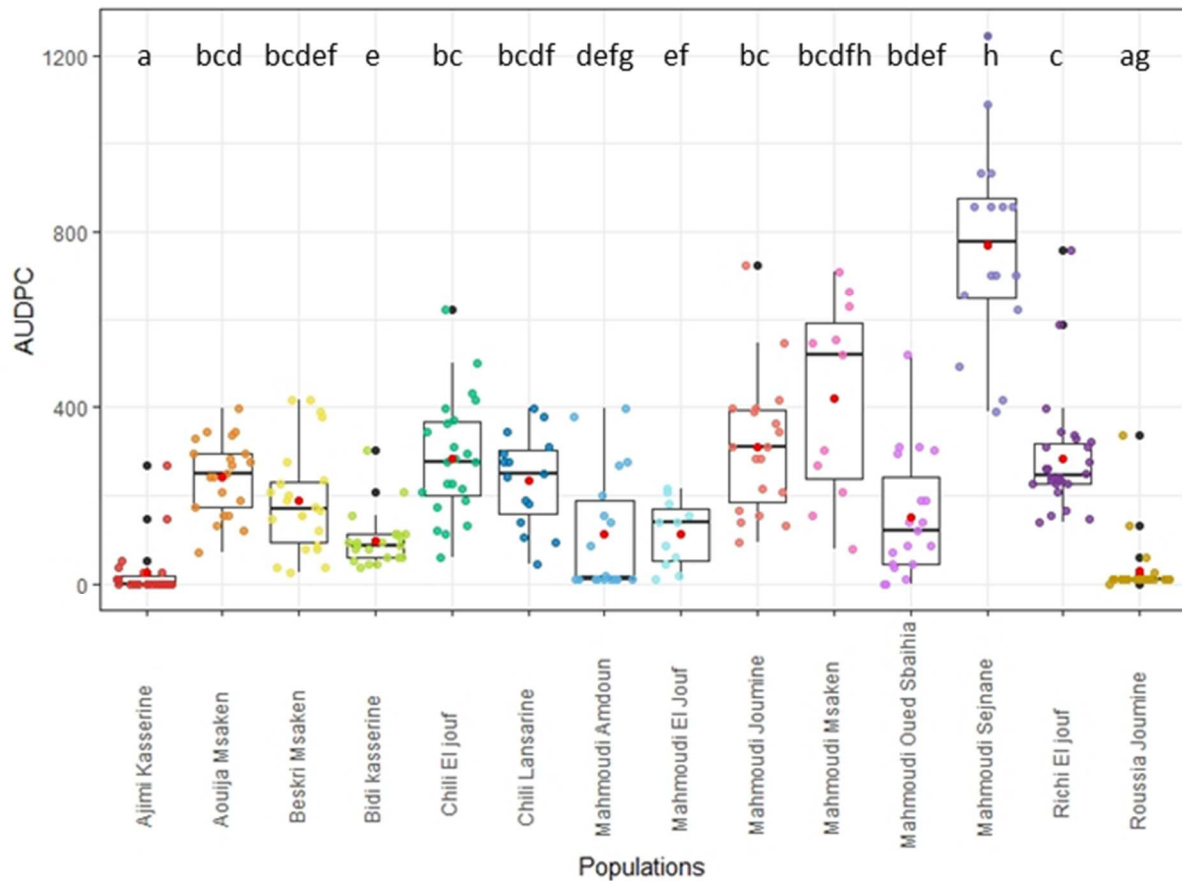
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1214 **Figure 4. Phylogenetic tree with the 14 studied durum wheat populations (in violet), the 40 landraces**
 1215 **collected from the USDA and the NGBT (in black) and three seed lots of the modern cultivar Karim**
 1216 **(in green), resulting from Neighbor-Joining cluster analysis based on Edward's distances (1000**
 1217 **bootstraps). Bootstrap support values expressed in percentages are indicated on the nodes only if**
 1218 **there are > 50%.**

