

1 **Assesment of phenology and morphological diversity of 3 species of Asteraceae:**
2 ***Anacyclus clavatus*, *Chamaemelum fuscatum* and *Leucanthemum parthenium***

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22 **Abstract**

23 **Aim:** Three species of Asteraceae: *Anacyclus clavatus*, *Chamaemelum fuscatum* and *Leucanthemum*
24 *parthenium* that have a wide range of uses in medicine and in industry were characterized by inter-
25 specific variations and phenological activities.

26 **Study Design:** Morphological characterization of these 3 species using 18 quantitative traits and
27 phenology study: vegetative period, flowering and fruiting time and seed formation for two consecutive
28 years.

29 **Place and Duration of Study:** Experimental plot at the Faculty of Sciences of Tunis, Tunisia- 2009-
30 2010.

31 **Methodology:** Measurements of the 18 morphological characters were performed on 3 samples of
32 *Anacyclus clavatus*, *Chamaemelum fuscatum* and *Leucanthemum parthenium* grown in the Faculty of
33 Sciences of Tunis, for each species, we have studied 10 individuals. Different phenological stages:
34 Vegetative period, Flowering and Fruiting of each species are studied.

35 **Results:** The phenological study show that the 3 species studied have distinct phenologies. The
36 longest phenological cycle is observed for *Leucanthemum parthenium*. Results of morphology study
37 showed significant differences to highly significant for the majority of the traits studied using variance
38 analysis. The comparison of means reveals that *Anacyclus clavatus* and *Chamaemelum fuscatum*
39 form a single group for most of the traits measured, while *Leucanthemum parthenium* is clearly
40 distinct from these two species. In addition, the principal component analysis confirms the results of
41 the variance analysis and the comparison of means.

42 **Conclusion:** The results of the phenological cycle's follow-up show that the 3 species studied have
43 distinct phenologies. The longest phenological cycle is observed for *Leucanthemum parthenium*. The
44 morphological study reveals that *Anacyclus clavatus* and *Chamaemelum fuscatum* form a single
45 group while *Leucanthemum parthenium* is clearly distinct from these two species.

46 **Keywords:** *Anacyclus clavatus*; *Chamaemelum fuscatum*; *Leucanthemum parthenium*;
47 morphological; phenology.

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49 **1. Introduction**

50 Phenological study is important in plant management and combating afforestation, honey analysis,
51 floral biology, estimation of reproductivity and regeneration [1]. It is important also in understanding
52 species interrelations and their interaction with the environment. Variations in phenophases among
53 individuals of different species have been linked to environmental perturbations [2]. A clear
54 understanding of phenological behavior on time of anthesis, time and duration of stigma receptivity,
55 fertilization, mode of pollination, seed development is necessary for breeding programs to obtain
56 better traits [3]. Thus plant phenological study has great significance because it not only provides
57 knowledge about the plant growth pattern but it also provides the idea on the effect of environment
58 and selective pressure on flowering and fruiting behavior [4].

59 Evaluation and characterization through morphological parameters of different crop germplasm is
60 therefore so much important for all plant breeders [5]. Therefore, it is important to make proper
61 strategies for the collection and evaluation of germplasm sources which are locally used in different
62 regions of the world and save them from being vanished [6]. To have a variety of better traits of any
63 crop we need information's about its genetic diversity [7]. Thus, characterization and estimation of
64 genetic diversity is an important step for the competent and successful maintenance and utilization of
65 different crop germplasm [8].

66 Genetic diversity is an inherited variation among and between populations, created, activated and
67 maintained by evolution [9]. Morphological traits provide a simple way of measuring genetic diversity
68 while studying genotype performance under normal growing conditions, but are influenced by
69 environmental factors ([10]; [11]). Plants have the potential to response to the changed environments
70 by changing their morphology and there for, the intra-specific variation in plant characteristics is
71 usually regarded as the adaptive mechanism to different environments [12].

72 The Asteraceae is one of the largest families, comprising 250.000 species. It is known for its wide
73 range of uses not only in medicine but also some plants are grown as ornamental plants such as
74 chamomile (*Leucanthemum parthenium*), others can provide different products: natural rubber,
75 colorants, insecticides and spices [13].

