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# Genetic variability, heritability and genetic advance of soybean (*Glycine max*) genotypes based on yield and yield-related traits

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## Abstract

The success of breeding programme relies on the variability present in the breeding material. Selection is also effective when there is significant amount of genetic variability among the individuals in a population. The study aimed at assessing genetic variability, heritability and genetic advance among yield and yield components of soybean genotypes. Eighty-two genotypes maintained at the Agricultural Research Council-Grain Crops were planted in an alpha lattice design, replicated twice in Potchefstroom and Brits in South Africa in 2017/18 growing season. Five plants per genotype were selected and yield and yield related traits were measured. Data were analysed through analysis of variance and genetic parameters were estimated. The results showed highly significant differences among the genotypes based on days to flowering, branch number per plant, hundred seed weight, pod weight per plant, pod length, seed number per plant, seed number per pod and grain yield. Heritability of yield and yield components were ranged from 5.9% to 100%. The most heritable traits were hundred seed weight (100 %), days to flowering (64.13 %) and seed number per pod (67.37%). The genotypic variance ranged from 0.3% to 33% while the phenotypic variation ranged from 1.5% to 44%. The phenotypic and genotypic coefficient of variations was high for grain yield, medium for days to flowering and lowest for hundred seed weight. The phenotypic coefficient of variations (PCV) was higher than genotypic coefficient of variation (GCV) for all characters indicating the influence of environmental factors. The expected genetic advance was high for most of the traits, medium for seed number per pod and hundred seed weight and low for pod number per plant. The expected genetic advance as percent of mean was variable and high for all traits. There was vast variation observed among the soybean genotypes evaluated based on the yield and yield related traits.

Keywords: Genetic variation, GCV, PCV, soybean, yield.

## Introduction

Soybean [Glycine max (L.) Merrill] belongs to genus Glycine. The genus Glycine wild is divided into two sub genera, Glycine and Soja. It is a diploidized allotetraploid (2n = 40) and autogamous plant. North Eastern (Manchuria) region of the China is believed to be the center of origin and diversification center of the cultivated soybean. Soybean is one of the most important oil and legume crop grown globally. It grows in the tropics and sub-tropics for human and animal consumption as well as industrial purposes. It contains 40 to 42% good quality protein and 18 to 22% oil, which comprise 85% unsaturated fatty acids. The oil from soybean is free from cholesterol, so it is highly desirable in the human diet (Aslam et al., 1995). Besides fixing the atmospheric nitrogen, this crop has the ability to grow in a wide range of environments, to reduce soil erosion, to suppress weeds and to suit inter and sequential cropping patterns.

According to FAO (2020) soybean is produced worldwide in an area of 124 921 956 hectares and the production recorded in 2018 was 348 712 311 tonnes. The highest producers were United States of America followed by Brazil and Argentina. Of the total world production, America produced 87.2%, Asia produced 9.3%, Europe was 2.7%, and 0.8% was produced in Africa. Africa's production was recorded to be 3 556 163 tonnes in area of 2 614 762 hectares in 2018. In South Africa, the production was estimated to be 1 540, 000 in the production area of 787 200 hectares. Soybean is produced in almost all provinces of South Africa.

It is a valued finding to understand genetic diversity and relationship for facilitating the transfer of useful genes among cultivated species and to maximize the use of available germplasm resources. The extent of genetic diversity in germplasm can be assessed through morphological characterization. The characterized material then helps the plant breeders to select the accessions to be utilized in hybridization programme (Ghafoor et al., 2002). An investigation of genetic relatedness at a broad level may provide important information about the historical relationship among different genotypes. It reveals genetic backgrounds and relationships of germplasm and also provides strategies to establish, unitize and manage crop core collections (Brown-Guedira et al., 2000). Therefore, the knowledge of the genetic variation within accessions from germplasm collections is essential to the choice of strategies to incorporate useful diversity in the program, to facilitate the introgression of genes of interest in commercial cultivars, to understand the evolutionary relations among accessions, to better sample germplasm diversity, and to increase conservation efficiency (Fu, 2003).

