Inheritance of Resistance to Sorghum Shoot Fly, Atherigona soccata

M. K. Dhillon,* H. C. Sharma, B. V. S. Reddy, Ram Singh, and J. S. Naresh

ABSTRACT

The sorghum shoot fly, Atherigona soccata Rond. (Diptera: Muscidae), is one of the most important pests of sorghum [Sorghum bicolor (L.) Moench], and host plant resistance is an important component for the management of this pest. Most of the sorghum hybrids currently under cultivation are based on cytoplasmic malesterility (CMS). To develop a strategy to develop sorghum hybrids with resistance to shoot fly, we studied the nature of gene action for resistance to this pest in F1 hybrids derived from shoot fly-resistant and -susceptible CMS and restorer lines. The hybrids based on shoot flyresistant CMS and restorer lines were glossy and trichomed and had lower proportion of plants with eggs (78.5% vs. 88.4 to 93.3%) and deadhearts (40.8% vs. 60.8 to 75.3%) than the hybrids based on other cross combinations, suggesting that resistance is required in both CMS and restorer lines for obtaining shoot fly-resistant hybrids. Proportional contributions of CMS lines for oviposition, deadhearts, leaf glossiness, and recovery resistance were greater than those of the restorer lines. The general (GCA) and specific combining ability (SCA) estimates suggested that inheritance for oviposition nonpreference, deadhearts, recovery resistance, and the morphological traits associated with resistance or susceptibility to A. soccata were governed by additive-type of gene action. The SCA effects and heterosis estimates indicated that heterosis breeding would not be rewarding in breeding for resistance to shoot fly.

S ORGHUM is an important crop in Asia, Africa, USA, Australia, and Latin America. It is grown on about 10.4 million hectares in India, with an annual grain production of 8 million megagrams (FAO, 2002). The productivity levels under subsistence farming conditions are quite low (500-800 kg ha⁻¹) mainly because of biotic and abiotic constraints. More than 150 species of insect pests damage the sorghum crop, of which sorghum shoot fly is most important in Asia, Africa, and the Mediterranean Europe (Sharma, 1993). Losses in grain yield because of shoot fly damage average about 5% in India (Jotwani, 1983). The shoot fly females lay white, elongated, cigarshaped eggs singly on the undersurface of the leaves, parallel to the midrib. After egg hatching, the larvae crawl to the plant whorl and move downward between the folds of the young leaves until they reach the growing point. When they feed, they cut the growing tip and the result is drying of the central leaf called "deadheart."

Published in Crop Sci. 46:1377–1383 (2006). Crop Breeding & Genetics doi:10.2135/cropsci2005.06-0123 © Crop Science Society of America 677 S. Segoe Rd., Madison, WI 53711 USA A number of genotypes with resistance to shoot fly have been identified, but the levels of resistance are low to moderate (Sharma et al., 2003a). Plant resistance to sorghum shoot fly appears to be a complex trait and depends on the interplay of a number of component characters (Dhillon, 2004).

The discovery of CMS (Stephens and Holland, 1954) made it easier to incorporate desired traits into hybrids (House, 1985). Because more than 75% of the area under sorghum cultivation in India is planted to high-yielding hybrids, and most of these hybrids are based on milo-cytoplasm, it is important to transfer genes conferring resistance to sorghum shoot fly into cytoplasmic male-sterile (A-lines), maintainer (B-lines), and restorer (R-lines) lines to develop hybrids with high grain yield and resistance to this pest.

There is no information on the interaction between shoot fly-resistant and -susceptible A-, B-, and R-lines relative to the expression and inheritance of resistance to *A. soccata* in F_1 hybrids. Since future breeding efforts will largely focus on high-yielding, shoot fly-resistant hybrids, the present studies were performed to understand the nature of gene action for components that contribute to resistance or susceptibility to *A. soccata*. Such an information will be useful in developing an appropriate strategy to produce shoot fly-resistant hybrids for cultivation by the farmers in the semiarid tropics.

MATERIALS AND METHODS

Experimental Material

The experimental material consisted of 12 restorer, 12 CMS, and their maintainer (5 shoot fly-susceptible and 7 shoot fly-resistant) lines selected at random from germplasm and breeding material maintained in the gene bank at the International Crops Research Institute for the Semi-Arid Tropics, Patancheru, Andhra Pradesh, India. The prehybridization evaluation of CMS, maintainer, and restorer lines for reaction to sorghum shoot fly (determined on the basis of percentage deadhearts) during the 2002 rainy season supplemented with an ex-ante inference was used to categorize the test material into shoot fly-resistant and -susceptible groups. The 144 F₁ hybrids were produced by crossing 12 CMS with 12 restorer lines in a line \times tester mating design during the 2002–2003 post-rainy season.

