

Phenotyping of Sorghum Lines for Resistance to African Stem Borer (*Sesamia calamistis*)

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Abstract

Stem borer (*Sesamia calamistis*) is a serious insect pest of sorghum (*Sorghum bicolor*) resulting in grain yield losses ranging between 15-80%. However, genotypes showing complete resistance to these borers have not been identified in Nigeria. Utilization of resistant varieties in combination with other methods of control would offer a sustainable strategy for *S. calamistis* management in sorghum production. The objective of this study was to validate the acclaimed resistance in the materials received from Kenya and India in Nigerian environment and to screen and ascertain the status of some Nigerian Sorghum to *Sesamia calamistis*. Eighty-eight sorghum lines were artificially infested with the eggs of the stem borers at two different environment (Field and Screen House) using alpha-lattice design, consisting of 11 plots in eight blocks, replicated twice. Data were collected on leaf feeding, number of dead-hearts, cumulative stem tunnel length, number of exit holes, and selected agronomic traits. There were significant ($p < 0.01$) differences among the test genotypes for all the traits measured. Based on the selection index, 15% of genotypes were categorized as resistant, 42% as moderately resistant, 33 as moderately susceptible and 10% as susceptible. 13 genotypes showed resistance across the environments (field and screen house): ICSB464, ICSL71086, SSV20041-2YELLOW, ICSL71018, ICSR94032, ICSV700, ICSL71193, ICSR94030, ICSL71253, ICSL71268, ICSL71023, ICSL71061 and ICSL71137 were resistant with selection index ranging from 0.0 to 0.5. These sorghum lines with various resistance to *S. calamistis* could be used as source of resistance and as parents in sorghum improvement programme in breeding for resistance to stem borer.

Keywords: Genotypes, novel source of resistance, *Sesamia calamistis*, *Sorghum bicolor*

DOI: 10.7176/JBAH/11-18-04

Publication date: September 30th 2021

1. Introduction

Sorghum [*Sorghum bicolor* (L.) Moench] is an important food crop and ranks fifth after wheat, rice, maize, and barley in total area of production globally (Kuhlman, *et al.*, 2010). Nearly 80% of the cultivated area lies in Asia and Africa and widely cultivated for food, forage, ethanol, and sugar production (Liu *et al.*, 2009). Sorghum production especially in tropical Africa is curtailed by a number of important anthropod pests, with the stem borers belonging to lepidoptera being the most important. Stem borers cause grain yield losses ranging from 15-80% depending on crop variety, phenological stage and agro ecological environment (Karaya *et al.*, 2009, Muturi *et al.*, 2012). The damage caused are leaf feeding, dead heart formation, exit holes and stem tunneling damage (Kishore *et al.*, 2007; and Muturi *et al.*, 2012). The spotted stem borer (*Chilo partellus* (Swinhoe)) Pyralidae, African stem borer (*Busseola fusca* Fuller), and African pink borer (*Sesamia calamistis*) are among the most damaging insect pests that greatly reduce sorghum grain yield in African environments (Sharma *et al.*, 2005; Mwimali *et al.*, 2015). Among the several stalk borer species, pink stem borer (*Sesamia calamistis* (Hampson)) is the most important pest of sorghum in the Nigerian savannah (Ajayi 1998, Anaso and Thilza, 2006). Grain yield loss of about 49% was reported in Nigeria (Ajayi 1991., Ajayi 1997). Total crop failure had also been reported in a few instances (Emmanuel and Chinwe, 2008). There is, however, limited germplasm with resistance to these pests. Several options for managing sorghum borers have potential to mitigate their damaging effects, but each option has its own limitations. Host plant resistance forms an important part of integrated pest management as it provides inherent control without environmental issues and is compatible with other pest management approaches (Singh *et al.*, 2012). Effective breeding methods for resistance to borer damage could, therefore, be designed by plant breeders using both improved and new sources of stem borer resistance.