Morphological characters have long been used to identify species and genera, to evaluate systematic relationships, and to discriminate between varieties (Smykal et al., 2008). In breeding practice and seed production, the role of morphological descriptors is very important, since the distinguishing between varieties can be done quickly and precisely. A few genes, thus easily observable and suitable for cultivar differentiation and identification, usually control qualitative traits. On the other hand, quantitative traits have more limitations in cultivar description, since they are affected by environment, developmental stage of the plant and the generation of selfing of breeding material. The quantitative traits include yield and yield parameters of which are observable for determination of yield performance of various genotypes. Their performance is also dependable on the environment and may be specific to a certain agro-ecology. Hence, determining the genetic parameters of genotypes is of paramount importance for parental selection and recommendations to the farmers.

For the effective selection of superior genotype to use in hybridization programme for the development of superior varieties, proper study of genetic variability due to genetic and non-genetic causes and other genetic parameters is necessary (Prasad et al., 2012). There has been many studies on genetic variation conducted by various researchers in soybean. Ravindra et al. (2017) and Baraska et al (2014) studied genetic variation on soybean yield and yield related traits where the authors estimated heritability, phenotypic and genotypic coefficient of variations, determination of genetic advance and genetic advance as a percent of mean. Neelima et al (2018) determined genetic variability, heritability and genetic advance on one hundred and twenty four soybean germplasm accessions. Genetic parameters like the genotypic coefficient of variation (GCV), phenotypic coefficient of variation (PCV), heritability and genetic advance (GA) are useful biometric tools for measuring genetic variability (Aditya et al., 2011). Hence, characterizing the genetic background of soybean and determining breeding values should be done before carrying out any improvement programme (Agong et al., 2001). The Agricultural Research Council-Grain Crops (ARC-GC) maintains more than two thousand soybean germplasm accessions in the gene bank. Some of the material has not been characterised for identification of good agronomic traits and yield parameters. Hence, the objective of the study was to assess genetic variability, heritability and yield and yield-related traits among soybean genotypes maintained by the ARC-GC.

## Results

## Analysis of variance

Results of analysis of variance for the nine agronomic traits for the eighty-two soybean genotypes are shown in Table 3. The data showed highly significant differences for site, genotype and their interaction for days to flowering, seed number per plant, and seed yield. Highly significant differences were also observed for site on branch number, pod number per plant, pod weight, seed number per plant, and seed yield per plant. Genotype was also significant for days to flowering, branch number, pod length, pod weight, seed number per pod, seed yield and seed number per plant. Highly significant differences were observed for site x genotype for days to flowering and significant differences were recorded for seed number per plant and seed yield per plant.

## Variance components, coefficients of variation, heritability and genetic advance analyses

Table 4 shows phenotypic, genotypic and environmental variances, and coefficient of variation, broad sense heritability and response to selection of the eighty-two soybean genotypes. Phenotypic variation ranged from 0.01 to 847.6. The phenotypic variation was high for grain yield per plant followed by days to flowering and lower in hundred seed weight. The genotypic variation was high for grain yield and lower for pod number per plant. It ranged from 0.00 to 4750.00. The phenotypic variances were higher than the genotypic variances. The environmental variance ranged from 0.01 to 372.60 with grain yield being the highest followed by days to flowering and the lowest variance was observed in hundred seed weight. Phenotypic coefficient of variation ranged between 0.85 and 44.89. The genotypic coefficient of variation also ranged between 0.84 and 33.60. The phenotypic and genotypic coefficients of variations were high for grain yield, followed by days to flowering and the lowest was hundred seed weight. Generally, the phenotypic variances of all the traits were higher than the genotypic variances indicating an influence of environmental effects in the expression of the traits to a certain extent.

Broad sense heritability (H<sup>2</sup>) of the traits under study ranged from 5.88 to 100%. The heritability was high for hundred seed weight (100%), followed by seed number per pod (67.37%) and days to flowering (64.13%). The lowest heritability values were observed for pod number per plant (5.88%), pod weight (20.81%), seed number per plant (21.85%) and branch number per plant (31.36%). The genetic advance (GA) ranged from 2.63 to 3365.87 with grain yield being the highest, followed by days to flowering and the lowest value was recorded for pod number per plant. The percent of genetic advance (GAM) was highest for pod weight (31488.75), followed by grain yield (8002.22), and seed number per pod (2632.79) and lowest for pod number per plant (128.27). High genetic advance as a percent of mean coupled with high heritability were recorded for hundred seed weight, seed number per pod and days to flowering. High heritability values coupled with high genetic advance values was recorded for days to flowering.