76 *A. clavatus* (*Anacyclus clavatus*), belonging to the Asteraceae family, is an herbaceous, annual and
77 spontaneous plant that is found almost everywhere in the Mediterranean region [14]. It's 20 to 50 cm
78 tall, hairy, green or whitish-green, with an upright or ascending stem, woolly and rowdy whose

79 branches are divorced. Leaves are bipinnate, long to very narrow segments terminated by a small
80 mucron [15]. The convex or somewhat conical receptacle carries triangular bracts, ovals in the shape
81 of sequins. The inflorescences have two types of hermaphrodite flowers: the central flowers are
82 yellow-colored and the peripheral flowers are tongued, long and white. They flourished from March to
83 June [14].

84 The fruits in the form of akene are small, very compressed cuneiform and of grey to beige colour
85 [15]. The number of chromosomes of this species is $2n = 18$ [16]. It's a plant that grows on the edges
86 of fields and roads and in the wastelands of the entire Mediterranean coast [15]. In Tunisia, it's is
87 located in the north (Kroumirie, Oued Medjerda and Cap Bon), and in the center. The use of this
88 species is very limited. The aerial part of *A. clavatus* is used as a powder against stomach pain. It
89 may also be one of the components of tobacco [17].

90 *C. fuscatum* (*Chamaemelum fuscatum*), belonging to the Asteraceae family, anthemidae tribe, and
91 Ormenis sub-section, is an annual, herbaceous, glabrous 30 cm rowing, ascending or upright. The
92 leaves are bipinnate. The heads are heterogeneous with yellow disc and white ligules; their flowering
93 is very early from November to April. The akene is very small, striated, tetragonal and brown to yellow
94 in colour. It's a very widespread plant on the banks of the seguias.

95 In Tunisia, *C. fuscatum* is found in the north (Ain Drahim, Kef), in the center (Sousse, Enfidha) and
96 in the South (Gabes). Internationally, It's located in the western Mediterranean basin of Spain, Greece
97 and North Africa (Tunisia, Morocco and Algeria) [15]. The number of chromosomes of this species is
98 $2n = 18$ [18]. It's known for its anti-malaria property and its protective effect against cell damage [19].

99 *L. parthenium* (*Leucanthemum parthenium*) belongs to the Asteraceae family too, the Anthemidae
100 tribe and the Asteroidea subfamily [20] and the *Leucanthemum* genus. This chamomile is a very
101 fragrant, perennial, rooted plant, with flowering stem erect without hair. The leaves are deeply divided
102 into 4 to 12 toothed segments. The internal tubular flowers are yellow and the ligulate external flowers
103 are white. They flourish from June to August in European conditions [14] and from July to October in
104 Iran [21]. The ripe fruits are brown, glandular and surmounted by a very short membranous crown.

105 *L. parthenium* is a medicinal plant used primarily for the prevention and reduction of migraine attacks
106 frequency, against stomach aches and malaria [22]. It's also known for its properties: antiseptic,

107 stomachic, antihysterical, vermifuge and insecticide. It's found spontaneously on the edges of roads
108 and often in the vicinity of dwellings and it can also be grown in gardens as an ornamental plant.
109 Internationally, *L. parthenium* is found almost all over Europe except the boreal zone and it is also
110 found in South-Western Asia [14].

111 However, there is little information on the morphological diversity and the phenology of *Anacyclus*
112 *clavatus*, *Chamaemelum fuscatum* and *Leucanthemum parthenium* and the potential of these species
113 in breeding programs. The aim of this study is to assess the variations in morphology and phenology
114 of *A. clavatus*, *C. fuscatum* and *L. parthenium*.

115 **2. Materials and methods**

116 **2.1. Plant material and experimental design**

117 Three species of Asteraceae have been studied in this work: *Anacyclus clavatus*, *Chamaemelum*
118 *fuscatum* and *Leucanthemum parthenium*. These species were grown on an experimental plot at the
119 Faculty of Sciences of Tunis, Tunisia under uncontrolled conditions. The seeds used originate from
120 Esbikha for *A. clavatus*, Haouz (Morocco) for *C. fuscatum* whereas the seeds of *L. parthenium* are
121 available in the laboratory of Genetics and Bioresources of the Faculty of Sciences of Tunis.

122 **2.2. Phenological characters**

123 Different phenological stages presented by the individuals of each species are defined:

124 **2.2.1. Vegetative period**

125 This stage spreads from the planting to the beginning of flowering. This is the phase of vegetative
126 growth.

127 **2.2.2. Flowering**

128 This is the period during which the flowers appear. The method of study is based essentially on the
129 visual observation of the appearance of the flowers.