Management of *S. calamistis* in sorghum has mainly focused on cultural control, burning of crop residues, intercropping and predominantly the use of pesticides (Amsalu *et al.*, 2008). However, use of these strategies would invariably increase the cost of cultivation of sorghum, which is not a feasible option for the resource poor farmers of the semi-arid tropics. Furthermore, the use of chemical pesticides could be harmful to both the environment and human health. Hence, the exploitation of host plant resistance is the only viable option both in terms of economic and environmental sustainability for controlling stem borer in sorghum (Tadele *et al.*, 2011). Considering multiple stem borer damage traits are useful since resistance to stem borers is quantitatively

inherited thus selecting for resistance based on a single parameter would not be effective. Although there have been some success through conventional breeding approaches in improving plant resistance to stem borer, but only low to moderate levels of resistance have been observed in the cultivated germplasm (Singh *et al.*, 2011) and the identified resistant sources in germplasm possess many undesirable agronomic characters (Singh and Rana, 1986).

High yielding varieties in Nigeria are susceptible to stem borers. Thus, there is the need to increase the levels of tolerance in elite genotypes without sacrificing grain and Stover yield. Much of the existing sorghum germplasm in Africa and Asia have not been evaluated for resistance to stem borers (Dhillon and Sharma, 2012). Therefore, it is important to identify sorghum genotypes with high levels of resistance and diversify the bases of resistance through screening more germplasm (Singh *et al.*, 2011). In Nigeria, little research attention has been accorded to stem borers like *S. calamistis* in cereals and there is paucity of information on the status (resistance/susceptibility) of the elite sorghum germplasm. Screening of all the elite germplasm becomes necessary in the present investigation as the first phase of resistance breeding programme. Such information will be useful in developing an appropriate strategy to produce stem borer-resistant open pollinated varieties and hybrids for cultivation by the farmers in the semi-arid tropics. Therefore, the objective of this study was to assess the levels of resistance of elite sorghum germplasm obtained from ICRISAT Kenya and India and Institute for Agricultural Research (IAR) released sorghum varieties.

MATERIALS AND METHODS

Experimental Sites

The research was conducted at the IAR screen house and research farm at Samaru, Zaria, Nigeria. The farm is located on 11°11'N, 7°38'E, 640 m asl, 1200 mm annual rainfall) in the Northern Guinea Savannah.

Experimental Procedures

Eighty-eight sorghum lines comprising forty recombinant inbred lines (RILs) from ICRISAT Nairobi, Kenya, eight varieties from ICRISAT Patancheru, India, twenty-eight elite germplasm lines from Nigeria and twelve IAR released varieties. These lines were planted in IAR research farm under rain-fed condition. These lines were arranged in an alpha-lattice design, consisting of 11 plots in eight blocks, replicated twice. Each plot consisted of a row, 5 m long; the inter and intra row spacing was 75 cm and 30 cm, respectively. Three seeds were planted and later thinned to two plants per stand after two weeks of sowing. At 16 days after sowing, the crops were sprayed with cypermethrin (synthetic pyrethroid) to minimize shoot fly infestation, since this insect interferes with screening for resistance to stem borers (Makueti *et al.*, 2012).

Field and Screen House Validation for Stem Borer Resistance Status

Twenty one days after sowing, plants were artificially infested with 50g of stem borer eggs placed between the stem and the last unfolded leaf sheath using a forcep. Eggs of *S. calamistis* used in this study were obtained from the International Institute for Tropical Agriculture (IITA) Ibadan. To avert drowning of eggs in the water held in leaf whorls, sorghum seedling whorls were tapped gently before infestation. Infestations were carried out early in the morning to encourage egg survival. All other recommended cultural practices were observed. At the screen house, the same source of eggs of *S. calamistis* used on the field were used. One sorghum line/variety was sown in one pot repeated twice. Three seeds were planted and later thinned to two plants per pot after 2 weeks of sowing. Infestation was carried out as described for field above.