## Clustering

A dendrogram was constructed and the genotypes were grouped into five clusters (Figure 1). The first two clusters had one genotype each, PR 154-22 and Santa Rosa, which shows that they are distantly related with other genotypes. The fourth clusters consisted of two genotypes which were also closely related to each other but distantly related with the others. The genotypes were F 82-7145 and F 82-7824. The genotypes in cluster two and five were subdivided into sub-clusters with the closely related being grouped together and the others between the sub-clusters being distantly related.

#### Discussion

The highly significant mean squares due to genotypes signified the presence of variability within the genetic materials under investigation. The significant differences due to sites indicates differences of the climatic conditions of the selected sites and the responses of the material grown. Significant interaction of genotypes by sites indicated the existence of a wide range of variations among genotypes and between sites selected and that different genotypes reacted differently to varying environments. This suggests that there is enough reasons for selection of traits from among the different sources of planting materials for yield and yield component traits. Similarly, Narayanankutty et al. (2005), and Jandong et al. (2019) recorded highly significant variation in most yield parameters among the soybean genotypes signifying the existence variability. According to plant breeders, it is important to obtain great genetic variability for selection in plants, imposition results that in fact promote significant genetic gains (Bernardo, 2010; Cruz et al., 2011).

Generally, the phenotypic variances of all the traits were higher than the genotypic variances indicating an influence of environmental effects in the expression of the traits to a certain extent. These results concur with the report by Neelima et al. (2018) and Ravidra et al. (2017). According to Deshmukh et al. (1986), PCV and GCV values greater than 20% are regarded as high, whereas values less than 10% are considered low and values between 10% and 20% considered medium. The phenotypic and genotypic coefficients of variations were high for grain yield, medium for days to flowering and lowest for hundred seed weight. This is attributed to the preponderance of additive gene action and possesses high selective value and thus, selection pressure could profitably be applied on this character for their improvement (Neelima et al 2018). High magnitude of GCV observed in grain yield indicates the presence of wide variation to be allowed for further improvement by selection (Jandong et al., 2020). These results are in agreement with the findings by Jandong et al. (2020), Neelima et al (2018), Lukman et al (2014), Mahbub et al. (2015), and Pushpa et al. (2013). The PCV was found to be higher than GCV, which indicates the important role of environment in the expression of the characters. The slight differences of PCV and GCV were seen in almost all the characters except the grain yield. This shows that the influence of environmental factors on the performance of phenotype is low (Aditya et al., 2011; Malek et al., 2014). Akram et al. (2016) stated that the low environmental factor causes a selection based on those characters would be effective. Other authors also reported small differences between the PCV and GCV (Mahbub et al., 2015; Nirmaladevi et al., 2015). The results of the phenotypic and genetic coefficient of variation show presence and wide genetic variation existing among the soybean genotypes maintained by the ARC-GC.

Heritability estimates are considered in understanding the pattern of inheritance of quantitative character like seed yield and yield components. Similarly, genetic advance is also a useful measure to predict gain in specified selection intensity. However, when it is considered along with heritability becomes more valuable to predict response to selection than the heritability estimates alone (Johnson et al., 1955). In this study, heritability was high for hundred seed weight, seed number per pod and days to flowering, which concurs to the reports from the study by Aditya et al. (2011), Aditya et al. (2011), Osekita and Olorunfemi (2014) and Kwusantoro et al. (2018). The high estimate of broad sense heritability revealed that the characters could be effectively used for cultivar development and hybridization program. Heritable variation of different yield parameters is useful for permanent genetic improvement (Singh, 2000). The most important function of heritability in the genetic study of quantitative characters is its predictive role to indicate the reliability of the phenotypic value as a guide to breeding value (Dabholkar, 1992; Falconer and Mackay, 1996). High heritability estimates for some of the phenotypic traits indicate less influence of environments, and so there is a good chance for the improvement of these traits through direct selection (Kumar et al., 2012).