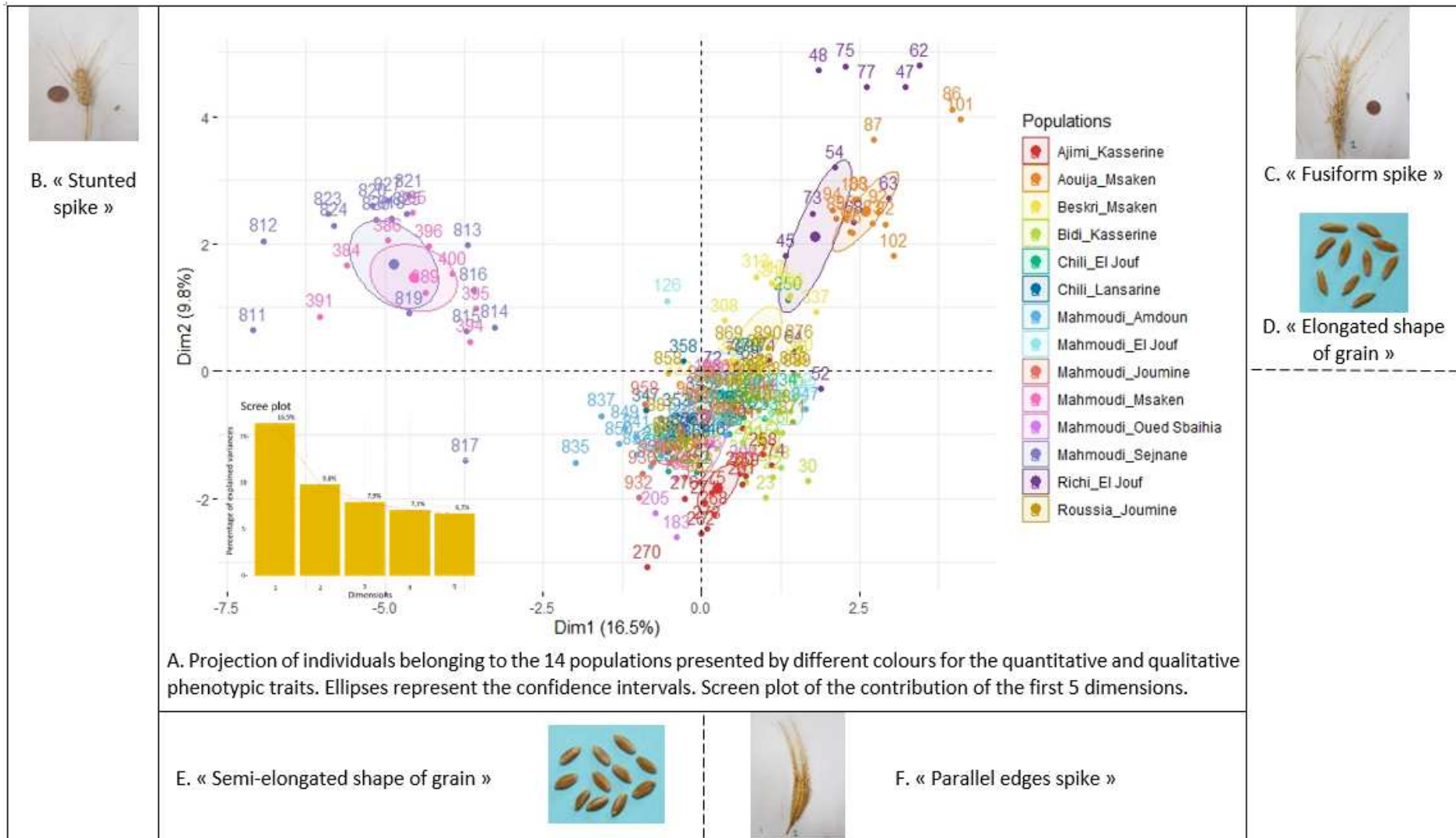
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1221 **Figure 5. *Septoria tritici* blotch (STB) AUDPC (area under the disease progress curve) for the 14 durum**
1222 **wheat populations obtained after their inoculation in field conditions at Kodia Bou Salem with the**
1223 ***Z. tritici* strain IPO91009. Means are represented by a red point. Populations with significantly**
1224 **different means are indicated by different letters after Kruskal-Wallis and Mann–Whitney tests at**
1225 **$p=0.05$.**

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1228 **Figure 6. A: Factor Analysis of Mixed Data (FAMD) of 15 quantitative and qualitative phenotypic traits measured in the 14 durum wheat populations. B-F:**
 1229 **Pictures of corresponding spike and grain shapes.**

