

1 Screening Groundnut (*Arachis hypogaea*) Genotypes for Resistance to Early and 2 Late Leaf Spot Diseases

3

4 ABSTRACT

5 **Background for the study:** Groundnut (*Arachis hypogaea* L.) is an important crop both in subsistence
6 and commercial agriculture in Ghana. Early leaf spot (*Cercospora arachidicola*) and late leaf spot
7 (*Cercosporidium personata*) are major limiting factors to groundnut productivity in Ghana.

8 **Aim:** The objective of the study was to screen groundnut genotypes for resistance to Early and Late leaf
9 spot diseases.

10 **Study Design:** The treatments were arranged in a randomized complete block design and replicated
11 three (3) times.

12 **Place and Duration of Study:** The research was conducted from May to December 2013 at the Savanna
13 Agricultural Research Institute experimental site at Nyankpala in the Northern Region. The site lies
14 between latitude 9° 25' 141 North and longitude 0° 58' 142 West and an altitude of 183m.

15 **Methodology:** The land was done using a tractor and field divided into plots of 2m x 5m with 1m interval
16 between plots. Sowing was done on 3rd June 2013. One seed was planted per hole at a depth of
17 approximately 4 cm. Early and late leaf spot ratings were recorded at 30, 60 and 90 days after planting,
18 using a ten-point scale. At pod maturity, plants from the middle two rows of each plot were hand-
19 harvested and weights of the above ground foliage and underground pods were oven-dried to obtain dry
20 haulm weight and pod yield respectively. Data collected were subjected to analysis of variance using
21 Genstat statistical package (12th edition). Means were separated using the least significant difference at
22 5%.

23 **Results:** There were significant differences ($P < .001$) among the groundnut genotypes in terms of severity
24 for both early and late leaf spot diseases. F-Mix, NC 7, PC 79-79, F-Mix x SINK 24 and NKATIE-SARI
25 had lowest score for both early and late leaf spot diseases. Among the 21 groundnut genotypes, F-Mix
26 recorded the highest pods yield of 1100kg/ha and haulm weight of 5867kg/ha followed by NC 7 with total
27 pods yield of 900kg/ha and haulm weight of 5373kg/ha. PC 79-79 had a total pods yield of 666.7kg/ha
28 and haulm weight of 4867kg/ha. The pods yield of F-Mix x SINK 24 was 533.3kg/ha and haulm weight of
29 4600kg/ha. NKATIE-SARI recorded pods yield of 500kg/ha and haulm weight of 4633kg/ha.

30 **Conclusions:** From the study, the genotypes F-Mix, NC 7, PC 79-79, F-Mix x SINK 24 and NKATIE-
31 SARI were found to be resistance to both early and late leaf spot disease whereas Chinese, Doumbala,
32 GM 120, GM 324 and ICGV 86015 were susceptible to both diseases.

33 **Key words:** Groundnut, leaf spot, genotypes, disease severity.

34 1.0 INTRODUCTION

35 Groundnut (*Arachis hypogaea* L.) is an important economic food and cash crop grown globally. The crop
36 contributes to the world diet and is used for food for people from both developed and developing
37 countries.

38 Groundnut also has other uses such as; source of cooking oil, solvents and medicine. It can also be
39 eaten raw, slightly cooked or when it is still fresh.

40 Groundnut vines (hay) can also serve as fodder for livestock especially during the dry season when there
41 is scarcity of green forage for livestock (Tsigbey *et al.*, 2004; Naab *et al.*, 2005).

42 According to Asiedu (1989), groundnut is a herbaceous plant of which there are two types, bunch and
43 runner types. Apart from the runner and bunch types, many intermediate forms or hybrids exist (Irvine,
44 1974).

45 According to FAO estimate, the average world production of groundnut pods in 1990 – 2003 was between
46 34.4 million tons/year from 24.4 million hectares of land (Ntare, 2007). The largest producers of
47 groundnuts are China and India followed by Sub-Saharan African countries, central and South America
48 (Johnson and Ives, 2001). Approximately 85% of the land area under groundnut production is in the
49 Sudan and Guinea Savannah zones (Amahene – Amankwa *et al.*, 1998). The total production in Sub-
50 Saharan Africa was 8.2 million tons/year from 9.5 million hectares of land (Ntare 2007). Groundnut is
51 cultivated in all agro ecological zones of Ghana. However, a large proportion (92%) of the production is in
52 Northern-Ghana (SRID, 2004; Twemboah, 2000).