130 **2.2.3. Fruiting**

131 This phase is characterized by the formation of the fruit. It begins with the formation of the first
132 seeds and ends with the general ripening of the seeds.

133 **2.3. Morphological traits**

134 In order to compare the various species studied, we describe the characters of their vegetative
135 part: The type of branching, the stem, the structure of the leaves, the structure of the inflorescences
136 and flowers, the structure of akene and the weight of 100 akenes.

137 Measurements of the morphological characters were performed on three samples of *Anacyclus*
138 *clavatus*, *Chamaemelum fuscatum* and *Leucanthemum parthenium* grown in the Faculty of Sciences
139 of Tunis, for each species, we have studied 10 individuals. The 18 morphological quantitative traits
140 were assessed to characterize and estimate genetic diversity among the 3 species studied, the
141 quantitative traits measured were:

- 142 • Length of main axis in cm: LAP
- 143 • Average length of primary branches in cm: LMRP
- 144 • Average length of branches in cm: LMRS
- 145 • Average length of the tertiary branches in cm: LMRT
- 146 • Length of main root in cm: LRP
- 147 • Number of leaves per plant: NF
- 148 • Average diameter of the receptacle in cm: DMR
- 149 • Average number of leaflets per leaf: NLL
- 150 • Average length of the leaf rachis in cm: LMRF
- 151 • Number of inflorescence per plant: NI
- 152 • Number of primary branches: NRP
- 153 • Number of secondary branches: NRS
- 154 • Number of tertiary branches: NRT
- 155 • Average number of ligules per head: NML
- 156 • Number of ligules of the main axis head: NLCAP
- 157 • Length of the smallest branch in cm: LPR
- 158 • Length of the longest branch in cm: LLR

- 159 • Weight of 100 akenes : $P_{100} A$

160 2.4. Data analysis

161 The evaluation of a collection of genetic resources is commonly based on the simultaneous
162 examination of many populations for various morphological characters. In this context, data on the
163 different morphological traits measured were:

- 164 • An analysis of variance with one classification criterion followed by a comparison of means.
- 165 • An estimate of the degrees of association between the different traits studied by the Pearson
166 correlation coefficient [23].
- 167 • A principal component analysis (PCA) based on the derivation of orthogonal variables [24].

168 In order to evaluate morphological diversity and to establish relationships among studied species,
169 several statistical procedures were conducted. Quantitative data were computed using the software
170 XLSTAT version 2011 to perform analysis of variance, comparison of mean using the Duncan test
171 and to calculate the Pearson correlation coefficient. Principal component analysis (PCA) was also
172 done using the software XLSTAT.

173 3. Results and discussion

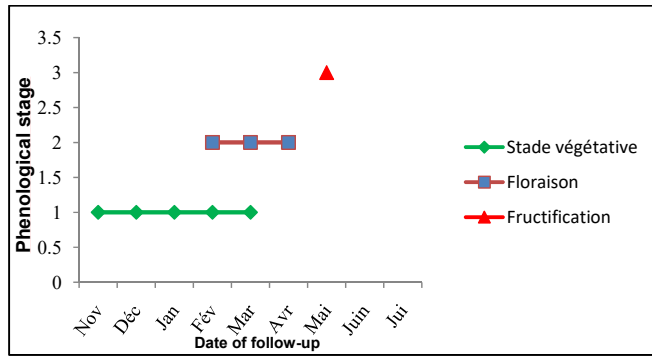
174 3.1. Phenology study

175 3.1.1. Vegetative period

176 The vegetative period is characterized by a strictly herbaceous development and extends from
177 seedling to full bloom. We divided this phase into 2 stages:

178 **Stage of germination:** it is characterized by the appearance of the primordial leaves. In all three
179 species, the germination begins after 10 days.

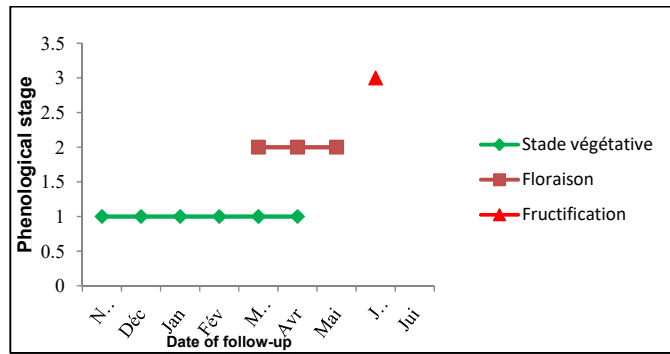
180 **Stage of foliage:** Observation of the phenological spectrum reveals that this stage is the longest of
181 the phenological cycle. This stage, which is characterized by the growth of the stems in length and by
182 the formation of the leaves, lasts 6 months for *Chamaemelum fuscatum* (Figure 1) and 7 months for
183 *Anacyclus clavatus* (Figure 2). *Leucanthemum parthenium* is a perennial herb plant (Figure 3).