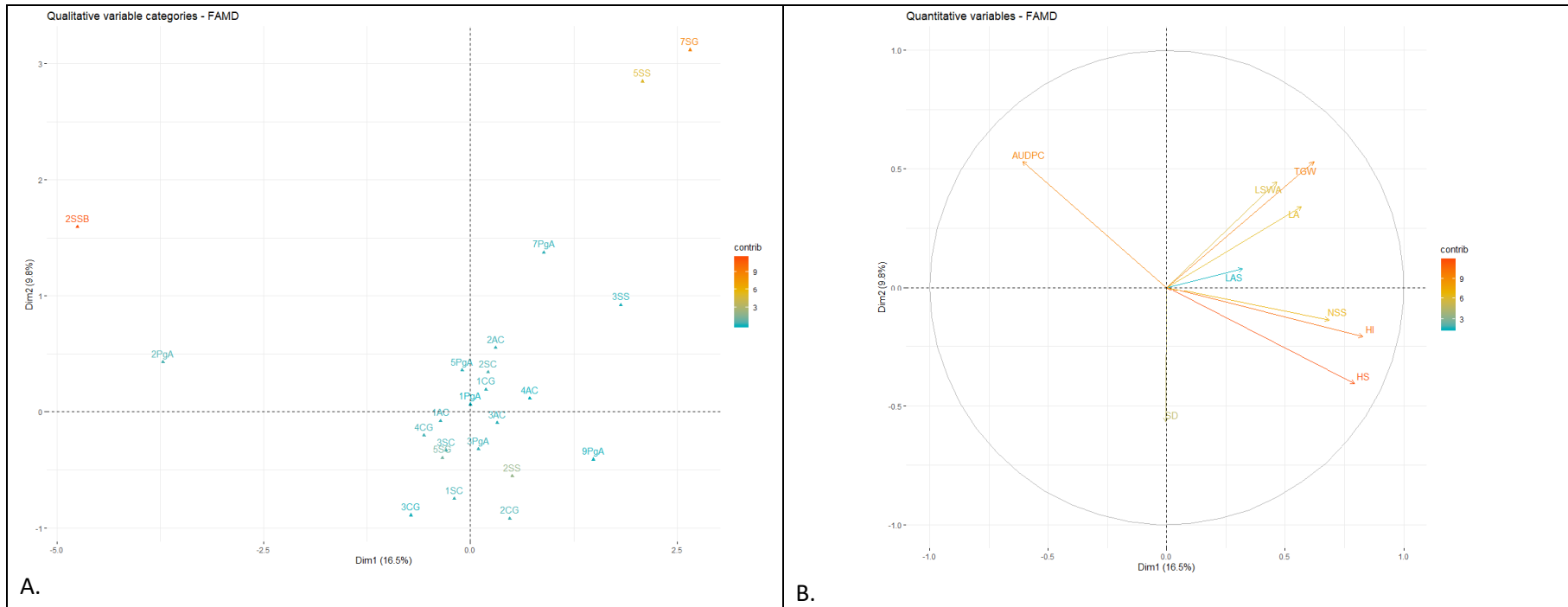
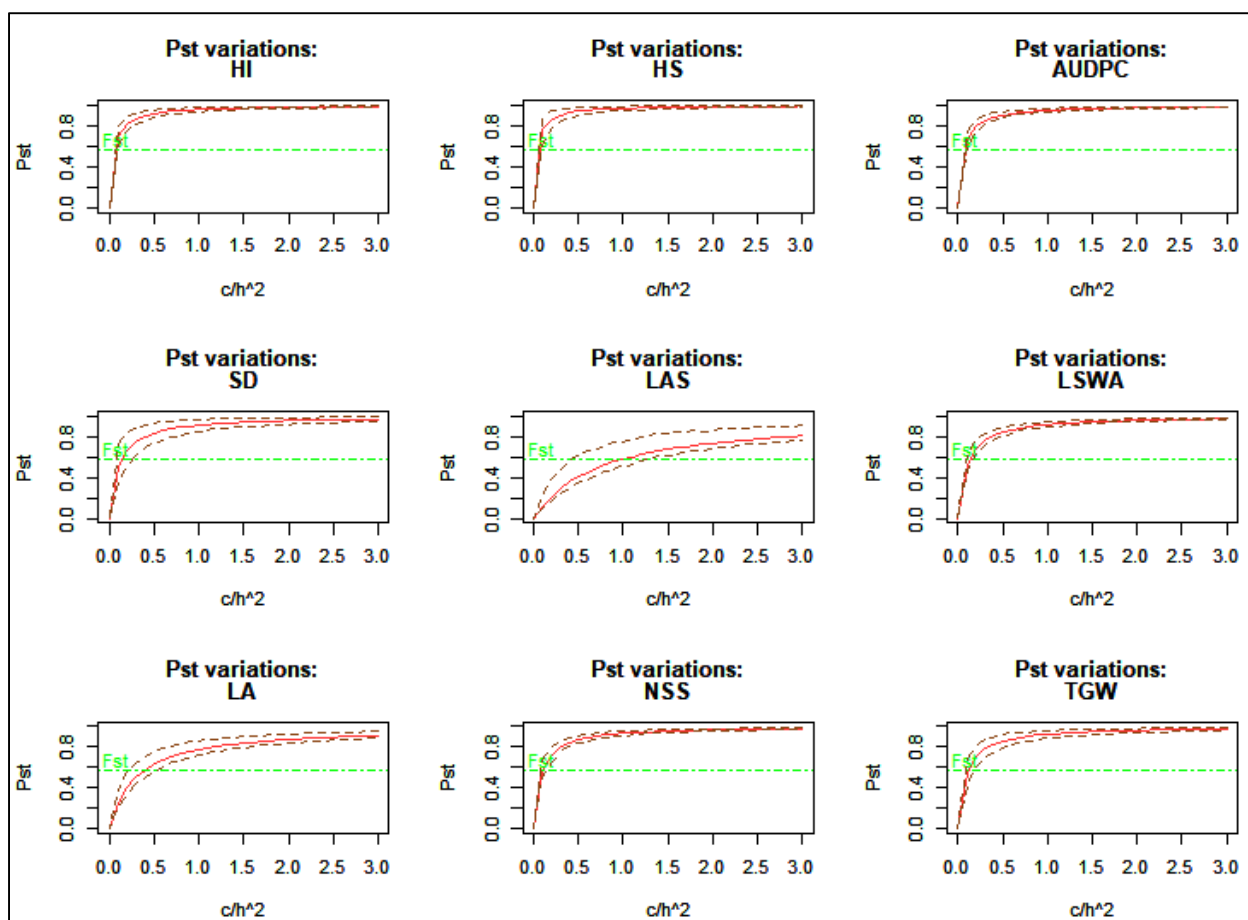


Figure 7. Output of the Factor Analysis of Mixed Data (FAMD). A. Categorical variable factor map projects the classes of qualitative variables in the plane of 1 and 2 dimensions underlining their contribution. Classes of Spike Shape: 1SS, 2SS, 2SSB, 3SS, 5SS; classes of Spike Colour: 1SC, 2SC, 3SC; classes of Awn Colour: 1AC, 2AC, 3AC, 4AC; classes of Anthocyanin colouration of Awns: 1PgA, 3PgA, 5PgA, 7PgA, 9PgA; classes of Colour of Grain: 1CG, 2CG, 3CG, 4CG; classes of Shape of Grain: 1SS, 2SS, 3SS. **B. Correlation circle represents the projection of quantitative variables on the 1 and 2 dimensions underlining their contribution.** HI: plant height; HS: heading date; AUDPC: area under disease progress curve; SD: spike density; LAS: length of awns in relation to spike; LSWA: length of spike without awns; LA: length of awns; NSS: number of spikelets per spike; TGW: thousand grains weight.