Parameters scored for Stem borer damage

On the field, 4 plants within each row were tagged for infestation while in the screen house two plants per pot were sampled and data were taken systematically from the marked plants. The observations were recorded on per plant basis at two and four weeks after the artificial infestation. Percentages of plants with leaf damage were computed by expressing the number of plants showing pinholes damage as a percentage of the total number of plants sampled (Muturi *et al.*, 2012). Dead heart incidence was computed by counting the number of plants showing dead heart damage and expressed as a percentage of the total number of plants sampled (Kumar *et al.*, 2005). At harvest, numbers of stem borer exit holes on the stem were counted on each sampled plant. The main stem of plants infested was split open from the base to the apex, and the cumulative tunnel length measured with a ruler in centimeters. Seedling vigour was scored at 2 weeks after sowing on a scale of 1 – 5, where 1 = low vigour (plants showing minimum growth, less leaf expansion and poor adaptation; 3=Moderate vigor; 5=high vigor tall plants with expanded leaves and robustness) (Kishore *et al.*, 2007, Muturi *et al.*, 2012). At physiological maturity, plant height was measured in centimeter from the base of the plant to the tip of the panicle using a calibrated pole. Days to panicle emergence was recorded as the number of days from the date of sowing to the date when 50% of panicle emerged in a plot. Days to 50% flowering was recorded as the number of days from the date of sowing to the date of anthesis of 50% of plants in a plot. After harvest, sorghum

panicles were sun-dried and hand threshed. Grain yield and hundred seed weight were recorded in grams for each of the sampled plants using an electric weighing balance.

Statistical Analysis

Data were first analyzed on individual location basis and combined across the 2 environments. The screen house and the field were equated to 2 environments. Data on percentages were angular transformed while those of counts were log transformed before the analysis of variance (Kishore *et al.*, 2007). The mean values of all the traits for each replicate were used to compute the analysis of variance using SAS GLM version 9.2. Treatment means were compared using a protected Fishers' least significant difference (LSD) test at $P = 0.05$. Selection index was calculated based on leaf damage (2nd and 4th week), dead heart (2nd and 4th week), stem tunneling and exit holes by adding the ratios between the values for each genotype and the overall mean for each parameter, and divide by 6 (number of damage parameters considered) (Tadele *et al.*, 2011). Genotypes were grouped into four categories namely resistant, moderately resistant, moderately susceptible and susceptible (Tadele *et al.*, 2011, Muturi *et al.*, 2012). The genotypes with selection index values less than 0-0.5 were regarded as resistant, 0.6-1.0 moderately resistance, 1.1-1.5 as moderately susceptible and those with a selection index greater than 2.1 as susceptible (Bergvinson *et al.*, 2004; Tadele *et al.*, 2011 and Muturi *et al.*, 2012).

RESULTS AND DISCUSSION

There were highly significant ($P = < 0.01$) differences for seedling vigor, heading date, days to 50% anthesis, plant height, panicle length, 100 grain weight, grain yield, percentage dead heart at 4 week, percentage leaf damage at 2 week, exit holes and tunnel length among the genotypes tested in the screen house (data not shown). In the field, mean squares due to genotypes were highly significant ($P = < 0.01$) for all the traits except for dead heart at 2 week and tunnel length (data not shown). For the combined analysis, environment mean squares were significant ($P = < 0.05-0.01$) differences for all traits except for dead heart at 4 week, exit holes and yield (Table1). There were highly significant ($P = < 0.01$) differences among genotypes for all traits measured except for leaf damage at 4 weeks, dead heart at 2weeks and tunnel length. The interactions between environment and genotype were significant for leaf damage at 2 week, dead heart at 2 and 4 weeks but highly significant ($p < 0.01$) for seedling vigor, days to 50% flowering, plant height, panicle length, 100 grain weight and exit hole (Table1).