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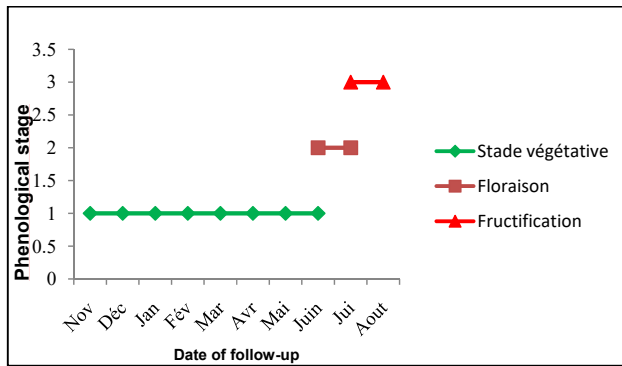
Fig.1. Phenological cycle of *Chamaemelum fuscatum*



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Fig.2. Phenological cycle of *Anacyclus clavatus*



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Fig.3. Phenological cycle of *Leucanthemum parthenium*

190 **3.1.2. Flowering**

191 Flowering is considered from the formation of the first flower until most flowers have evolved this
192 period differs from one species to another: For *Chamaemelum fuscatum*, the flowering period ranges
193 from mid-February to the end of April (Figure 1). For *Anacyclus clavatus*, this period extends from the
194 end of March to mid-May (Figure 2). For *Leucanthemum parthenium*, the first flower blooms in early
195 June and full bloom is observed around mid-July (Figure 3).

196 Flowering appears to be highly favoured during the rainy season for *Anacyclus clavatus* and
197 *Chamaemelum fuscatum*, only *Leucanthemum parthenium* flowers during the dry season. We find
198 that the species *Chamaemelum fuscatum* characterized by a very early flowering date has a spread
199 flowering period. In addition, the species *Leucanthemum parthenium* characterized by a late flowering
200 date has a relatively short flowering stage and this to escape the water stress.

201 **3.1.3. Fruiting**

202 It is the formation of fruit in the form of akene. We have noticed that the appearance of the first
203 akene coincides with the peak of flowering, while the full fructification characterized for the 3 species
204 by the change of color flowers in tubes from yellow to light grey and the fall of the white ligules is
205 generally obtained after two weeks of the appearance of the first fruit (Figure 1, 2 and 3).

206 In fact, the study of [25] reveals that **akenes** of *A. clavatus* that germinated earlier produced plants
207 with higher biomass and higher reproductive effort. In addition, this work show that the phenology of
208 *Anacyclus clavatus* **akene** germination was the main factor affecting post dispersal life-history traits
209 related to competitive ability and reproductive success.

210 In addition, the study of [26] showed a high phenological diversity for the four phenological patterns
211 (buds, flowers, fruits and seeds) among fifteen leguminous plant species growing in Amritsar.

212 **3.2. Morphology study**

213 **3.2.1. Study of vegetative part**

214 A comparative morphological characteristics of the 3 species studied is shown in Table 1.

215 **Table 1:** Main distinctive characteristics of 3 species studied.

| Species | NR | Leafs | Flowers | Akenes | P ₁₀₀ A in mg | DR (cm) |
|--------------------------------|-----|--|----------------------------|--------------------|-----------------------------|-------------|
| <i>Anacyclus clavatus</i> | T+5 | Dark green bipinnate | White ligulated flowers | Beige | 45.23 | 1.56 ± 0.01 |
| <i>Chamaemelum fuscatum</i> | T+5 | Green bipinnate | Flowers in yellow tubes | Brown to yellow | 26.63 | 0.67 ± 0.05 |
| <i>Leucanthemum parthenium</i> | T+3 | Greenish- yellowish divided into wide | White ligulated flowers | Brown | 9.96 | 0.65 ± 0.02 |

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217 **NR:** number of ramifications, **P₁₀₀ A:** weight of 100 akenes, **T:** number of branches, **DR:** diameter of
218 the receptacle.