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1233 **Figure 8. Comparison of phenotypic differentiation of the 14 durum wheat populations (P_{st} : red solid**
1234 **line; dotted red lines represents the 95% CI). The neutral genetic differentiation (F_{st} - dotted green line**
1235 **for $F_{st}=0.57098$ was calculated using the package hierfstat with Rstudio version 3.5.2), while the ratio**
1236 **c/h^2 ranged from 0 to 3 where c represents the proportion of the total variance and h^2 the heritability.**
1237 **The lowest value of c/h^2 for which P_{st} exceeds F_{st} = critical value of c/h^2 can be considered an indication**
1238 **of the robustness of using P_{st} as an alternative for Q_{st} (Brommer, 2011).**

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1240 **Tables**

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1242 **Table 1: Polymorphism of the 9 SSR (microsatellite) markers used to characterize the 14 durum wheat populations:** number
 1243 of alleles (N_a), number of private alleles (N_{ap}), mean observed heterozygosity (H_o), mean expected heterozygosity (H_s), fixation
 1244 index (F_{is}) following Nei (1987) expected heterozygosity over all (H_e), and Polymorphism Information Content (PIC) values.

| | Xgpw2103 | Xgpw2239 | Xgpw4082 | Xgpw7148 | Xgwm193 | Xgwm285 | Xgwm372 | Xgwm4004 | Xgwm413 |
|----------------------------|----------|----------|----------|----------|---------|---------|---------|----------|---------|
| N_a | 4 | 3 | 6 | 6 | 9 | 8 | 12 | 5 | 8 |
| N_{ap} | 2 | 0 | 3 | 2 | 5 | 3 | 4 | 1 | 3 |
| H_o | 0.005 | 0.005 | 0.003 | 0.008 | 0.006 | 0.014 | 0.007 | 0.005 | 0.012 |
| H_s | 0.147 | 0.224 | 0.198 | 0.267 | 0.192 | 0.353 | 0.323 | 0.313 | 0.317 |
| F_{is} | 0.965 | 0.977 | 0.986 | 0.969 | 0.971 | 0.961 | 0.977 | 0.984 | 0.962 |
| H_e | 0.396 | 0.561 | 0.524 | 0.413 | 0.528 | 0.691 | 0.643 | 0.665 | 0.738 |
| PIC | 0.322 | 0.423 | 0.466 | 0.387 | 0.492 | 0.639 | 0.618 | 0.600 | 0.704 |

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1247 **Table 2: Analysis of Molecular Variance (AMOVA) based on SSR (microsatellite) markers and using the F_{st}**
1248 **measure, for 335 individuals belonging to 14 durum wheat populations.**

| Source | df | SS | MS | Estimated variance | Variance (%) |
|----------------------------|-----|----------|--------|--------------------|--------------|
| Between populations | 13 | 934.965 | 71.920 | 1.459 | 54% |
| Within populations | 321 | 782.721 | 2.438 | 1.204 | 45% |
| Within individuals | 335 | 10.500 | 0.031 | 0.031 | 1% |
| Total | 669 | 1728.187 | | 2.694 | 100% |

df: degree of freedom; SS: sums of squares; MS: mean square; variance (%): percentage of total variance contributed by each component.

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1250 **Table 3: Genetic diversity of the 14 durum wheat populations as evaluated from 9 SSR (microsatellite) markers**