The test material (12 A-, B-, and R-lines, and their 144 F_1 hybrids), along with shoot fly-resistant (IS 18551) and susceptible (Swarna) checks, was planted in a randomized complete block design (RCBD) in three replications during the 2003 rainy (July–November), early post-rainy (September–January), and late post-rainy (October–March) seasons using the interlard fish-meal technique (Soto, 1974). Each genotype was sown in four-row plots of 2-m row length; the rows were 75 cm apart. The seed was sown with a four-cone planter at a depth of 5 cm. The plants were thinned 1 wk after seedling emergence to maintain a spacing of 10 cm between plants.

Data were recorded in the central two rows on oviposition and deadheart formation at 14 d after seedling emergence

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(DAE). Recovery resistance was assessed on a scale of 1 to 9 (1 = 80%) plants with 2 to 3 uniform productive tillers; and 9 = <20% damaged plants with 1 to 2 productive tillers) at maturity. Data were also recorded on leaf glossiness on a scale of 1 to 5 [1 = highly glossy (light green, shining, narrow, and erect leaves),and 5 = nonglossy (dark green, dull, broad, and drooping leaves)] (Sharma and Nwanze, 1997). At 12 DAE, trichome density was recorded on central portion of the fifth leaf (from the base) from three seedlings in each genotype at random. The leaf pieces were cleared in acetic-lactic acid (2:1) (Maiti et al., 1980; Dhillon, 2004) and observed for the presence of trichomes and trichome density on the abaxial (lower) and adaxial (upper) surfaces of the leaves. The leaf sections were mounted on a slide in a drop of lactic acid and observed under a microscope at a magnification of 10×. Pigmentation of the plumule and leaf sheath was assessed on a 1-to-5 rating scale (1 = plumule or leafsheath with a deep pink pigment, 2 = plumule or leaf sheath with a pink pigment, 3 = plumule or leaf sheath with light pink pigment, 4 = plumule or leaf sheath with very light pink pigment, 5 = plumule or leaf sheath of green color) at 5 DAE. Chlorophyll content (g m⁻²) was measured with a chlorophyll meter (SPAD-502, Minolta Corporation) on the flag leaf (three leaves per plot) at 80 d after seedling emergence (Yamamoto et al., 2002). Waxy bloom was recorded on a scale of 1 to 5 (1 = stem)and leaves without a wax layer, and 5 = stem and leaves covered with a dense layer of wax) at 50% flowering.

Statistical Analyses

The data were subjected to analyses of variance. The parents were classified as resistant and susceptible on the basis of their reaction to shoot fly. Genetic analyses were performed via a line \times tester method, as suggested by Kempthrone (1957), using GenStat Release 6.0. The sum of squares due to F_1 hybrids was partitioned into sum of squares due to females, males, and females \times males, which was used to estimate the additive and dominance components of the variation. Simple correlations, multiple regression, and stepwise regression analyses were performed to understand the association between the morphological traits and resistance to sorghum shoot fly (Sharma et al., 2003b).

The main effects of CMS and restorer lines were equivalent to general combining ability (GCA), and the effects of a CMS line with a specific restorer were equivalent to specific combining ability (SCA) (Hallauer and Miranda, 1981). The heritability [Narrow-sense heritability = ${(Vgca)/(Vgca + Vsca + Vs$ VE)} \times 100; where, Vgca = general combining ability variance, Vsca = specific combining ability variance, and VE = error variance], and the proportional contribution of females, males, and their interaction [contribution of females = $\{SS (females)/$ SS (crosses) \times 100; contribution of males = {SS (males)/SS (crosses) \times 100; contribution of females \times males interaction = [SS (females \times males)/SS (crosses) \times 100]] to total variability of each trait/character was also computed.

RESULTS AND DISCUSSION

Oviposition Preference and Deadheart Formation

Deadheart formation due to shoot fly damage was greatest during the rainy season, followed by early postrainy, and the late post-rainy seasons, but there was no change in the relative resistance-susceptibility rankings of the material tested across seasons. The shoot flyresistant CMS lines were preferred for oviposition (76.3% plants with eggs) and suffered more deadheart incidence (40.1% deadhearts) than the maintainer lines (68.4% plants with eggs and 32.5% plants with deadhearts), while the differences among the shoot flysusceptible CMS and maintainer lines were not apparent (Table 1). Preference of CMS lines for oviposition and more deadhearts formation as compared with the maintainers suggested that the resistance-susceptibility to A. soccata was influenced by factors associated with male-sterility or fertility restoration cytoplasm in sorghum. Similar findings have earlier been reported in case of sorghum midge, Stenodiplosis sorghicola (Coq.) (Sharma et al., 1994). The hybrids based on shoot flyresistant CMS and restorer lines had significantly lower proportion of plants with eggs and deadhearts than the

Table 1. Reaction of cytoplasmic male-sterile (A), maintainer (B), and restorer (R) lines of sorghum and their F1 hybrids to Atherigona soccata, and the morphological traits associated with resistance to this insect across seasons (ICRISAT, Patancheru, India).