53 It is produced both as a commercial and a subsistence crop (Tsigbey *et al.*, 2004). Groundnut yields are
54 however very low in Ghana averaging less than 1000kg/hectare in comparison with an average of
55 2500kg/hectare obtained in developed countries (Shokes and Culbreath, 1997). In Ghana, the common
56 limiting factor to groundnut production is the prevalence of diseases, most importantly early leaf spot
57 caused by *Cercospora arachidicola* and late leaf spot caused by *Cercosporidium personata* (Frimpong *et al.*,
58 2006a). Both diseases are distributed widely and occur in epidemic proportion in northern Ghana
59 (Nutsugah *et al.*, 2007). The incidence of early and late leaf spot on susceptible groundnut genotypes can
60 lead to total defoliation which can drastically reduce yield (Shew *et al.*, 1995). High defoliation can also
61 affect hay quality of vines fed to livestock (Tsigbey *et al.*, 2004).

62 Leaf spot disease appear as reddish brown to black necrotic spot on leaves. The necrotic spot in early
63 leaf spot are bounded by a yellow halo which separates it from late leaf spots which have no halo.
64 Both diseases reduce the leave area available for photosynthesis and cause premature defoliation
65 and yield loss as much as 70% in West Africa (Shokes and Culbreath, 1997; Waliyar *et al.*, 2000).
66 Annual yield losses of up to 50% attributable to the disease can occur in northern Ghana (Nutsugah *et al.*
67 *et al.*, 2006; Tsigbey *et al.*, 2001). The most common method of control is by the use of fungicides.
68 However, in Ghana, a large number of farmers do not practice any control for these diseases in their
69 groundnut farms largely due to inadequate resources to use the appropriate chemical control and/or
70 difficulty in obtaining fungicides (Allen 1983; Nutsugah *et al.*, 2007). Moreover, these chemicals leave
71 toxic residues in the environment (Maloy 1993; Ihejirika *et al.*, 2006a).

72 Gibbons (2002), discovered that application of chemicals against leaf spot as well as against other
73 diseases should be reduced to a minimum on health and environmental grounds. Tuormaa (2006)
74 reported that, world health organization (WHO) report estimated that there were between 800,000 and
75 1,500,000 cases of unintentional pesticides poisoning in the world, resulting to about 3,000 and
76 28,000 deaths.

77 A case study finding showed highly disturbing levels of pesticides misuse and abuse leading to
78 poisoning of families and livestock (Adolpus, 2007).

79 Due to the above mentioned reasons, the use of disease – resistant cultivars is the best and
80 economically way to control diseases of food crops (Mallikarjuna *et al.*, 2004; Jyosthna *et al.*, 2004). It
81 is the most cost efficient of all the control measures (Driscoll, 1990).

82 Resistant cultivars save time, effort and money which would have been spent in controlling plant
83 diseases. The environment also gains because there will be no application of pesticides (Maloy,
84 1993). Subrahmanyam *et al.*, (1982) reported that some genotypes of groundnut are resistant to
85 cercospora leaf spot (early and late leaf spot).

86 The objective of this study was to screen groundnut genotypes for resistance to early and late leaf spot
87 diseases.

88

89 **2.0 Materials and Methods**

90 **2.1 Experimental Site.**

91 The research was conducted from May to December 2013 at the Savanna Agricultural Research Institute
92 experimental site at Nyankpala in the Northern Region. The site lies between latitude 9° 25' 141 North
93 and longitude 0° 58' 142 West and an altitude of 183m.