Mean Performances

Across the two environments, the mean performance for leaf damage at 2 week ranged from 1.9% for KAT487 to 52.2% for Samsorg17 with a mean of 34.2%. (Table 2). Leaf damage at 4 weeks range from 0.0% for ICSL71086, ICSR94032 to 48.2% for Samsorg 3 with an average of 16.2%. Genotypes observed to show low leaf damage at 4 weeks were ICSL71086, ICSR94032, ICSL71268, SSV20041-1 White and ICSB464 (in increasing order of leaf damage incidence). Dead heart at 2 weeks ranged from 0.0 % to 64.8% with an average of 8.8%. Samsorg 17 recorded the highest number of dead heart (64.8%) at 2 weeks. At 4 weeks, it ranged from 0.0% to 100% with a mean of 26.5%. Genotypes ICSB 464, ICSV700, ICSL71193, ICSR94030, ICSL71253, ICSL71023 and ICSL71018 suffered the least dead heart damage at 4 weeks. ICSR94032, ICSL71086 and SSV20041-2 Yellow recorded fewer exit holes of 0.5, 1.3 and 1.5 respectively. Stem tunneling length range from 10.8cm for ICSL71137 to 97.5cm for SSV 20041-1 White, with a mean of 49.4cm.

Seedling vigour scores ranged from 2.0 to 4.0 with an average of 3.0. Genotypes ICSL71018, ICSL71253, ICSL71268, ICSL71005, SAMSORG 3, SAMSORG 8, SAMSORG17, SAMSORG42 and SAMSORG 44 had greater vigour scores (were more vigorous) than ICSR94030, ICSL71193 and ICSL71061 (Table2). Days to panicle emergence (HD) were longest for SAMSORG 17 (92 days) while ICSL71061 took 68 days for the panicle to emerge. The mean heading date observed was 76 days. Days to 50% flowering ranged from 70 to 94 days for ICSL71061 and SAMSORG 17 respectively with a mean of 78 days. Plant height ranged from 93.8 cm for ICSL 71137 to 223.4 cm for Samsorg8 with a mean of 143 cm. Panicle length ranged from 14.2 cm for ICSV700 to 25.1cm for Samsorg8 and closely followed by

Table 1. Combined means squares for the reaction of sorghum to stemborer (*S. calamistis*) and agronomic traits.

Source of variation	DF	SV	AN	HD	PLT HT, cm	PAL, cm	100-GW (g)	Yield, (g)	LD2, %WK	%LD4, %WK	DH2, %WK	DH, %4WK	E H	TL cm
Loc	1	1.16**	819.9**	946.23**	25170.6**	66.04*	4.33**	2785.0	7.79**	0.75**	2.27**	1.56	0.05	24951.7**
Rep (Loc)	2	0.13	5.11	1.933	358.6	3.20	0.38*	1772.0	0.02	0.04	1.80**	3.74**	0.04	605.4
Block (Loc x Rep)	28	0.23	3.93	3.353	318.9	5.65	0.07	652.0	0.05	0.03	0.09	0.14	0.08	1427.1
Entry	87	1.58**	112.7**	115.07**	2926.9**	45.05**	0.99**	1537**	0.09**	0.04	0.12	0.34**	0.15**	1077.8
Loc x Entry	87	0.64**	27.6**	30.197**	1003.7**	23.84**	0.24**	1067.0	0.08*	0.04	0.13*	0.22*	0.10**	942.8
Error	14	0.17	3.8	2.6414	225.4	8.16	0.08	804.1	0.05	0.03	0.09	0.16	0.06	873.2

*and ** significance at 0.05 and 0.01 levels of probabilities, respectively.

SV=Seedling Vigor, HD=Heading Date, AN=Days to 50%Anthesis, PL HT=Plant Height (cm), PAL=Panicle Length (cm),100GW=100Grain Weight(g), Yield(g),%DH2WKS= Percentage dead heart at 2 weeks, %DH4WKS = Percentage dead heart at 4 weeks,%LD2WKS= Percentage leaf damage at 2 weeks,%LD4WKS=Percentage leaf damage at 4 weeks, EH=Exit Holes and TL=Tunnel Length in centimeters.