| Populations | Number of lineages ¹ | N _a ² | H _s ³ | H _o ⁴ | Number of MLG ⁵ | Number of eMLG ⁶ | R ⁷ | Shannon index H ⁸ | Stoddart and Taylor's index (G) ⁹ | Simpson's index LAMDA ¹⁰ | Eveness E _s ¹¹ |
|--------------------------------------|---------------------------------|-----------------------------|-----------------------------|-----------------------------|----------------------------|-----------------------------|----------------|------------------------------|--|-------------------------------------|--------------------------------------|
| Ajimi Kasserine | 32 | 2.333 | 0.138 | 0.010 | 10 | 5.19 | 0.29 | 1.466 | 2.60 | 0.615 | 0.480 |
| Aouija Msaken | 28 | 2.556 | 0.159 | 0 | 5 | 2.86 | 0.148 | 0.608 | 1.35 | 0.260 | 0.420 |
| Beskri Msaken | 26 | 2.333 | 0.382 | 0 | 9 | 5.48 | 0.320 | 1.528 | 2.96 | 0.663 | 0.545 |
| Bidi kasserine | 25 | 2.667 | 0.268 | 0.036 | 12 | 7.36 | 0.458 | 1.977 | 4.37 | 0.771 | 0.542 |
| Chili El jouf | 29 | 2.111 | 0.096 | 0 | 4 | 2.60 | 0.107 | 0.545 | 1.33 | 0.250 | 0.460 |
| Chili Lansarine | 28 | 2.444 | 0.185 | 0 | 11 | 6.93 | 0.370 | 1.894 | 4.04 | 0.753 | 0.539 |
| Mahmoudi Amdoun | 21 | 2.444 | 0.367 | 0 | 7 | 5.30 | 0.300 | 1.402 | 2.74 | 0.635 | 0.568 |
| Mahmoudi El Jouf | 13 | 2.444 | 0.332 | 0.009 | 6 | 6.00 | 0.416 | 1.285 | 2.45 | 0.592 | 0.554 |
| Mahmoudi Joumine | 19 | 2.333 | 0.285 | 0 | 8 | 6.25 | 0.388 | 1.587 | 3.19 | 0.687 | 0.565 |
| Mahmoudi Msaken | 15 | 1.333 | 0.057 | 0 | 4 | 3.72 | 0.214 | 0.857 | 1.77 | 0.436 | 0.569 |
| Mahmoudi Oued Sbaihia | 22 | 2.889 | 0.407 | 0.015 | 11 | 7.26 | 0.476 | 1.895 | 4.10 | 0.756 | 0.549 |
| Mahmoudi Sejnane | 16 | 2.111 | 0.267 | 0 | 7 | 6.22 | 0.400 | 1.667 | 4.27 | 0.766 | 0.760 |
| Richi El jouf | 31 | 3.000 | 0.348 | 0.022 | 7 | 4.10 | 0.200 | 1.283 | 2.80 | 0.643 | 0.691 |
| Roussia Joumine | 30 | 3.111 | 0.342 | 0 | 17 | 9.61 | 0.551 | 2.625 | 11.25 | 0.911 | 0.801 |
| ¹² F _{st} =0.572 | | | | | | | | | | | |

1: Number of genotyped lineages by population after eliminating hexaploid individuals and genotypes containing missing value for at least one marker. 2: Mean average of alleles. 3: Mean expected heterozygosity. 4: Mean observed heterozygosity. 5: Number of MultiLocus Genotypes. 6: Number of expected MLG at the smallest sample size based on rarefaction. 7: Genotypic Richness (Dorken & Eckert, 2001). 8: Shannon-Wiener Index of MLG diversity (Arnaud-Haond & al., 2007; Shannon 2001). 9: Stoddart and Taylor's index of MLG diversity (Stoddart & Taylor, 1988). 10: Simpson's Index (Simpson, 1949). 11: Evenness, E_s (Pielou, 1975; Grünwald & al., 2003). 12: F_{st} over all loci (Weir & Cockrham, 1984).

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Table 4: Matrix of F_{st} between the 14 durum wheat populations

| | Ajimi Kasserine | Aouija Msaken | Beskri Msaken | Bidi Kasserine | Chili El Jouf | Chili Lansarine | Mahmoudi Amdoun | Mahmoudi El Jouf | Mahmoudi Joumine | Mahmoudi Msaken | Mahmoudi Oued Sbaihia | Mahmoudi Sejnane | Richi El Jouf |
|--------------------------|--------------------|------------------|------------------|-------------------|------------------|--------------------|--------------------|---------------------|---------------------|---------------------|-----------------------------|---------------------|------------------|
| Aouija Msaken | 0.808* | | | | | | | | | | | | |
| Beskri Msaken | 0.688* | 0.570* | | | | | | | | | | | |
| Bidi kasserine | 0.039 | 0.733* | 0.579* | | | | | | | | | | |
| Chili El Jouf | 0.845* | 0.778* | 0.236* | 0.765* | | | | | | | | | |
| Chili Lansarine | 0.788* | 0.705* | 0.192* | 0.699* | 0.035* | | | | | | | | |
| Mahmoudi Amdoun | 0.679* | 0.590* | 0.343* | 0.576* | 0.489* | 0.409* | | | | | | | |
| Mahmoudi El Jouf | 0.722* | 0.618* | 0.047 | 0.591* | 0.149* | 0.088* | 0.315* | | | | | | |
| Mahmoudi Joumine | 0.758* | 0.656* | 0.053 | 0.650* | 0.139* | 0.111 | 0.346* | 0.033 | | | | | |
| Mahmoudi Msaken | 0.887* | 0.757* | 0.579* | 0.804* | 0.862* | 0.768* | 0.703* | 0.698* | 0.704* | | | | |
| Mahmoudi Oued Sbaihia | 0.614* | 0.541* | 0.112* | 0.490* | 0.208* | 0.140* | 0.247* | 0.016 | 0.108 | 0.615* | | | |
| Mahmoudi Sejnane | 0.737* | 0.731* | 0.559* | 0.638* | 0.772* | 0.689* | 0.587* | 0.586* | 0.641* | ² 0.800* | 0.527* | | |
| Richi El Jouf | 0.680* | 0.270* | 0.233* | 0.591* | 0.354* | 0.287* | 0.315* | 0.188* | 0.253* | 0.575* | 0.178* | 0.570* | |
| Roussia Joumine | 0.707* | 0.680* | 0.506* | 0.606* | 0.724* | 0.667* | 0.612* | 0.559* | 0.577* | 0.645* | 0.525* | 0.647* | 0.569* |

*Significance of the F_{st} at a threshold of 5%.