Genetic advance measures the expected genetic progress that would result from selecting the best performance genotypes for a given characters (Allard, 1960). The genetic gain that can be obtained for a particular trait through selection is the product of its heritability, phenotypic standard deviation and selection differential (Burton, 1952). High heritability coupled with high genetic advance was observed for days to flowering, on the other hand, high heritability coupled with low genetic advance were observed for hundred seed weight and seed number per pod. High genetic advance as a percent of mean coupled with high heritability were recorded for hundred seed weight, seed number per pod and days to flowering. High heritability values coupled with high genetic advance values was recorded for days to flowering, which suggested that this trait can be considered as a favourable attribute for the improvement through selection and this may be due to additive gene action Panse (1957) and thus, could be improved upon by adapting selection without progeny testing. Additionally, high heritability values coupled with low genetic advances were recorded for hundred seed weight and seed number per pod revealing the predominance of non-additive gene action. High heritability coupled with high genetic advance as a percent of mean was recorded for hundred seed weight, seed number per pod and days to flowering. Similar results were reported by Bangar et al. (2003), Hina Kausar et al. (2005), Abady et al. (2013), Chandel et al. (2013), Amit et al. (2014), Mahbub et al. (2015), Yadawad et al. (2015) and Savita and Koti (2016). The estimate of heritability along with genetic advance gives the idea about the genetic architecture of the population.

Several researchers (Baraskar et al., 2014; Chandrawa et al., 2017; Malek et al., 2014; Mishra et al., 2015; and Reni and Rao, 2013) have also calculated genotypic and phenotypic components of variance, heritability and genetic advance for different yield characters in soybean. The authors revealed that selection was effective for a population with broad genetic variability and character with high heritability. Heritability provides useful information on the magnitude of the inheritance of traits from original parents to their progenies, while genetic advance is helpful in finding the actual genetic gain expected under selection (Larik et al., 2000, Nwangburuka and Denton, 2012, Ogunniyan and Olakojo, 2015).

The most diverse genotypes were PR 154-22, Santa Rosa, F 82-7145 and Vicoja among others. Clustering of genotypes aid in selecting the most diverse genetic material, potential parents for hybridization and for transgressive segregation and prospects of getting high hetersosis. Many authors have used clustering in soybean for assessment of presence of genetic diversity (Marconato et al., 2016; Ibiduni et al., 2020). The findings in this study concur with the reports given by Ojo et al., (2012) and Dayaman et al. (2009) who found six cluster groups among 45 genotypes of soya bean and Mehetre et al. (1994) who reported nine distinct clusters in 43 accessions of soya bean.

## Table 1. Agro-morphological characters measured in the study.

Trait	Abbreviation	Description
Days to 50% flowering	DFW	The number of days from date of sowing to the stage where 75% of the buds have fully flowered.
Branch number	BRN	Number of branches counted per plant
Hundred seed weight (g)	HSW	Weight of hundred seeds measured in grams
Pod Number per plant	PNP	Number of pods produced per plant
Pod weight (g)	PWT	Weight of pods produced per plant
Pod length (mm)	PDL	Length of the ten selected pods per plant
Seed number per pod	SNP	Number of seeds produced per pod
Seed number per plant	SNT	Number of seeds produced per plant
Yield per plant (g)	GYD	Total weight of seeds per plant.