Samsorg 44(24.9cm) with an average of 19.5cm.Hundred grain weight ranged from 0.2 g for ICSL 71219 to 4.1 g for Yar Washa with a mean was 2.1g. High grain yield (> 59 g) were observed on ICSL71137, Samsorg17, Samsorg72.3, Samsorg44 and SSV20041-1 White while lower grain yield (<20g) was recorded on ICSL71018, KAT487, and ICSL71086.

Table 2. Mean Performance of 13 resistant and nine Susceptible sorghum genotypes to *Sesamia calamistis* and some agronomic traits evaluated under infestation across two environments in 2014.

Line	LD2W, % 2WK	LD4W,% 4WK	DH2W, % 2WK	DH4W,% 4WK	EH	TL, cm	SI	CAT	SV	HD	AN	PLANTHT, cm	PAL, cm	GW, kg	Yield, kg
ICSB464	28.4	5.0	0.0	0.0	2.7	29.0	0.2	R	3.0	81.1	81.8	125.4	19.8	1.8	54
ICSL71086	17.2	0.0	0.0	25.4	1.3	20.2	0.2	R	3.0	71.9	74.8	100.7	17.5	2.3	17.7
SSV20041-2 YELLOW	5.1	8.6	0.0	9.6	1.5	51.0	0.3	R	3.0	87.0	88.4	196.5	17.3	3.0	40.5
ICSL71018	34.6	8.6	0.0	0.4	5.8	21.7	0.3	R	4.0	75.6	77.8	119.5	17.0	2.4	8.7
ICSR94032	26.7	0.0	0.0	8.7	0.5	45.5	0.4	R	2.0	89.0	90.1	141.9	21.3	1.6	42
ICSV700	44.1	13.0	0.0	0.0	5.2	18.2	0.4	R	3.0	82.9	85.2	155.9	14.2	1.8	34.6
ICSL71193	44.7	11.6	0.0	0.0	5.8	42.8	0.5	R	2.0	72.0	74.6	132.0	19.6	2.1	29.7
ICSR94030	40.8	17.1	0.0	0.0	5.2	33.2	0.5	R	3.0	78.6	79.1	113.2	24.8	1.3	31.4
ICSL71253	22.3	13.9	0.0	0.0	10.0	39.7	0.5	R	4.0	73.4	75.4	129.9	20.5	2.2	33.3
ICSL71268	24.8	0.2	0.0	11.0	6.9	37.9	0.5	R	4.0	69.2	70.9	125.0	15.8	1.8	43.3
ICSL71023	25.8	9.6	0.0	0.0	5.9	41.9	0.5	R	2.0	73.2	76.4	115.7	17.1	2.2	41.4
ICSL71061	40.6	6.1	0.0	8.9	6.8	50.4	0.5	R	3.0	67.9	69.9	135.6	17.0	1.7	22.5
ICSL71137	29.7	17.1	0.0	34.8	4.1	10.8	0.5	R	3.0	76.5	78.1	93.8	15.8	1.8	59.2
SAMSORG44	37.3	9.9	32.3	49.2	6.6	53.4	1.6	S	4.0	75.9	77.5	141.7	24.9	2.6	82.1
SAMSORG8 (KSV14)	34.0	28.5	28.0	59.6	3.6	26.5	1.6	S	4.0	84.0	85.7	223.4	25.1	3.2	72.3
SSV20041-1 WHITE	21.6	2.2	33.0	53.4	8.2	97.5	1.6	S	3.0	86.1	87.2	204.8	22.9	3.3	90.6
SAMSORG3 (KSV4)	38.8	48.2	28.0	47.1	4.4	45.5	1.8	S	4.0	75.3	77.4	154.7	20.6	1.9	56.1
ICSL71005	43.3	13.9	30.6	69.7	7.3	83.5	1.9	S	4.0	72.4	74.8	145.6	20.6	1.9	31.2
MAIMADARA-2	47.9	42.4	33.7	55.0	3.5	27.2	1.9	S	3.0	74.8	76.0	147.3	21.6	1.8	52.0
SAMSORG42	22.1	19.2	42.4	69.9	7.7	72.4	2.0	S	4.0	88.4	90.1	163.1	22.3	3.1	58.9
KAT487	1.9	25.1	53.8	86.2	2.8	44.6	2.1	S	3.0	86.6	88.8	154.5	24.6	2.5	16.6
SAMSORG17	52.2	22.2	64.8	100.0	3.1	57.9	2.7	S	4.0	92.2	94.1	142.7	18.3	3.0	60.2
Mean	34.2	16.2	8.8	26.5	5.6	49.4	1.0		3.3	76.1	78.0	143.0	19.5	2.1	43.2
CV	51.5	84.9	229.7	95.1	50.6	59.7			12.5	2.1	2.5	10.4	14.6	13.3	65.7
LSD	34.3	27	39	49.3	5.6	57.9			0.8	3.2	3.8	29.4	5.6	0.5	55.6