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Table 5. Estimates of Shannon–Weaver Diversity Index H' , mean H' and standard error ($\pm SE$) of each of the 14 populations and seven genetic groups¹ based on the evaluation of lineages with 15 phenotypic traits.

| | HI | HS | AUDPC | SS | SC | SD | LAS | LSWA | LA | AC | PgA | NSS | CG | SG | TGW | H' mean $\pm SE$ | |
|-----------------------------|-----------------------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|--------------------|---------------------------------|
| Populations | H' | | | | | | | | | | | | | | | | |
| | Bidi Kasserine | 0.75 | 0.00 | 0.17 | 0.00 | 0.57 | 0.69 | 0.00 | 0.72 | 0.37 | 0.69 | 1.13 | 0.83 | 0.40 | 0.00 | 0.49 | 0.45\pm0.09 |
| | Richi El Jouf | 0.48 | 0.00 | 0.83 | 0.50 | 1.02 | 0.81 | 0.21 | 0.71 | 0.74 | 0.75 | 1.11 | 0.71 | 0.17 | 0.91 | 0.90 | 0.66\pm0.08 |
| | Aouija Msaken | 0.17 | 0.00 | 0.62 | 0.44 | 0.84 | 0.70 | 0.00 | 0.57 | 0.61 | 1.09 | 0.78 | 0.84 | 0.00 | 0.25 | 0.65 | 0.50\pm0.09 |
| | Mahmoudi El Jouf | 0.63 | 0.48 | 0.30 | 0.00 | 1.24 | 0.53 | 0.00 | 0.68 | 0.68 | 0.66 | 0.92 | 0.82 | 0.00 | 0.00 | 0.74 | 0.51\pm0.10 |
| | Mahmoudi Oued Sbaihia | 0.70 | 0.00 | 0.66 | 0.17 | 1.34 | 0.61 | 0.00 | 0.73 | 0.46 | 1.25 | 0.79 | 0.68 | 0.55 | 0.00 | 0.91 | 0.59\pm0.11 |
| | Chili El Jouf | 0.57 | 0.00 | 0.82 | 0.14 | 0.90 | 0.47 | 0.00 | 0.47 | 0.79 | 0.74 | 0.72 | 0.41 | 0.70 | 0.00 | 0.69 | 0.50\pm0.08 |
| | Ajimi Kasserine | 0.74 | 0.00 | 0.09 | 0.00 | 1.15 | 0.26 | 0.00 | 0.41 | 0.79 | 0.84 | 0.88 | 0.58 | 0.54 | 0.00 | 0.29 | 0.44\pm0.10 |
| | Beskri Msaken | 0.75 | 0.00 | 0.46 | 0.72 | 1.13 | 0.56 | 0.00 | 0.62 | 0.76 | 0.76 | 0.78 | 1.06 | 0.68 | 0.00 | 0.75 | 0.60\pm0.09 |
| | Chili Lansarine | 0.62 | 0.00 | 0.66 | 0.00 | 1.25 | 0.35 | 0.00 | 0.43 | 0.65 | 1.27 | 0.71 | 0.80 | 0.65 | 0.00 | 0.76 | 0.54\pm0.11 |
| | Mahmoudi Msaken | 0.52 | 0.94 | 0.77 | 0.00 | 0.84 | 0.26 | 0.00 | 0.00 | 0.96 | 0.32 | 0.26 | 0.57 | 0.32 | 0.00 | 0.45 | 0.41\pm0.09 |
| | Mahmoudi Sejnane | 0.53 | 1.38 | 1.08 | 0.00 | 0.82 | 0.61 | 0.49 | 0.61 | 0.78 | 0.39 | 0.87 | 0.21 | 0.00 | 0.00 | 0.35 | 0.54\pm0.10 |
| | Mahmoudi Amdoun | 0.84 | 0.00 | 0.53 | 0.00 | 1.35 | 0.71 | 0.28 | 0.00 | 0.83 | 0.74 | 0.93 | 0.92 | 1.13 | 0.00 | 0.99 | 0.62\pm0.12 |
| | Roussia Joumine | 0.90 | 0.00 | 0.14 | 0.00 | 1.08 | 0.51 | 0.35 | 0.74 | 0.86 | 0.00 | 0.84 | 0.88 | 1.13 | 0.00 | 0.71 | 0.54\pm0.11 |
| | Mahmoudi Joumine | 0.65 | 0.00 | 0.85 | 0.00 | 1.30 | 0.39 | 0.57 | 0.98 | 0.92 | 0.93 | 0.79 | 0.46 | 0.98 | 0.00 | 0.85 | 0.64\pm0.10 |
| | mean H' | 0.63 | 0.20 | 0.57 | 0.14 | 1.06 | 0.53 | 0.14 | 0.55 | 0.73 | 0.75 | 0.82 | 0.70 | 0.52 | 0.08 | 0.68 | |
| Genetic groups ¹ | H' | | | | | | | | | | | | | | | | |
| | G1 | 0.60 | 0.00 | 0.60 | 0.36 | 1.43 | 0.70 | 0.48 | 0.89 | 0.89 | 1.07 | 0.72 | 0.92 | 1.06 | 0.22 | 0.58 | 0.70\pm0.09 |
| | G2 | 0.93 | 0.00 | 0.14 | 0.00 | 1.08 | 0.51 | 0.38 | 0.74 | 0.97 | 0.00 | 0.84 | 0.88 | 1.13 | 0.00 | 0.29 | 0.53\pm0.11 |
| | G3 | 0.82 | 0.00 | 0.15 | 0.00 | 1.07 | 0.42 | 0.00 | 0.60 | 0.65 | 1.12 | 1.14 | 0.86 | 0.39 | 0.00 | 0.58 | 0.52\pm0.11 |
| | G4 | 0.80 | 0.49 | 0.93 | 0.87 | 0.99 | 0.93 | 0.14 | 0.95 | 0.93 | 1.12 | 0.95 | 1.28 | 0.55 | 0.81 | 1.03 | 0.85\pm0.07 |
| | G5 | 0.80 | 0.05 | 0.75 | 0.27 | 1.13 | 0.72 | 0.00 | 0.44 | 0.72 | 1.06 | 1.00 | 0.97 | 0.80 | 0.00 | 0.97 | 0.64\pm0.10 |
| | G6 | 0.37 | 0.58 | 1.14 | 0.00 | 0.86 | 0.62 | 0.54 | 0.58 | 0.81 | 0.43 | 0.88 | 0.23 | 0.00 | 0.00 | 0.00 | 0.47\pm0.09 |