94 **2.2. Field Experiment and Design.**

95 The land was ploughed and harrowed using a tractor and divided into plots of 2m x 5m with 1m interval
96 between plots. The treatment were arranged in a randomized complete block design and replicated three
97 (3) times. Sowing was done manually on 3rd June 2013. One seed was planted per hole at a depth of
98 approximately 4 cm. A plot consisted of six rows with spacing of 0.5 m between rows and 0.2 m between
99 plants in a row. Weed control was done using hoe to remove weeds and hand pulling of weeds where
100 necessary.

101 **2.2.1. Source of Groundnut genotypes.**

102 Twenty-one (21) groundnut genotypes obtained from CSIR-Savanna Agricultural Research
103 Institute and its collaborators from Burkina Faso were used for the study. The groundnut
104 genotypes used and their source are listed in table 1 below.

105

106 **Table 1.0 groundnut genotypes and their sources.**

Number	Genotype	Sources
1	NC 7	Tamale-Ghana
2	NKATIE-SARI	Tamale-Ghana
3	ICGV 86015	Tamale-Ghana
4	FDRS × F-MIX	Tamale-Ghana
5	ICGV 92096	Tamale-Ghana
6	F-MIX × SINK 24	Tamale-Ghana
7	CHINESE	Tamale-Ghana

8	F-MIX	Tamale-Ghana
9	DOUMBALA	Burkina Faso
10	TS-32-1	Burkina Faso
11	PC 79-79	Burkina Faso
12	GM 204 (123)	Burkina Faso
13	ICGV 86124	Tamale-Ghana
14	ICGV 97188	Tamale-Ghana
15	ICGV 96814	Tamale-Ghana
16	ICGV 86015	Tamale-Ghana
17	GM 656	Burkina Faso
18	GM 155	Burkina Faso
19	GM120	Burkina Faso
20	GM 663	Burkina Faso
21	GM 324	Burkina Faso

107 2.3. Data Collection and Analysis

108 Data was collected on leaf spot disease severity, number of pods per 5 plants, haulm weight, pod yield
109 and maturity rating. Data on leaf spot severity were taken at 30, 60 and 90 days after planting using the
110 Florida scale of (1-10) based on visual observations where 1 = no disease, 2 = very few lesions on leaves
111 in lower canopy, 3 = few lesions on leaves in lower and upper canopy, 4 = some lesions on leaves in
112 lower and upper canopy with $\leq 10\%$ defoliation, 5 = lesions noticeable in upper canopy and $\leq 25\%$
113 defoliation, 6 = lesion numerous with $\leq 50\%$ defoliation, 7 = lesions very numerous with $\leq 75\%$ defoliation,
114 8 = numerous lesions on few remaining leaves with $\leq 90\%$ defoliation, 9 = remaining leaves covered with
115 lesions with $\leq 95\%$ defoliation, and 10 = plants defoliated or dead (Chitekaet *al.*, 1997). Ten plants were
116 sampled per plot. Data also was taken on incidence of rossete as additional data.

117 At pod maturity, plants from the middle two rows of each plot were hand-harvested and weights of the
118 above ground foliage and underground pods were oven-dried to obtain dry haulm weight and pod yield
119 respectively. Data collected were subjected to analysis of variance and the mean difference separated
120 using the least significant difference test at LSD of $p \leq 0.05$ using Genstat statistical package (4th edition).

121 3.0 RESULTS

122 3.1 Disease score of early and late leaf Spot

123 Table 2 Effect of groundnut genotype on severity score for early and late leaf spot diseases.

TREATMENTS	DISEASE SCORE
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	EARLY LEAF SPOT	LATE LEAF SPOT
CHINESE	5.133 a-d	6.933 a
DOUMBALA	4.533 d-f	6.200 bc
F-MIX	4.200 f	5.200 de
F-MIX × SINK 24	4.600 c-f	5.333 de
FDRS × F-MIX-39	4.867 a-e	5.933 bc
GM 120	5.267 ab	6.467 ab
GM 155	5.200 abc	6.200 bc
GM 204 (123)	5.267 ab	6.267 bc
GM324	5.333 a	6.333 b
ICGV 86015	5.000 a-e	6.267 bc
ICGV 86024	5.267 ab	6.267 bc
ICGV 92096	4.867 a-e	6.133 bc
ICGV 96814	4.400 ef	6.067 bc
ICGV 97188	5.133 a-d	6.133 bc
ICGV 86124	5.133 a-d	6.133 bc
TS-32-1	5.133 a-d	6.133 bc
NC 7	4.133 f	5.333 de
NKATIE-SARI	4.667 b-f	5.067 e
PC 79-79	4.600 c-f	5.733 cd
GM 656	5.067 a-d	6.067 bc
GM 663	5.000 a-e	6.000 bc
P value	<.001	<.001
CV%	2.6	0.5
SED	0.3039	0.2786