SV=Seedling Vigor, HD=Heading Date, AN=Days to 50%Anthesis, PL HT=Plant Height (cm), PAL=Panicle Length (cm),100GW=100Grain Weight(g), Yield(g),%DH2WKS= Percentage dead heart at 2 weeks, %DH4WKS = Percentage dead heart at 4 weeks,%LD2WKS= Percentage leaf damage at 2 weeks,%LD4WKS=Percentage leaf damage at 4 weeks, EH=Exit Holes and TL=Tunnel Length in centimeters

This study identified sorghum genotypes with resistance to *S. calamistis* based on leaf damage, dead heart formation, stem tunneling and exit holes following artificial infestation of seedling whorls with stem borer eggs. Multiple stem borer damage was considered because resistance to stem borers is a multi-mechanism quantitative trait, and thus, selecting for resistance based on a single parameter would not be effective (Singh *et al.*, 2011). Sorghum being an indigenous crop to Africa has co-evolved with native insect pests and it is likely that some varieties are tolerant or resistant to these insect pests (Muturi *et al.*, 2012). The analysis of variance revealed significant variation among the genotypes for all characters examined. Out of the 22 genotypes reported, 13(59%) of the genotypes with selection index from 0.0 to 0.50 showed resistance across the environment (Table 2). ICSB464, ICSL71086, SSV20041-2YELLOW, ICSL71018, ICSR94032, ICSV700, ICSL71193, ICSR94030, ICSL71253, ICSL71268, ICSL71023, ICSL71061, and ICSL 71137(in an increasing order). Sorghum resistance

to *C. partellus* based on reduced dead heart damage was reported by Sharma *et al.*, (2006). Genotypes: ICSB464, ICSL71086, ICSL71018, ICSR94032, ICSV700, ICSL71193, ICSR94030, ICSL71268, ICSL71023, ICSL71137, ICSL71061 and ICSL71253 was previously reported to be resistant to *Chilo partellus* and *Busseola fusca* in sorghum (Chinwada *et al.*, 2001; Sharma *et al.*, 2007, Muturi *et al.*, 2012) were also found to be resistant to *Sesamia calamistis* in this study. This implies that these genotypes have multiple resistance to the three species of stem borers, thus can be used as resistant check in screening trials and as potential donors in the breeding for multiple resistance to the multiple agents of stem borer infection.

Genotypes with selection index 0.60 to 0.10 were ICSV93046, ICSL71077, ICSL71244, ICSR94031, ICSL71258, DALWANDA, ICSL71168, KAURA MAIGUNDUMA, ICSL71180, ICSL71140, YAR GUMEL, ICSB472, ICSL71138, MACIA, ICSL71054, ICSB484, 89MW1005, MORI JIGAWA, SAMSORG6 (KSV12), IS36555, ICSL71219, ICSL71160, SAMSORG14, ICSL71185, ICSL71088, ICSL71247, ICSL71112 AMARYA DA ANGO, and ICSL71085 (data not shown). Out of the 29 moderately resistant genotypes, 6 were elite germplasm: DALWANDA, KAURA MAIGUNDUMA, YAR GUMEL, MACIA, MORI JIGAWA, AMARYA DA ANGO, 2 regional germplasm: 89MW1005, IS36555, and 2 IAR released varieties SAMSORG 6 and SAMSORG14.