| Hybrid and parent groups | Plants with eggs (%) | Deadhearts (%) | Recovery resistance score† | Leaf glossiness score ‡ | Trichome density | | Pigmentation score§ | | Chlononhull | XX 7 |
|--|----------------------|----------------|-------------------------------|----------------------------|---------------------|---------|---------------------|-------------|-------------|--------------|
| | | | | | abaxial | adaxial | Plumule | Leaf sheath | content | bloom score¶ |
| RA# | 76.3 | 40.1 | 6.0 | 2.4 | 62.8 | 96.0 | 2.7 | 3.3 | 43.3 | 4.9 |
| RB†† | 68.4 | 32.5 | 5.6 | 2.0 | 79.2 | 120.2 | 2.7 | 3.4 | 41.2 | 4.9 |
| SA‡‡ | 88.4 | 64.5 | 7.1 | 4.8 | 0.2 | 0.4 | 3.6 | 4.1 | 47.7 | 5.0 |
| SB§§ | 90.6 | 67.4 | 7.3 | 4.8 | 4.6 | 10.6 | 4.1 | 4.3 | 47.2 | 4.9 |
| RR¶¶ | 74.3 | 39.3 | 4.4 | 1.7 | 94.8 | 137.6 | 2.0 | 2.5 | 45.2 | 4.4 |
| SR## | 90.6 | 69.6 | 6.4 | 4.0 | 1.6 | 3.7 | 3.1 | 4.1 | 46.3 | 5.0 |
| $\mathbf{RA} 	imes \mathbf{RR}$ ††† | 78.5 | 40.8 | 4.7 | 2.2 | 94.5 | 141.1 | 2.2 | 2.9 | 46.4 | 4.7 |
| $RA \times SR^{\dagger\dagger\dagger}$ | 88.4 | 60.8 | 5.7 | 3.9 | 19.8 | 31.7 | 2.7 | 3.8 | 47.5 | 4.9 |
| $SA \times RR^{\dagger\dagger}$ | 93.0 | 69.1 | 4.8 | 4.5 | 32.3 | 49.9 | 2.1 | 3.0 | 48.8 | 5.0 |
| $SA \times SR^{\dagger\dagger\dagger}$ | 93.3 | 75.3 | 5.8 | 4.7 | 1.2 | 1.9 | 3.1 | 4.0 | 49.0 | 5.0 |
| LSD ($P = 0.05$) | 12.99 | 5.50 | 0.76 | 0.54 | 12.35 | 16.37 | 0.67 | 0.61 | NS | NS |

* Recovery resistance score (1 = >80% plants with 2 to 3 uniform productive tillers, and 9 = < 20% damaged plants with 1 to 2 productive tillers). \ddagger Leaf glossiness score (1 = highly glossy, and 5 = nonglossy).

Present growth is a score (1 - dark green pink color, and 5 - light yellow color).
Waxy bloom score (1 = stem without wax layer, and 5 = stem densely covered with wax layer).

#RA = Shoot fly-resistant A-lines. **†† RB = Shoot fly-resistant B-lines.**

 $\ddagger SA = Shoot fly-susceptible A-lines.$ §§ SB = Shoot fly-susceptible B-lines.

¶¶ RR = Shoot fly-resistant R-lines.

SR = Shoot fly-susceptible R-lines.

††† Hybrid combinations.

hybrids on the basis of other cross combinations. Hybrids based on shoot fly-resistant CMS and-susceptible restorer lines had comparatively lower percentage of plants with eggs and deadhearts than the hybrids based on shoot fly-susceptible CMS and resistant or susceptible restorer lines. The hybrids based on shoot fly-susceptible CMS and shoot fly-resistant restorer lines were as susceptible as the hybrids based on shoot flysusceptible CMS and restorer lines, suggesting the influence of cytoplasmic factors on expression of shoot fly resistance in sorghum because of the interaction between cytoplasmic and nuclear genes (Dhillon, 2004). Similar results on the influence of CMS on genotypic susceptibility have also been reported by Xu and Song (1997) and Xu et al. (1998) for bacterial leaf blight, caused by Xanthomonas oryzae pv. oryzae (Ishiyama) Swings et al. = X. campestris pv. oryzae (Ishiyama) Dye in rice (Oryza sativa L.) and by Sharma et al. (1996) for sorghum midge in sorghum.

Recovery Resistance

Shoot fly-resistant CMS and maintainer lines showed better recovery resistance (score 5.6–6.0) as compared with the susceptible CMS and maintainer lines (score 7.1–7.3) (Table 1). Hybrids based on shoot fly-susceptible CMS and restorer lines showed poor recovery resistance than the hybrids based on other cross combinations. Varieties with high recovery resistance are known to yield more under shoot fly infestation (Rana et al., 1985).