124 Genotypes with different letters in a column are significantly different (protected LSD test, P<0.05)

125 The results indicated that there was significant difference ($P < .001$) among the groundnut genotypes in
 126 the severity of both early and late leaf spot diseases (Table 1). The genotypes NC7, F-MIX, ICGV
 127 96814, PC 79-79, NKATIE-SARI, F-MIX \times SINK-24 and FDRS-F-MIX-39, had lower disease severity
 128 scores for early leaf spot and so have some level of resistance to early leaf spot while the genotypes
 129 CHINESE, ICGV 86015, ICGV 86124, ICGV 97188, TS-32-1, GM 656, GM 155, GM 120, GM 204 (123)
 130 and GM 324 had higher disease severity score for early leaf spot and so were susceptible to early leaf
 131 spot disease. Chinese and Doumbala were used as susceptible checks while Nkatie-SARI was the
 132 resistant check.

133 For late leaf spot, the genotypes NKATIE-SARI, F-MIX, F-MIX \times SINK 24, NC7 and PC 79-79 recorded
 134 the lowest disease severity scores while the genotypes GM 656, ICGV 96814, TS-32-1, ICGV 92096,
 135 ICGV 97188, ICGV 86124, ICGV 86015, GM 663, GM 155, ICGV 86024, GM 204 (123), GM 120, GM
 136 324, DOUMBALA and CHINESE had the highest disease severity score and so exhibit some level of
 137 susceptibility to late leaf spots.

138 **3.2 Total number of pod/plant, number of matured pod/plant and dry pod weight.**

139 Table 3 shows the effect of groundnut genotypes on total number of pods/plant; number of matured
 140 pods/plant and dry pod weight

TREATMENTS	Total number of pods/plant	Number of matured pods/plant	Dry pod weight (kg/Ha)
CHINESE	5.67 j	3.67 i	500.0 b-f
DOUMBALA	10.67 gh	7.667 fgh	866.7 a-d
F-MIX	19.67 a	17.00 a	1100.0 a
F-MIX \times SINK 24	17.00 ab	14.33 abc	533.3 b-f
FDRS-F-MIX-39	15.33 bcd	12.67 bcd	566.7 b-f
GM 120	7.67 ij	4.00 i	300.0 efg
GM 155	14.33 b-e	10.33 def	533.3 b-f
GM 204 (123)	11.67 e-h	9.00 efg	400.00 efg
ICGV 86024	11.67 e-h	7.67 fgh	600.0 b-f
GM 656	11.33 fgh	8.00 efg	233.3 fg
GM 663	10.33 hi	7.67 fgh	366.7 efg
GM 324	7.6 ij	4.33 hi	366.7 efg
ICGV 86015	13.67 c-f	10.33 def	700.0 a-e
ICGV 86124	13.33 d-g	11.33 bcd	933.3 ab

ICGV 97188	13.33 d-g	10.00 d-g	500.0 b-f
ICGV IS 96814	10.33 hi	6.67 ghi	566.7 b-f
ICGV IS 92096	12.00 e-h	9.00 efg	466.7 c-f
TS-32-1	16.33 bc	10.67 def	433.3 d-g
NC 7	17.00 ab	14.67 abc	900.0 abc
NKATIE-SARI	17.00 ab	15.00 ab	500.0 b-f
PC 79-79	16.33 bc	13.33 a-d	666.7 a-f
P value	<.001	<.001	0.004
CV%	5	4.4	12.6
SED	1.462	1.745	217.9