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220 The inflorescences and the flowers

221 The inflorescence of *Anacyclus clavatus*, *Chamaemelum fuscatum* and *Leucanthemum parthenium*
222 is a flower head containing two types of flowers: yellow flowers tubulated in the center and white
223 flowers ligated at the periphery. The flowers of the 3 species have the same floral biology, but show a
224 difference in floral structure. Indeed, the liguled flowers of *Chamaemelum fuscatum* are long and
225 beaked at the tip, while those of two other species are similar; they are short and more or less
226 rounded.

227 The diameter of the receptacle varies from one species to another. It is 0.65 ± 0.02 cm in
228 *Leucanthemum parthenium*, 0.67 ± 0.05 cm in *Chamaemelum fuscatum* and 1.56 ± 0.01 cm in
229 *Anacyclus clavatus*.

230 Fruit

231 The fruits differ between the 3 species studied. The fruit of *Anacyclus clavatus* (Figure 4) is an
232 indelible akene, beige at maturity, of rectilinear shape to flattened cone. This akene is surrounded by
233 two membranous wings, clear, very thin, parchment and truncated at the apex. In the case of an
234 akene without these wings, the fruit appears mottled and has four longitudinal ribs.

235

236 The fruit of *Chamaemelum fuscatum* (Figure 5) is an indehiscent akene, very small, not marginated,
237 flattened ovoid, raised by 3 ribs weak and finely striated. Their color is brown to yellow at maturity.

238 The fruit of *Leucanthemum parthenium* (Figure 6) is an indehiscent akene, very small, brown at
239 maturity, glandular and surmounted by a very short membranous crown and crenate.

240

241 **Weight of 100 akenes**

242 The mean weight of 100 akenes of *A. clavatus* is 45.23 mg. For *C. fuscatum*, it is 26.63 mg. An
243 average weight of 9.96 mg was calculated in *L. parthenium* (Table 1).

244 **3.2.2. Analysis of morphological variability**

245 **3.2.2.1. Analysis of variance**

246 The analysis of variance with one classification criterion (species effect) showed highly significant
247 differences between the three species studied (Table 2) for the majority of the quantitative traits

248 measured such as: Length of the longest branch (LLR), Length of the smallest branch (LPR), number
 249 of secondary branches (NRS), number of primary branches (NRP), mean leaf spine length (LMRF),
 250 average number of leaflets (NLL), mean diameter of the receptacle (DMR), length of the main root
 251 (LRP), mean length of the tertiary branch (LMRT), average length of secondary branch (LMRS),
 252 average length of primary branch (LMRP) and length of the main axis (LAP). The difference between
 253 the three species is not significant for: The number of the principal axis head ligules (NLCAP), the
 254 average number of ligules per capitule (NML) and the number of tertiary branches (NRT). This result
 255 reflects a phenotypic heterogeneity between the 3 species studied, taking into account the measured
 256 parameters.

257 **Table 2:** Results of the variance analysis of the 17 morphological traits measured.

| Characters | df | Average square | F _{obs} | Pr > F |
|------------|----|----------------|------------------|--------------------|
| LAP | 2 | 3730,630 | 68,058 | < 0,0001 HS |
| LMRP | 2 | 982,641 | 26,382 | < 0,0001 HS |
| LMRS | 2 | 862,412 | 52,589 | < 0,0001 HS |
| LMRT | 2 | 360,894 | 26,359 | < 0,0001 HS |
| LRP | 2 | 40,961 | 11,73 | 0,000 HS |
| NF | 2 | 338256,13 | 5,355 | 0,011 S |
| DMR | 2 | 2,701 | 108,846 | < 0,0001 HS |
| NLL | 2 | 150,633 | 75,039 | < 0,0001 HS |
| LMRF | 2 | 11,796 | 36,769 | < 0,0001 HS |
| NI | 2 | 30601,433 | 2,983 | 0,068 NS |
| NRP | 2 | 185,633 | 14,312 | < 0,0001 HS |
| NRS | 2 | 14770 | 15,244 | < 0,0001 HS |
| NRT | 2 | 4548,433 | 0,867 | 0,432 NS |
| NML | 2 | 226,9 | 1,258 | 0,3 NS |
| NLCAP | 2 | 0,7 | 1,086 | 0,352 NS |
| LPR | 2 | 15,74 | 22,619 | < 0,0001 HS |
| LLR | 2 | 935,217 | 8,415 | 0,001 HS |