Morphological Traits

Shoot fly-resistant CMS, maintainer, and restorer lines were glossy (score 1.7–2.4), while the shoot flysusceptible CMS, maintainer, and restorer lines were nonglossy (score 4.0–4.8) (Table 1). The hybrids based on the glossy CMS and restorer lines showed the same level of glossiness as the parents, whereas the hybrids based on the nonglossy CMS lines and glossy or nonglossy restorer lines were nonglossy. However, the hybrids based on glossy CMS lines and nonglossy restorer lines were intermediate in expression of leaf glossiness. Expression of leaf glossiness in F_1 hybrids was influenced more by the CMS than by the restorer lines, possibly because of interaction of cytoplasmic and nuclear genes (Dhillon, 2004).

The adaxial surface of the leaf had more trichomes than the abaxial leaf surface. The shoot fly-resistant CMS, maintainer, and restorer lines were trichomed, whereas the shoot fly-susceptible CMS, maintainer, and restorer lines were nontrichomed, except the restorer Swarna (Table 1). The hybrids based on the trichomed CMS and restorer lines had more trichomes compared with the parents. Hybrids based on nontrichomed CMS and trichomed restorer lines had fewer trichomes than the trichomed parent, except the hybrids based on restorer parent SFCR 125. The hybrids based on nontrichomed CMS and restorer lines were nontrichomed, while the hybrids based on trichomed CMS and nontrichomed restorer lines were nontrifor the hybrids based on ICSV 91011 and Swarna as restorer parents. The restorer lines showed a greater influence on expression of trichomes in the F_1 hybrids. The level of resistance to shoot fly was higher when both glossy and trichome traits occurred together (Agrawal and House, 1982; Dhillon et al., 2005).

The intensity of leaf sheath and plumule pigmentation was greater for shoot fly-resistant CMS, maintainer, and restorer lines as compared with the susceptible ones (Table 1). The hybrids based on tan-type (nonpigmented) CMS and restorer parents were also tan type. The restorer lines influenced expression of pigmentation in plumule and leaf sheath. Purple-pigmented sorghums have been reported to be tolerant to shoot fly damage (Singh et al., 1981), but genetically diverse sorghum material used in the present studies did not support this hypothesis. The chlorophyll content of the flag leaf at 80 DAE in shoot fly-resistant CMS and maintainer lines was lower than that of the shoot fly-susceptible CMS and maintainer lines (Table 1). There were no differences in the chlorophyll content of shoot fly-susceptible CMS, maintainer, and restorer lines, and the F_1 hybrids. Shoot fly-resistant lines showed lower chlorophyll content than the susceptible ones. Mate et al. (1996) also reported higher chlorophyll content in the shoot flysusceptible than the shoot fly-resistant sorghum genotypes. The differences were nonsignificant for waxy bloom among the hybrids and their parents.

Association between Shoot Fly Damage Parameters and Morphological Traits

Resistance to shoot fly (deadhearts, plants with eggs, and recovery resistance) was associated with leaf glossiness $(r = 0.45^{**}$ to $0.88^{**})$ (** P = 0.01), trichome density ($r = -0.50^{**}$ to -0.74^{**}), leaf sheath pigmentation ($r = 0.19^{**}$ to 0.54**), and waxy bloom ($r = 0.27^{**}$ to 0.41**). Chlorophyll content was significantly and positively associated with oviposition $(r = 0.42^{**})$ and deadhearts ($r = 0.45^{**}$). Pigmentation on plumule was positively associated with recovery resistance (r = 0.43^{**}). Leaf glossiness, trichome density on both the leaf surfaces, leaf sheath pigmentation, and waxy bloom were significantly and positively correlated ($r = 0.24^{**}$ to 0.96**) with one another. Chlorophyll content showed significant and negative association with leaf glossiness, trichome density, and waxy bloom. Morphological traits and plants with eggs explained 86.0% of the total variation in shoot fly deadhearts [Deadhearts (%) = - $74.4 + 0.27X_1 + 3.26X_2^* + 0.74X_3 - 0.76X_4 + 1.23X_5^* - 0.05X_6 + 0.02X_7 + 0.54X_8 (R^2 = 86.0\%) (* P = 0.05)]$ [where, X_1 = chlorophyll content; X_2 = leaf glossiness; X_3 = leaf sheath pigmentation; X_4 = plumule pigmentation; X_5 = plants with eggs; X_6 = trichome density (adaxial); X_7 = trichome density (abaxial); and X_8 = waxy bloom.]. Adjusted R-squared stepwise regression analyses indicated that leaf glossiness (X_2) , trichome density on adaxial surface of leaves (X_6) , and plants with eggs (X_5) explained 86.0% of the variation in deadhearts $[Deadhearts(\%) = -65.12 + 3.28X_2 + 1.31X_5 - 0.03X_6]$ $(R^2 = 86.0\%)$].