141 Genotypes with different letters in a row are significantly different (Protected LSD test, $P < 0.005$).

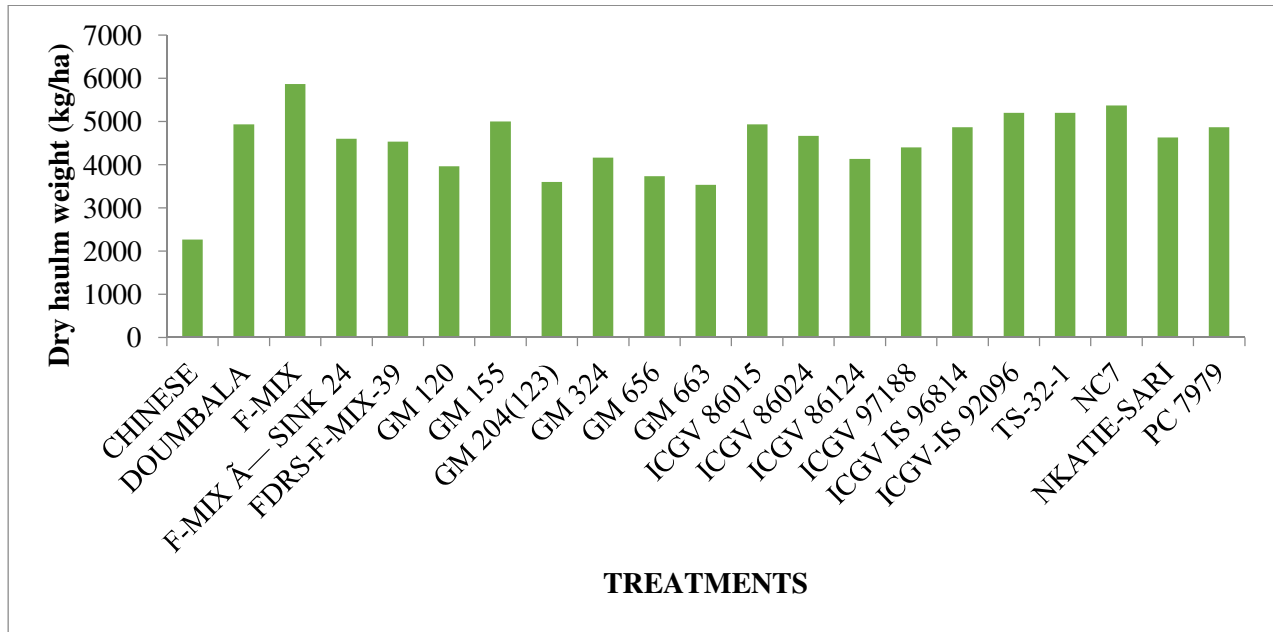
142 The genotypes F-MIX, NC 7, NKATIE-SARI, F-MIX × SINK 24, PC 79-79, TS-32-1, FDRS-F-MIX-39 and
143 GM 155 recorded the highest total number of pods/ plant whereas CHINESE, GM 120, GM 324, GM 663,
144 DOUMBALA, GM 656 and GM 204 (123) had the lowest total number of pods/plant.

145 Also, F- MIX, NKATIE-SARI, NC 7, F-MIX × SINK 24, PC 79-79 and FDRS-F-MIX-39 had the highest
146 number of matured pods while Chinese, GM 120, GM 324, ICGV 96814, Doumbala and GM 663
147 recorded the lowest number of matured pods.

148 The genotypes F-Mix, NC 7, ICGV 86124, ICGV 86015, and PC 79-79 recorded the highest dry pod
149 weight while Chinese, GM 656, GM 120, and GM 324 which recorded the lowest dry pod weight (kg/ha).

150 **3.3 Dry haulm weight (kg/ha)**

151 The results indicated that there were significant differences ($P < 0.005$) dry haulm weight among the
152 groundnut genotypes (Fig 1).