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 259 **df** : degree of freedom; **F_{obs}** : F observed ; **HS**: highly significant; **S**: significant ($P < 0.05$) ; **NS**: no
 260 significant ($P \geq 0.05$).
 261

262 **3.2.2.2. Comparison of means**

263 According to the Duncan test, we distinguish 5 types of groups (Table 3). Comparison of means
 264 shows that *A. clavatus* and *C. fuscatum* are distinctly different from *L. parthenium* for: the length of the
 265 main axis (LAP), the mean length of the secondary branch (LMRP), the average length of the tertiary
 266 branch (LMRT), Root length (LR), number of leaves (NF), number of primary branches (NRP) and
 267 number of secondary branches (NRP).

268 *A. clavatus* is distinguished from *L. parthenium* and *C. fuscatum* for the mean diameter of the
 269 receptacle (DMR), the length of the smallest branch (LPR) and the length of the longest branch (LLR).
 270 In fact, the three species did not differ significantly in the mean diameter of the receptacle (DMR), the
 271 length of the smallest branch (LPR) and the length of the longest branch (LLR).

272 The parameters discriminating the three species are: the average length of the primary branch
 273 (LMRP), the mean number of leaflets per leaf (NMf) and the average length of the spine (LMRF). For
 274 the number of inflorescence per plant (NI), *Anacyclus clavatus* is not significantly different from
 275 *Chamaemelum fuscatum* or *Leucanthemum parthenium*. Therefore, *Anacyclus clavatus* and
 276 *Chamaemelum fuscatum* are much alike for more than half the morphological characters studied.
 277 Most of the highest averages of the morphological traits are observed in *Anacyclus clavatus*, while the
 278 majority of the lowest averages are observed in *Leucanthemum parthenium* (Table 3).

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284 **Table 3:** Comparison of means of the 3 species studied using the Duncan test.

| Traits | <i>Anacyclus clavatus</i> | <i>Chamaemelum fuscatum</i> | <i>Leucanthemum parthenium</i> |
|--------|---------------------------|-----------------------------|--------------------------------|
| LAP | 19,8 B | 20,71 B | 53,7 A |
| LMRS | 20,6 A | 17,91 A | 3,39 B |
| LMRT | 12,12 A | 12,5 A | 1,91 B |
| LR | 8,1 B | 7,72 B | 11,4 A |
| NF | 629,5 A | 524,5 A | 271,7 B |
| NRP | 11,4 B | 11,9 B | 19,1 A |

| | | | | |
|--------------|---------------|---------|---------|-----|
| NRS | 39,6 A | 29,6 B | 100,6 A | 285 |
| DMR | 1,56 A | 0,67 B | 0,65 B | |
| LPR | 3,21 A | 1,4 B | 0,8 B | 286 |
| LLR | 46,69 A | 29,97 B | 29,91 B | |
| NRT | 53,7 A | 37,3 A | 79,6 A | 287 |
| NML | 11,7 A | 19,9 A | 11,6 A | |
| NLACP | 13,3 A | 13,4 A | 12,9A | 288 |
| LMRP | 36,12 A | 24,34 B | 16,42 C | |
| NMf | 15,6 A | 10,9 B | 7,9 C | 289 |
| LMRF | 4,36 A | 3,19 B | 2,19 C | |
| NI | 116,5 A and B | 82,4 B | 190,6 A | 290 |

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296 **3.2.2.3. The Matrix of correlation coefficients**

297 The matrix of correlation coefficients between the characters studied (Table 4) shows: A positive
 298 correlation of the following traits: LMRP and LMRS correlate positively with each other and with all the
 299 parameters of LMRT, NF, DMR, NLL, LPR and LLR ; The character LAP is strongly correlated
 300 positively with the parameters LR, LMRF, NI, NRP and NRT ; A highly significant positive correlation
 301 between LMRF with NI, NRP and NRS ; NI correlates strongly with the parameters : NRP, NRS and
 302 NRT and weakly with LLR ; NRP is strongly correlated with NRS and weakly correlated with the
 303 characters NRT and LPR. The LAP has a highly significant negative correlation with the parameters
 304 (LMRS, LMRT, NLL) and significant with the characters (LMRP, NF, DMR, LPR); LMRS. It is
 305 important to note that NLCAP and NML are not correlated with any of the other characters and that
 306 LMRP is the most positively correlated with the other traits (Table 4).