General Combining Ability (GCA) Effects

Oviposition Preference and Deadheart Formation

The mean squares for hybrids, CMS lines, restorers, CMS lines \times restorers, and for their interactions with the environments were statistically significant at P = 0.05 or 0.01 for percentage plants with eggs and deadhearts, except environments × CMS lines × restorers for deadhearts (Table 2). The GCA effects for oviposition preference and deadheart incidence for the shoot flyresistant CMS and restorer lines were significant and negative, except in few cases; whereas such effects for shoot fly-susceptible parents were positive (Table 3). Similar observations have been reported by Sharma et al. (1977) and Hallali et al. (1982). The parental performance was a good indicator of hybrid behavior as has been reported by Rao et al. (1974). The proportional contribution of CMS lines for percentage plants with eggs and deadhearts was greater than that of the restorer lines (Table 3). The narrow-sense heritabilities for oviposition and deadhearts were 15.0 and 12.8%, respectively, and estimates of additive variances for these traits were greater than their dominance variances (Table 4), suggesting the role of additive-type of gene action conditioning these traits. As oviposition nonpreference and deadheart formation being governed by additive-type of gene action, resistance in both male and female parents would be required for obtaining hybrids with resistance to shoot fly. Similar inheritance pattern of these traits was reported by Rao et al. (1974), Hallali et al. (1983), and Agrawal and Abraham (1985). Ravindrababu and Pathak (2000) suggested additive and epistatic gene effects for resistant \times resistant crosses whereas for resistant \times susceptible crosses additive, dominance, and epistatic (additive \times dominance) effects were important for resistance to shoot fly.

Recovery Resistance

The mean squares for recovery resistance for environments \times hybrids, environments \times CMS lines, and environments \times restorers were significant at P = 0.05 or

Table 2. Mean squares for shoot fly (*Atherigona soccata*) oviposition preference, deadhearts, and recovery resistance in F_1 sorghum hybrids and their parents (ICRISAT, Patancheru, India).

| Source of variation | df | Plants with eggs (%) | Deadhearts (%) | Recovery resistance |
|--|------|-------------------------|-------------------|------------------------|
| Environments (E) | 2 | 54 008.6** | 130 408.3** | 417.3** |
| Hybrids | 143 | 555.6** | 2246.7** | 7.7** |
| Males (M) | 11 | 3732.2** | 17069.6** | 11.8** |
| Females (F) | 11 | 1241.0** | 6751.8** | 59.1** |
| Males \times females | 121 | 200.2** | 489.7* | 2.7** |
| Environments \times hybrids | 284 | 209.4** | 219.9** | 2.6** |
| Environments \times females | 22 | 999.3** | 324.6** | 7.2** |
| Environments × males | 22 | 444.9** | 369.8** | 7.4** |
| $\mathbf{E} \times \mathbf{F} \times \mathbf{M}$ | 240 | 115.8** | 196.6 | 1.7 |
| Error | 854 | 69.1 | 158.0 | 1.6 |
| Proportional | cont | ribution (%) | to total variance | |
| Females | | 52.01 | 58.44 | 11.76 |
| Males | | 17.30 | 23.12 | 59.06 |
| Females $	imes$ males | | 30.69 | 18.44 | 29.18 |

* F test significant at 0.05 probability level.

** F test significant at 0.001 probability level.

| Table 3. G | eneral com | bining abili | ty (GCA) | effects of 12 |
|------------|-----------------------|--------------|----------------|----------------|
| cytoplasm | nic male-steri | le (CMS) an | d restorer lii | ies of sorghum |
| for shoot | t fly (Atheri | igona soccal | ta) ovipositi | on preference |
| (plants v | with eggs), | deadhearts, | and recov | ery resistance |
| (ÎCRISA) | F , Patancheru | ı, India). | | · |