| | | | | | | | | | | | | | | | | |
|---------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|------------------|
| G7 | 0.00 | 0.00 | 0.31 | 0.00 | 0.46 | 0.62 | 0.00 | 0.62 | 0.62 | 0.00 | 0.00 | 0.00 | 0.00 | 0.91 | 0.62 | 0.28±0.09 |
| mean H' | 0.62 | 0.16 | 0.58 | 0.21 | 1.00 | 0.65 | 0.22 | 0.69 | 0.80 | 0.68 | 0.79 | 0.73 | 0.56 | 0.28 | 0.58 | |

¹ Genetic groups defined by STRUCTURE software at K=7.

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Table 6. Pairwise populations significantly different quantitative traits¹ at $p < 0.001$

| | Bidi Kasserine | Richi El Jouv | Aouija Msaken | Mahmoudi El Jouv | Mahmoudi Oued Sbaihia | Chili El Jouv | Ajimi Kasserine | Beskri Msaken | Chili Lansarine | Mahmoudi Msaken | Mahmoudi Sejnane | Mahmoudi Amdoun | Roussia Joumine |
|------------------------------|--------------------|----------------------|----------------------------|------------------|------------------------|-----------------|------------------------|-------------------------|-----------------|----------------------|----------------------|-----------------|-----------------|
| Richi El Jouv | SD, LSWA | | | | | | | | | | | | |
| Aouija Msaken | SD, LSWA | | | | | | | | | | | | |
| Mahmoudi El Jouv | | | SD | | | | | | | | | | |
| Mahmoudi Oued Sbaihia | NSS | SD | SD, LA, LSWA | | | | | | | | | | |
| Chili El Jouv | NSS | SD | SD, LSWA | | | | | | | | | | |
| Ajimi Kasserine | | AUDPC, SD, LSWA, TGW | AUDPC, SD, LSWA, TGW | | | AUDPC | | | | | | | |
| Beskri Msaken | NSS | | | | | | | AUDPC, SD, LSWA | | | | | |
| Chili Lansarine | NSS | NSS, SD | NSS, SD | | | NSS | | NSS, AUDPC | | | | | |
| Mahmoudi Msaken | NSS, HS | NSS, HI, HS, LSWA | NSS, HI, HS, LSWA | NSS | NSS, HI, HS | NSS, HI, HS | NSS, HS, AUDPC, SD | NSS, HS | NSS | | | | |
| Mahmoudi Sejnane | NSS, HS, AUDPC, SD | NSS, HI, HS, TGW | NSS, HI, HS, LA, LSWA, TGW | NSS, HI, AUDPC | NSS, HI, HS, AUDPC, SD | NSS, HI, HS, SD | NSS, HI, HS, AUDPC, SD | NSS, HI, HS, AUDPC, TGW | HI, HS | | | | |
| Mahmoudi Amdoun | NSS | TGW | NSS, HI, LA, TGW | | | HI | NSS, SD | | | HS, NSS | HS, AUDPC | | |
| Roussia Joumine | NSS, SD, LSWA | AUDPC, TGW | HI, AUDPC, TGW | | SD, LSWA | AUDPC, SD, LSWA | SD, LSWA | | AUDPC | NSS, HS, AUDPC, LSWA | NSS, HS, AUDPC, LSWA | | |
| Mahmoudi Joumine | NSS | | LA, LSWA | | | | AUDPC, SD | | | NSS, HS | NSS, HS, TGW | | AUDPC, LA, LSWA |

Colour gradient: cells are getting darker as the number of phenotypic traits significantly different between the two populations is increasing. ¹ Abbreviations: Table ESM3

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1258 **Table 7. P_{st} - F_{st} comparison for quantitative traits measured in 273 lineages belonging to the 14 durum wheat populations.**