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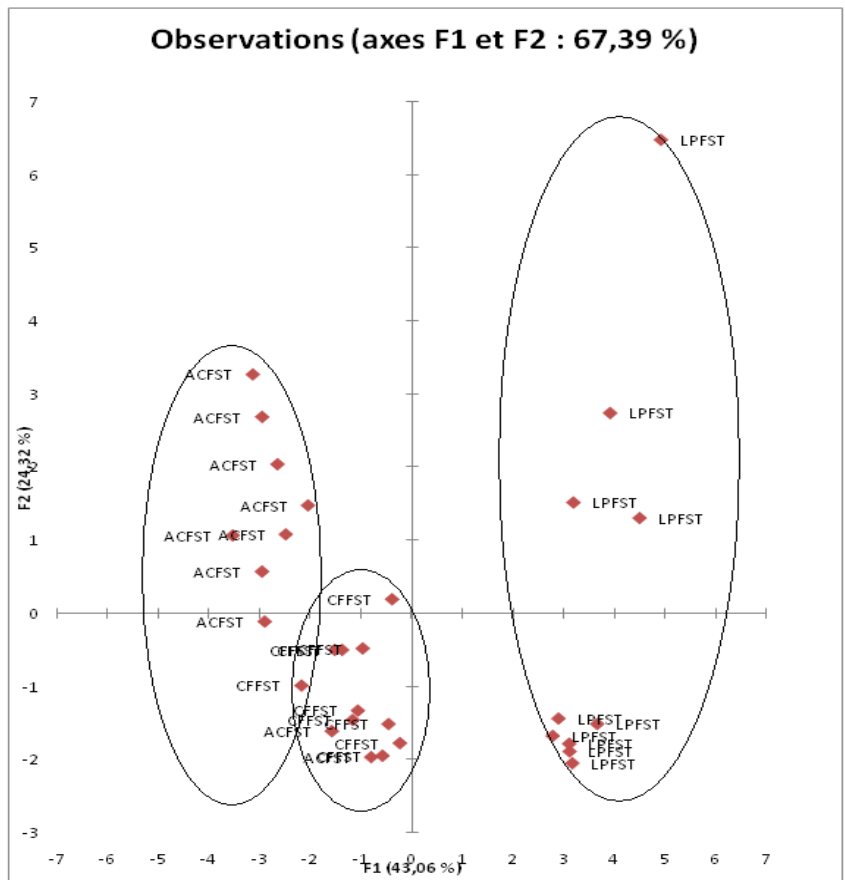
308 **3.2.2.4. Principal component analysis**

309 The graphical representation of the individuals dispersion of the 3 species studied reveals a
310 homogeneous grouping of the species studied forming 3 clear groups (Figure 7).

311 Indeed, there is a slight overlap between the two groups: *Anacyclus clavatus* and *Chamaemelum*
312 *fuscatum*, whereas, *Leucanthemum parthenium* group seems very distinct from the two others
313 species. These results confirm those of the variance analysis which showed a strong resemblance
314 between *Anacyclus clavatus* and *Chamaemelum fuscatum*.

315 It is also observed that the individuals of the species *Chamaemelum fuscatum* occupy a rather
316 restricted part of the plane and are located entirely in the negative part of the two axes F1 and F2.
317 While, the individuals belonging to *Anacyclus clavatus* are scattered on the two axes (F1 and F2) with
318 a trend towards the positive values of the F1 axis (Figure 7).

319 Furthermore, individuals of *Leucanthemum parthenium* are the best dispersed on the 2 axes (F1 and
320 F2) with a tendency towards the negative values of F1 axis (Figure 7).



321

322 **Figure 7:** Principal components analysis of the 3 species studied using XLSTAT software

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Table 4: Matrix of correlation coefficients of the different morphological parameters.