| Genotypes | Plants with eggs (%) | Deadhearts (%) | Recovery resistance |
|---|----------------------|---------------------|---------------------|
| | Shoot fly-resist | tant CMS lines | |
| SPSFR 94011A | -4.68** | -8.41** | -0.26* |
| SPSFR 94006A | -4.32** | -8.98** | -0.19 |
| SPSFR 94007A | -2.20** | -5.74** | -0.13 |
| SPSFR 94010A | -0.96 | -2.73* | 0.64** |
| SPSFR 94034A | -4.86** | -10.07** | 0.12 |
| SP 55299A | -3.31** | -7.88 ** | -0.29** |
| SP 55301A | -10.36** | -21.69** | -0.01 |
| | Shoot fly-suscep | tible CMS lines | |
| SPSFR 94012A | 4.74** | 8.35** | -0.52** |
| 296A | 6.20** | 11.21** | 0.16 |
| Tx 623A | 7.40** | 17.94** | 0.44** |
| CK 60A | 6.47** | 15.40** | -0.18 |
| ICSA 42A | 5.87** | 12.60** | 0.22 |
| SE ± (GCA) | 0.80 | 1.21 | 0.12 |
| $SE \pm (g_i - g_j)$ | 1.13 | 1.71 | 0.17 |
| | Shoot fly-resista | nt restorer lines | |
| ICSV 705 | -4.70** | -5.56** | 0.24* |
| ICSV 700 | -0.99 | -2.07 | -0.80** |
| ICSV 708 | -1.44 | -4.48** | -0.83** |
| PS 30710 | -0.26 | -6.29** | -0.81** |
| IS 18551 | -4.49** | -8.60** | -0.45** |
| SFCR 151 | -3.16** | -7.08** | 0.73** |
| SFCR 125 | -2.38** | -7.71** | -0.95** |
| | Shoot fly-suscept | ible restorer lines | |
| ICSV 91011 | 1.90* | 3.21** | 0.08 |
| CS 3541 | 3.40** | 6.67** | 0.33** |
| MR 750 | 4.50** | 12.16** | 0.80** |
| ICSV 745 | 4.46** | 12.27** | 0.97** |
| Swarna | 3.16** | 7.49** | 0.73** |
| SE ± (GCA) | 0.80 | 1.21 | 0.12 |
| $\mathbf{SE} \pm (\mathbf{g}_i - \mathbf{g}_j)$ | 1.13 | 1.71 | 0.17 |

* GCA effects significant at P = 0.05.

** GCA effects significant at P = 0.01.

0.01 (Table 2). The proportional contribution of restorer lines was more than that of the CMS lines for recovery resistance. The GCA effects for recovery resistance were significant and negative for the CMS lines SPSFR 94011, SPSFR 94012, and SP 55299 but significant and positive for SPSFR 94010 and Tx 623 (Table 3). The GCA effects for recovery resistance of the shoot flyresistant restorer lines were significant and negative, whereas those of the shoot fly-susceptible restorers were

Table 4. Additive and dominance variances, and narrow-sense heritability for oviposition preference, deadhearts, recovery resistance, leaf glossiness, trichome density, plant pigmentation, chlorophyll content, and waxy bloom in sorghum (ICRISAT, Patancheru, India).

| | Genotypic | Naman aanaa | | |
|----------------------------|--------------|--------------|------------------|--|
| Traits | $\sigma^2 A$ | $\sigma^2 D$ | heritability (%) | |
| Plants with eggs (%) | 42.34** | 14.56** | 15.0 | |
| Deadhearts (%) | 211.50** | 36.86* | 12.8 | |
| Recovery resistance | 1.36** | 0.12 | 22.0 | |
| Leaf glossiness | 0.94** | 0.20* | 60.5 | |
| Trichome density (adaxial) | 3343.78** | 784.42** | 1.0 | |
| Trichome density (abaxial) | 1583.1 | 500.13** | 1.1 | |
| Plumule pigmentation | 0.54** | 0.13** | 45.7 | |
| Leaf sheath pigmentation | 0.50** | 0.06** | 39.9 | |
| Waxy bloom | 0.05** | 0.03** | 35.1 | |
| Chlorophyll content | 5.35** | 2.84** | 23.9 | |

* Significant at P = 0.05.

** Significant at P = 0.01.

significant and positive, except in a few cases. Additivetype of gene action was involved in recovery resistance (Table 4). However, earlier studies suggested predominance of additive, nonadditive, and epistatic (resistant \times susceptible) type of gene action for recovery resistance (Starks et al., 1970; Sharma et al., 1977; Borikar and Chopde, 1982).

GCA Effects for Morphological Traits Associated with Resistance to *Atherigona soccata*

The mean squares for hybrids, CMS lines, restorers, CMS lines \times restorers, and for their interactions with the environments were statistically significant at P = 0.05 or 0.01 for leaf glossiness and trichome density on abaxial and adaxial leaf surfaces, except the mean squares for environments \times CMS lines for trichomes on the abaxial leaf surface (Table 5). The proportional contribution of CMS lines for leaf glossiness was more than that of the restorer lines, whereas the reverse was true for trichome density. The GCA effects for leaf glossiness of shoot flyresistant CMS and restorer lines were significant and negative, whereas those for the shoot fly-susceptible CMS and restorer lines were significant and positive (although there were a few exceptions), but the reverse was true for trichome density (Table 6). Leaf glossiness has been reported to be controlled by a single recessive gene (Tarumoto, 1980) and is simply inherited (Agrawal and House, 1982). Trichomes and leaf glossiness are independently inherited and apparently have an additive effect in reducing shoot fly incidence (Hallali et al., 1982; Maiti et al., 1984). The narrow sense heritability for leaf glossiness was 60.5 and 1.0% for trichome density.