Genotypes with selection index 1.1 to 1.5 were: SAMSORG41, KAURA KADUNA-1, ICSL71001, ICSL71055, IS30768, SAMSORG40, RIB*98-SB-F3-78, ICSL71187, ICSV745, SAMSORG5, ICSL71080, FARA 2 BAUCHI, IS17562, ICSL71016, ICSL71213, MAIMADARA-1, ICSL71246, ICSL71215, KL-2, SAMSORG38, ICSL71007, DANJIBE, RIBDAHU, KL-1, ZAUNA INUWA, TWIN SEEDED, SAMSORG39, SAMSORG17, and YAR WASHA (data not shown). Some genotypes previously reported as resistant from India (Sharma *et al.*, 2007) and Kenya (Muturi *et al.*, 2012) were found to be moderately susceptible to this pest at the test sites in Nigeria. This could be attributed to the insect species and genotype by environment interactions that influenced expression of resistance to damage by *S. calamistis*.

Genotypes with selection index 1.6 to 2.0 were SAMSORG44, SAMSORG8, SSV20041-1 WHITE, SAMSORG3, ICSL71005, MAIMADARA-2, SAMSORG 42, KAT487 and SAMSORG17. (Table 2). Genotype ICSL71005 previously reported resistant to *C. partellus* in India (Sharma *et al.*, 2007) and to *B. fusca* in Zimbabwe (Chinwada *et al.*, 2001), was found susceptible to *S. calamistis* in Nigeria. This could be attributed to the insect species and genotype by environment interactions.

Conclusion

This study demonstrated that there are genotypic differences in resistance/susceptibility to damage by *S. calamistis*. Resistance to *S. calamistis* is polygenic, thus, the use of numerous traits facilitate identification of superior genotypes. The sorghum materials could be grouped into resistant, moderately resistant, moderately susceptible and susceptible. This study identified sorghum genotypes with resistance to *S. calamistis*. The 13 Sorghum genotypes resistant to *S. calamistis* identified can serve as donor parents in the breeding for stem borer resistance. Their use as potential donor parents is further buttressed by the fact that 12 out of the 13 resistant parents possess multiple resistance to the causative agents of stem borer: - *Chilo partellus*, *Busseola fusca* and *Sesamia calamistis* in sorghum. In addition, eight elite germplasm and two IAR varieties were identified to be moderately resistant. Cultivation of genotypes with resistance to stem borers would greatly improve food security and income of the resource poor farmers in areas prone to African pink borer.

Recommendations

The genotypes identified to have multiple resistance can be used in breeding for multiple resistance to the multiple agents of infection. Genotypes that showed combined resistance to the three borers and with good agronomic performance may be deployed to areas where these borers exist. However, breeding for resistance to these borers should continue besides deployment of these stem borer resistant varieties. The most susceptible genotype, SAMSORG 3, SAMSORG 8, SAMSORG17, SAMSORG 42 and SAMSORG 44 could be utilized as a susceptible check in screening for resistance to *S. calamistis* and improved further to incorporate resistance. The 13 Sorghum genotypes resistant to *S. calamistis* identified can serve as potential donor parents in the breeding for stem borer resistance, also they can be used as resistant check in screening trials. The nine elite germplasm and three IAR varieties identified to be moderately resistant can be capitalized on as takeoff genotypes to build upon their resistance.

ACKNOWLEDGEMENTS

This study is part of a PhD research project of the first author conducted with financial support from the Harnessing Opportunities for Productivity Enhancement (HOPE2) Project under the Institute for Agricultural Research (IAR) Plant Science Department, A.B.U, Zaria. We are grateful to Dr C, Tom Hash from ICRISAT India and Dr Mary Mgonja ICRISAT -Nairobi for providing some of sorghum seeds for evaluation. Authors appreciate technical support provided by Mr. Hassan Tush and Ezekiel. Department of Plant Science, Ahmadu

Bello University Zaria. I also thank The Director General, NABDA, for granting me study leave.

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