153

154 **Fig 1 Effect of groundnut genotypes on dry haulm weight (kg/ha)**

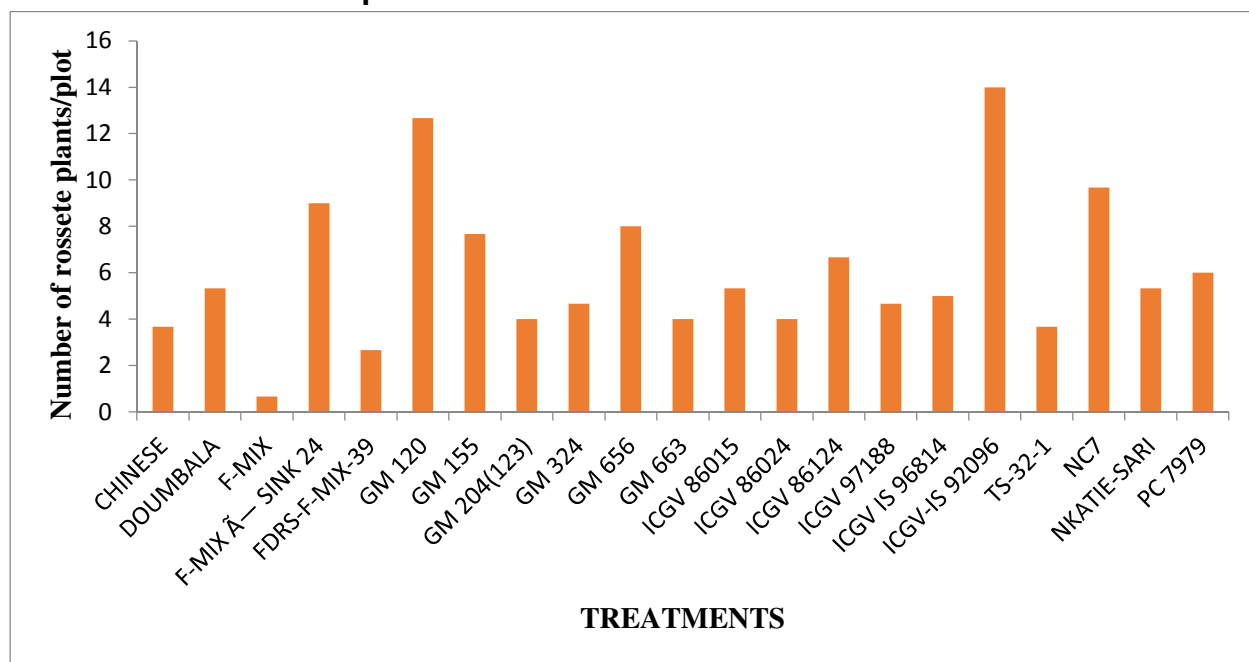
155

156 For dry haulm weight (kg/ha) as expected, the genotypes F-mix, NC 7, TS-32-1, ICGV 86015 and ICGV
 157 IS 92096 recorded the highest dry haulm weight (kg/ha) while Chinese, GM 204 (123) and GM 663
 158 recorded the lowest dry pod weight.

159

160

161 **3.4 Number of Rossete plants/10 m²**



162
163 **Fig 2 incidence of Rossete on the groundnut genotypes**

164
165 The results show no significant difference ($P > 0.05$) among the groundnut genotypes on the number of
166 Rossete plants/ 10 m² (Fig 2).

167 **4.0 DISCUSSION**

168 **4.1 Severity of Early and Late leaf spot disease score.**

169 Early and late leaf spot diseases are the most widespread foliar diseases of groundnut (*Arachis*
170 *hypogaea* L.) (Smith et al., 1992), accounting for pod yield losses of up to 50% in severe
171 epidemics (Jackson et al., 1969; Nutsugah et al., 1998, 2006; Tsigbey et al., 2001).
172 Management strategies for leaf spot epidemics rely on reducing initial inocula via crop rotation
173 or on reducing the rate of disease spread via resistant cultivars and fungicide applications
174 (Shokes and Culbreath, 1997). Crop rotation for 2 to 3 years is recommended, because this
175 may delay the development of a leaf spot epidemic by 2 to 3 weeks (Kokalis-Burelle, et al.,
176 1997). However, because of the rapid rate of increase of leaf spot diseases, crop rotation alone
177 is insufficient for control, hence the need for the identification of groundnut genotypes resistant
178 to leaf spot diseases.