The mean squares for CMS lines, restorers, and CMS lines \times restorers were significant at P = 0.05 or 0.01 for plumule and leaf sheath pigmentation, waxy bloom, and chlorophyll content (Table 5). The proportional contribution of restorer lines for pigmentation of plumule and leaf sheath, and waxy bloom was more than that of CMS lines, whereas the reverse was the case for chlorophyll content. The GCA effects for pigmentation of the plumule

and leaf sheath of the shoot fly-resistant CMS and restorer lines were significant and negative, whereas those of shoot fly-susceptible CMS and restorers were significant and positive, except in a few cases (Table 6). The GCA effects for chlorophyll content in the CMS lines SPSFR 94010, SP 55299, and SP 55301, and the restorer lines IS 18551, SFCR 151, and ICSV 745 were significant and negative, whereas those for the CMS lines Tx 623 and CK 60, and the restorer lines PS 30710, SFCR 125, CS 3541, and Swarna were significant and positive (Table 6). The GCA effects for waxy bloom in the CMS lines SPSFR 94007, SP 55299, and SP 55301, and the restorer lines ICSV 700 and IS 18551 were significant and negative, whereas those for Tx 623A, CK 60A, ICSA 42A, and Swarna were significant and positive (Table 6). The narrow-sense heritability for plumule and leaf sheath pigmentation, chlorophyll content, and waxy bloom varied between 23.9 to 45.7%, and estimates of additive variances for these traits were greater than their dominance variances (Table 4), suggesting the role of additive-type of gene action conditioning these traits.

Specific Combining Ability (SCA) Effects

The SCA effects for oviposition and deadhearts were nonsignificant, and there was positive midparent heterosis for oviposition preference and deadheart incidence, except in a few cases, suggesting that heterosis breeding would not be rewarding in breeding for resistance to shoot fly.

The SCA effects for recovery resistance were nonsignificant and showed positive heterosis, suggesting the involvement of additive-type of gene action for recovery resistance. The SCA effects for glossiness and trichome density were also nonsignificant.

CONCLUSIONS

Preference of CMS lines for damage by shoot fly as compared with the maintainers suggested the influence of factors associated with male-sterility or fertility

Table 5. Mean squares for leaf glossiness, trichome density, plant pigmentation, chlorophyll content, and waxy bloom in F₁ sorghum hybrids and their parents (ICRISAT, Patancheru, India).

| | | | Trichome density | | Pigmentation | | | |
|--|------------|-----------------|------------------|-----------------|--------------|-------------|------------------------|------------|
| Source of variation | df | Leaf glossiness | Abaxial | Adaxial | Plumule | Leaf sheath | Chlorophyll content | Waxy bloom |
| Environments (E) | 2 | 10.7 | 32 299.4** | 6177.7 | - | _ | - | - |
| Hybrids | 143 | 10.0** | 19 147.7** | 36 064.4** | 5.0** | 0.1 | 335.4** | 0.3 |
| Males | 11 | 66.2** | 66 411.6** | 137 577.6** | 11.4** | 11.5** | 103.0** | 1.3** |
| Females | 11 | 40.0** | 116 518.6** | 239 879.9** | 10.0** | 7.8** | 135.3** | 0.7** |
| Males $	imes$ females | 121 | 2.2** | 5977.5 ** | 8164.4** | 0.9** | 0.5** | 22.9** | 0.2** |
| Environments $	imes$ hybrids | 284 | 0.6** | 2397.1** | 2569.6** | † | † | † | † |
| Environments × females | 22 | 2.3** | 2141.7 | 2681.6** | † | † | † | † |
| Environments $	imes$ males | 22 | 1.0** | 4705.5** | 2513.1** | † | † | † | † |
| $\mathbf{E} \times \mathbf{F} \times \mathbf{M}$ | 240 | 0.4* | 2211.2** | 2569.5** | † | † | t | † |
| Error | 854 (334)‡ | 0.4 | 1476.3 | 1104.7 | 0.5 | 0.3 | 14.3 | 0.1 |
| | | Prop | ortional contrib | ution (%) to to | tal variance | | | |
| Females | | 50.91 | 26.71 | 29.44 | 32.30 | 31.14 | 27.64 | 17.82 |
| Males | | 30.73 | 46.85 | 51.34 | 37.00 | 46.25 | 21.04 | 32.49 |
| Females $	imes$ males | | 18.36 | 26.44 | 19.22 | 30.70 | 22.61 | 51.32 | 49.69 |

* F test significant at 0.05 probability level.

** F test significant at 0.001 probability level.

† ANOVA for single season.