1259 Lower values of the critical c/h^2 indicate a more robust inference of local adaptation.

| Traits ^μ | P_{st} at $c/h^2 = 1$ [§] (null assumption) | F_{st} | 95% CI lower | 95% CI upper | Critical c/h^2 [£] |
|---------------------|---|----------|--------------|--------------|-------------------------------|
| | | 0.57098 | 0.5218 | 0.6180 | |
| HI | 0.9580 | | 0.9353 | 0.9760 | 0.1118 |
| HS | 0.9692 | | 0.9417 | 0.9881 | 0.1001 |
| AUDPC | 0.9506 | | 0.9319 | 0.9680 | 0.1182 |
| SD | 0.9073 | | 0.8432 | 0.9625 | 0.3008 |
| LAS | 0.5784 | | 0.5066 | 0.7535 | |
| LSWA | 0.9123 | | 0.8904 | 0.9399 | 0.1991 |
| LA | 0.7732 | | 0.7252 | 0.8585 | 0.6129 |
| NSS | 0.9299 | | 0.9064 | 0.9537 | 0.1670 |
| TGW | 0.9220 | | 0.8813 | 0.9551 | 0.2179 |

^μ Abbreviations: Table ESM3

[§] $c/h^2 = 1$ is the null assumption meaning that the proportion of phenotypic variance caused by additive genetic effects is the same for between-population and within-population variances (Brommer, 2011)

[£] critical c/h^2 ratio was calculated according to the formula (Brommer, 2011).

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1263 **Table 8. List of the 14 durum wheat populations: locality, suspected origin/year and year of sampling.**

| Populations | Locality-Gouvernorate | Suspected origin/year of landrace* | Year of sampling |
|-----------------------|------------------------------|---|-------------------------|
| Ajimi Kasserine | Kasserine-Kasserine | Unavailable | 2015 |
| Aouija Msaken | Msaken -Sousse | North Africa, 1909 | 2015 |
| Beskri Msaken | Msaken-Sousse | Algeria, 1909 | 2015 |
| Bidi Kasserine | Kasserine-Kasserine | North Africa, 1908 | 2015 |
| Chili El Jouv | El Jouv -Zaghouan | France, 1932 | 2015 |
| Chili Lansarine | Lansarine-Manouba | France, 1932 | 2015 |
| Mahmoudi Amdoun | Amdoun-Beja | Tunisia, 1983 | 2017 |
| Mahmoudi El Jouv | El Jouv -Zaghouan | Tunisia, 1983 | 2015 |
| Mahmoudi Joumine | Joumine-Bizerte | Tunisia, 1983 | 2017 |
| Mahmoudi Msaken | Msaken-Sousse | Tunisia, 1983 | 2015 |
| Mahmoudi Oued Sbaihia | Oued Sbaihia -Zaghouan | Tunisia, 1983 | 2015 |
| Mahmoudi Sejnane | Sejnane-Bizerte | Tunisia, 1983 | 2017 |
| Richi El Jouv | El Jouv-Zaghouan | Tibar, Tunisia 1908/1909 | 2015 |
| Roussia Joumine | Joumine-Bizerte | Bizerte, Tunisia, 1927 | 2017 |

*Deghaïs & al. 2003, 2007; Ammar & al. 2011

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Table 9. Description of the panel of 9 SSR (microsatellite) markers used for population genetics.

| Markers | Chromosomes | Size Range (bp) ¹ | Dye colour | Motif | Primers F | Primers R | Tm ² | Reference |
|----------|-------------|------------------------------|------------|---|----------------------|-----------------------|-----------------|-----------------------|
| Xgwm413 | 2B | 88-110 | Ned | (GA) ₂₀ | TGCTTGCTAGATTGCTTGGG | GATCGTCTCGTCCTTGCA | 60°C | Röder & al., 1998 |
| Xgpw7148 | 3B | 87-100 | Fam | (AG) ₂₁ | GCACACAACGACACTTGCTT | GCTTAGCTGCTTGCTTTGTG | 60°C | Sourdille & al., 2004 |
| Xgwm193 | 6BS | 155-192 | Ned | (CT) _{21imp} (CA) ₈ | CTTTGTGCACCTCTCTCTCC | AATTGTGTTGATGATTTGGGG | 60°C | Korzun & al., 1997 |
| Xgpw2239 | 4AS | 191-197 | Vic | (CT) ₁₉ | CAACCATATGCCAGGAGAC | TGTTGCTGTCTGAAACAGGG | 60°C | Sourdille & al., 2004 |
| Xgwm285 | 3B | 209-235 | Ned | (GA) ₁₉ | ATGACCCTTCTGCCAAACAC | ATCGACCGGGATCTAGCC | 60°C | Somers & al., 2004 |
| Xgpw4082 | 4B | 211-235 | Fam | (TG) ₂₃ (GA) _{16.5} | CTTTCTTCCCCTCCTGTCC | ATCATCACAATGCAGCGAG | 60°C | Sourdille & al., 2004 |
| Xgpw4004 | 5A | 209-256 | Pet | Unavailable | CGCCTCGGATTCTATTCTTG | CTTACTGCGGCCTTGAGTTG | 60°C | GrainGenes |
| Xgpw2103 | 7AL (7B) | 230-308 | Vic | (TC) ₁₆ | CGTATGCAGCATGGCATC | GCTATGTTGTGTGGCATTGG | 60°C | Sourdille & al., 2004 |
| Xgwm372 | 2A | 296-331 | Fam | (GA) ₂₆ | AATAGAGCCCTGGGACTGGG | GAAGGACGACATTCCACCTG | 60°C | Röder & al., 1998 |

¹Size range (bp: base pairs) for each marker corresponds to the minimum and maximum size identified in all the durum wheat populations; ²Tm: melting temperature.