179 NKATIE-SARI was released as a groundnut cultivar relatively resistant to both early and late
180 leaf spots compared to the susceptible cultivar Chinese and Doumbala by CSIR- Savanna
181 Agricultural Research Institute. The present study demonstrated that the groundnut genotypes
182 FDRS-F-MIX-39 and GM 663 were as resistant to early and late leaf spots as NKATESARI.

183 However, F-mix, ICGV-IS 92096, ICGV-IS 96814, F-mix × Sink 24, and PC 79-79 appeared to
184 be more resistant to early and late leaf spot than NKATIE-SARI.

185 The genotypes GM 120, GM 324 and TS 32-1 were highly susceptible to both early and late leaf
186 spots diseases.

187 **4.2 Total number of pods/plant**

188 As expected, the relatively resistant genotypes F-Mix, F-Mix × SINK 24, FDRS-F-Mix-39, NC 7, NKATIE-
189 SARI and PC 79-79 had the highest total number of pods/plant. The genotypes Chinese, GM 120 and
190 GM 324 recorded the lowest total number of pods/plant due to their susceptibility to *Cercospora* leaf spot
191 disease. Bdliya (2007) observed that *Cercospora* leaf spot is disastrous to groundnut especially towards
192 pod formation stage of the crop causing low seed and haulm yield. The diseases cause premature leaves
193 defoliation and a reduction in the photosynthetic area of the leaf surface which resulted in the low yield in
194 the susceptible genotypes (Shokes and Culbreath 1997; Waliyar *et al.* 2000). Another reason for the
195 reduced groundnut yield in the susceptible genotypes might be the shedding of numerous pods as the
196 weakened pegs break when the plants are dug and inverted (Hagen, 1998).

197 **4.3 Number of matured pods/plant**

198 The genotypes F-Mix, NKATIE-SARI, NC 7 and F-Mix × SINK 24 that showed some level of resistance to
199 leaf spots also had improved number of matured pods, apparently due to high production of
200 photosynthate.

201 **4.4. Dry pod weight**

202 They improved sink source relationship due to leaf retention in the relatively resistant genotypes also
203 resulted in increased dry pod weight.

204 **4.5. Dry haulm weight (kg/ha)**

205 The genotypes F-Mix, NC 7, TS-32-1, ICGV 86015 and ICGV 92096 recorded the highest dry haulm
206 weight because they exhibited some level of resistance to both early and late leaf spot disease as
207 compared to Chinese, GM 204 (123) and GM 663 which recorded the least dry haulm weight (kg/ha)
208 probably because of loss of leaves to early and late leaf spot disease. This agrees with Bdliya (2007) who
209 observed that *Cercospora* leaf spot is disastrous to groundnut especially towards pod formation stage of
210 the crop causing low seed and haulm yield.

211 **5.0 CONCLUSION AND RECOMMENDATION**

212

213 From the study, the genotypes F-Mix, NC 7, PC 79-79, F-Mix × SINK 24 and NKATIE-SARI were found to
214 be resistance to both early and late leaf spot disease whereas Chinese, Doumbala, GM 120, GM 324 and
215 ICGV 86015 were very susceptible to both diseases.

216 The study also revealed that the genotypes F-Mix, NC 7, PC 79-79, F-Mix × SINK 24 and NKATIE-SARI
217 recorded the highest total number of pods/plant as well as the highest number of matured pods/plant. For
218 dry pod weight, the genotypes F-Mix, ICGV 86124 and ICGV 86015 had the highest dry pod weight.

219 During the study there was high number of rosette plants observed, however there was no significant
220 difference among the genotypes.

221 The recommendation was that, farmers should use F-Mix, NC 7, F-Mix × SINK 24 and Nkatie-SARI
222 among others that exhibited some level of resistance to both early and late leaf spot diseases and were
223 also high yielding. This will save them the cost of fungicide spray against the diseases and also improve
224 their income from the increased yields. Also the susceptible genotypes especially Chinese that are widely
225 cultivated should be improved upon through further crosses with the resistant genotypes to enhance their
226 resistance to the leaf spot pathogen.

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