| Traits | LAP | LMRP | LMRS | LMRT | LR | NF | DMR | NMF | LMRF | NI | NRP | NRS | NRT | NML | NLCAP | LPR | LLR |
|--------|--------|--------|--------|--------|--------|--------|--------|--------|--------|--------|--------|--------|--------|--------|-------|-------|-----|
| LAP | 1 | | | | | | | | | | | | | | | | |
| LMRP | -0.536 | 1 | | | | | | | | | | | | | | | |
| LMRS | -0.810 | 0.842 | 1 | | | | | | | | | | | | | | |
| LMRT | -0.766 | 0.707 | 0.918 | 1 | | | | | | | | | | | | | |
| LR | 0.607 | -0.270 | -0.572 | -0.544 | 1 | | | | | | | | | | | | |
| NF | -0.388 | 0.797 | 0.763 | 0.764 | -0.281 | 1 | | | | | | | | | | | |
| DMR | -0.451 | 0.679 | 0.496 | 0.315 | -0.290 | 0.271 | 1 | | | | | | | | | | |
| NMF | -0.670 | 0.691 | 0.662 | 0.522 | -0.511 | 0.423 | 0.798 | 1 | | | | | | | | | |
| LMRF | 0.790 | -0.266 | -0.629 | -0.677 | 0.451 | -0.269 | -0.048 | -0.283 | 1 | | | | | | | | |
| NI | 0.532 | 0.220 | -0.123 | -0.143 | 0.417 | 0.377 | -0.176 | -0.195 | 0.523 | 1 | | | | | | | |
| NRP | 0.803 | -0.291 | -0.579 | -0.594 | 0.575 | -0.135 | -0.410 | -0.572 | 0.673 | 0.528 | 1 | | | | | | |
| NRS | 0.826 | -0.119 | -0.494 | -0.461 | 0.603 | 0.014 | -0.314 | -0.455 | 0.701 | 0.872 | 0.774 | 1 | | | | | |
| NRT | 0.410 | 0.329 | -0.007 | -0.004 | 0.303 | 0.462 | -0.104 | -0.080 | 0.373 | 0.946 | 0.473 | 0.798 | 1 | | | | |
| NML | -0.130 | 0.052 | 0.090 | 0.095 | 0.014 | 0.269 | -0.160 | -0.031 | -0.136 | 0.025 | 0.171 | -0.172 | 0.048 | 1 | | | |
| NLCAP | -0.179 | 0.282 | 0.282 | 0.325 | -0.357 | 0.254 | 0.153 | 0.161 | -0.267 | 0.006 | -0.058 | -0.016 | 0.075 | -0.020 | 1 | | |
| LPR | -0.529 | 0.576 | 0.492 | 0.378 | -0.385 | 0.289 | 0.762 | 0.787 | -0.142 | -0.224 | -0.478 | -0.387 | -0.153 | -0.114 | 0.247 | 1 | |
| LLR | -0.184 | 0.868 | 0.597 | 0.465 | -0.051 | 0.722 | 0.541 | 0.485 | 0.088 | 0.495 | 0.058 | 0.248 | 0.526 | -0.094 | 0.194 | 0.396 | 1 |

332 In fact, the morphological study of [27] showed variations among the 33 accessions of *Ricinus*
333 *communis* L. from Andaman and Nicobar Islands for all the 18 traits studied. This work reveals also
334 that plant height exhibited high significant positive correlations with the number of nodes on the main
335 stem. In addition, the cluster analysis based on morphological traits grouped the 33 accessions of
336 *Ricinus communis* L. into two major clusters [27].

337 Furthermore, the study of [28] was found a significant amount of genetic variability for all the twenty
338 morphological parameters studied among safflower germplasm. In addition, this work reveals that
339 seed yield plant had high significant and positive correlation with branches plant, capitulum plant,
340 seeds capitulum and 100 seed weight. Furthermore, the hierarchical cluster analysis based on agro-
341 morphological parameters divided the 121 accessions of safflower into 5 main clusters [28].

342 The morphological study of [29] in rice varieties showed high phenotypic variability ($P < 0.0001$) for
343 the characters: leaf length and leaf width, primary branching, maturity and grain thickness. In addition,
344 this work revealed a positive and strong correlation (0.77) between the height at maturity and leaf
345 length. The cluster analysis of this morphological study based on Euclidian distances between the 98
346 genotypes of Rice has allowed identifying three major clusters.

347 **4. Conclusion**

348 The phenological study shows that the 3 species studied have distinct phenologies. The longest
349 phenological cycle is observed for *Leucanthemum parthenium*. The variance analysis showed
350 significant differences to highly significant for the majority of the traits studied. Furthermore, this study
351 allowed us to validate the morphological and phenological approach as tools for selection of suitable
352 genotypes. This genetic diversity will be more evidenced using molecular markers. Although, the
353 morphological descriptors of *Anacyclus clavatus*, *Chamaemelum fuscatum* and *Leucanthemum*
354 *parthenium* must be completed by a molecular analysis using RAPD, SSR or AFLP to understand the
355 genetic organization of these species in Tunisia.

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