* Value in the parenthesis is error degrees of freedom for pigmentation, waxy bloom, and chlorophyll content.

| | | Trichome density | | Pign | entation | | |
|----------------------|-----------------|------------------|-----------------|--------------------|-------------|---------------------|------------|
| Genotypes | Leaf glossiness | Abaxial | Adaxial | Plumule | Leaf sheath | Chlorophyll content | Waxy bloom |
| | | | Shoot fly-res | istant CMS line | s | | |
| SPSFR 94011A | -0.49** | 36.49** | 46.75** | -0.44** | -0.41** | 0.11 | 0.01 |
| SPSFR 94006A | -0.73** | 21.52** | 38.37** | -0.45** | -0.48** | -0.66 | -0.07 |
| SPSFR 94007A | -0.58** | 18.14** | 21.12** | -0.37** | -0.42** | 0.62 | -0.18** |
| SPSFR 94010A | -0.43** | 1.68 | 6.46* | 0.52** | 0.30** | -1.53* | 0.04 |
| SPSFR 94034A | -0.57** | 23.48** | 35.33** | -0.18 | 0.17 | -0.47 | 0.09 |
| SP 55299A | -0.62** | 12.61** | 24.72** | -0.62** | -0.59** | -1.32* | -0.24** |
| SP 55301A | -0.95** | 17.28** | 18.34** | 1.17** | 0.88** | -2.68** | -0.21** |
| | | | Shoot fly-susc | eptible CMS lin | es | | |
| SPSFR 94012A | 0.77** | -29.11** | -42.42** | -0.42** | -0.20* | -1.15 | 0.09 |
| 296A | 0.94** | -23.37** | -34.29** | 0.38** | 0.02 | 1.07 | 0.09 |
| Tx 623A | 0.89** | -24.17** | -36.03** | 0.08 | 0.21* | 1.24** | 0.12* |
| CK 60A | 0.82** | -18.26** | -26.4** | -0.01 | -0.15 | 5.00** | 0.12* |
| ICSA 42A | 0.94** | -36.29** | -51.97** | 0.34** | 0.67** | -0.24 | 0.12* |
| SE ± (GCA) | 0.06 | 3.70 | 3.20 | 0.11 | 0.10 | 0.63 | 0.05 |
| $SE \pm (g_i - g_j)$ | 0.08 | 5.23 | 4.52 | 0.16 | 0.14 | 0.89 | 0.07 |
| - | | | Shoot fly-resis | stant restorer lin | es | | |
| ICSV 705 | -0.46** | 11.01** | 15.38** | -0.06 | -0.06 | -1.12 | -0.02 |
| ICSV 700 | -0.34** | 45.79** | 56.07** | -0.83** | -0.60** | -0.68 | -0.32** |
| ICSV 708 | -0.47** | 44.88** | 72.3** | -0.15 | -0.65** | 0.83 | 0.09 |
| PS 30710 | -0.49** | 27.11** | 34.86** | -0.09 | -0.51** | 2.43** | 0.09 |
| IS 18551 | -0.77** | 33.23** | 45.32** | -0.31** | -0.40** | -3.29** | -0.46** |
| SFCR 151 | 0.05 | 5.58 | 17.91** | 0.10 | 0.10 | -1.67** | 0.09 |
| SFCR 125 | -0.64** | 0.32 | 2.97 | -0.51** | -0.51** | 1.41* | 0.07 |
| | | | Shoot fly-susce | ptible restorer li | nes | | |
| ICSV 91011 | 0.16** | -23.92** | -36.82** | 0.96** | 0.74** | 0.41 | 0.09 |
| CS 3541 | 0.66** | -34.96** | -50.12** | 0.38** | 0.55** | 1.17* | 0.09 |
| MR 750 | 0.72** | -37.73** | -54.93** | -0.37** | 0.05 | 0.67 | 0.07 |
| ICSV 745 | 0.68** | -37.38** | -56.08** | 1.08** | 1.08** | -1.74** | 0.07 |
| Swarna | 0.92** | -33.92** | -46.86** | -0.20 | 0.21 | 1.59* | 0.12* |
| SE ± (GCA) | 0.06 | 3.70 | 3.20 | 0.11 | 0.10 | 0.63 | 0.05 |
| $SE \pm (g_i - g_i)$ | 0.08 | 5.23 | 4.52 | 0.16 | 0.14 | 0.89 | 0.07 |

Table 6. General combining ability (GCA) effects of 12 cytoplasmic male-sterile (CMS) and restorer lines of sorghum for leaf glossiness, trichome density, plant pigmentation, chlorophyll content, and waxy bloom (ICRISAT, Patancheru, India).

* GCA effects significant at P = 0.05.

** GCA effects significant at P = 0.01.

restoration cytoplasm on resistance–susceptibility to *A.* soccata in sorghum. The equal levels of susceptibility in case of shoot fly-susceptible CMS and shoot fly-resistant or susceptible restorer lines-based hybrids suggested the influence of cytoplasmic factors on expression of shoot fly resistance in sorghum because of interaction of cytoplasmic and nuclear genes. Resistance is required in both male and female parents for obtaining hybrids with resistance to *A. soccata*. Studies on nature of gene action and regression analyses indicated that expression of leaf glossiness, trichomes, plants with eggs, and deadhearts in F_1 hybrids was conditioned by additive-type of gene action, and these can be used as morphological markers to select for resistance to sorghum shoot fly.

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