## Table 2. List of germplasm used in the study.

Serial Number	Genotype	Serial Number	Genotype
1	12950	42	Hernon
2	14/6/32	43	HERON
3	165/5	44	IBIS
4	21/6/23/2	45	Impala
5	23545	46	Jackson
6	26090	47	Jupiter
7	28/6/54	48	Mack 6
8	40913	49	Maksura
9	42897	50	Mojiana
10	42912	51	MTD 63
11	44000	52	N69-2774
12	44007	53	ND 85
13	54 S 116	54	ORIBI
14	54 S 219	55	PI 170889/(R56-49)
15	54 S 95	56	PR 133 (484)
16	69 S 10	57	PR 144-4
17	69 S 13 Seleksie	58	PR 144-9
18	69 S 19	59	PR 145-2
19	85/05/	60	PR 154-13
20	89/05/	61	PR 154-14
21	AGS 239	62	PR 154-22
22	B 66 S 365	63	PR 154-47
23	B 66S 385	64	PR 164-20
24	Chippewa 63	65	PR 164-22
25	Coc Chum	66	PR 165-3
26	Columbia M 8 A	67	PR 165-31
27	Crawford	68	PR 165-50
28	D64-4636	69	PR 165-52
29	D66-8666	70	R 5-4-2 M
30	DB 1601	71	Rhosa ligte hilum
31	DUNDEE	72	S4-A,P,4
32	EGRET	73	Santa Rosa
33	Essex	74	Shelby
34	F 82-7145	75	Soja (pautena)
35	F 82-7656	76	SSS 2
36	F 82-7824	77	SSS 3
37	Grant	78	TN 81-46
38	Gx Gous	79	UFV-1
39	Hampton 266 A	80	Vicoja
40	Hawkeye	81	Yeluanda
41	Hawkeye (USSR)	82	Yubelejuaja



Figure 1. A dendrogram showing clustering of soybean genotypes based on yield and yield components.

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SOV	DF	DFL	HSW	BRN	PDL	PNP	PWT	SNT	SNP	SYD
Site	1	56195.97***	145.5ns	38.694***	10.80ns	175456***	81304.0***	1398926***	0.0000ns	38338.5***
Genotype	81	276.42***	248.7ns	3.710***	37.48**	5951ns	1283.9***	13898*	0.5079***	691.5**
Site*genotype	89	188.89***	186.3ns	1.919ns	24.83ns	5632ns	1144.5ns	15209**	0.0000ns	670.7**
Residual	143	66.55	234.2	2.474	22.73ns	6546.0	876.5	9799	0.1572	425.7

SOV = Source of variance, DF = degrees of freedom, ns = non-significant, \*\*\* = significant at P $\leq$ 0.001, \*\* = significant at P  $\leq$ 0.025 and \* = significant at P  $\leq$ 0.05.

Table 4. Phenotypic, genotypic and environmental variances and coefficient of variation, broad sense heritability and response to selection of the nine yield related traits of soybean genotypes.

Trait								
	δ₂p	δ₂g	δ₂e	PCV	GCV	H <sup>2</sup>	GA	GAM
HSW	0.01	0.01	0.01	0.85	0.84	100.00	19.41	1545.73
DFW	168.63	108.14	60.49	15.51	12.42	64.13	1717.93	2450.50
BRN	2.95	0.93	2.03	7.36	4.12	31.36	111.17	2037.82
PNP	0.05	0.00	0.04	1.51	0.37	5.88	2.63	128.27
PWT	0.06	0.01	0.04	1.81	0.82	20.81	550.84	31488.75
PDL	21.05	5.6	15.45	7.36	3.79	26.60	251.80	647.77
SNP	0.28	0.19	0.09	3.17	2.59	67.37	73.52	2632.79
SPT	0.06	0.01	0.04	1.56	0.73	21.85	10.63	466.48
YDP	847.60	475.00	372.60	44.89	33.60	56.04	3365.87	8002.22

## Materials and methods

## Plant material, trial layout and management

Eighty-two genotypes maintained by the Agricultural Research Council-Grain Crops are shown in Table 2. The genotypes were planted in 2017/18 growing season in Potchefstroom (26.7145° S. 27.0970° E) and Brits (25.6100° S. 27.7960° E) in the North West Province of South Africa. The trials were laid out in alpha lattice designs replicated two times. Each plot consisted of two 4 m length rows with a spacing of 75 cm between the rows and 10 cm between the plants. Fertilizer 2:3:4 was applied before planting. The plants were irrigated using sprinklers three times a week. The pre- and post-emergent herbicides (Metagan and Basagran) were used to control the weeds as well as manual weeding. Lime Ammonium Nitrate was top dressed 45 days after germination, before the plants flower. The cultural practices were applied as per soybean planting recommendations.

#### Data Collection

Five plants were evaluated from each replication of each genotype. The characters shown in Table 1 were measured.

## **Data Analysis**

Analysis of variance was performed for all traits in order to test the significance of variation among the genotypes using GenStat version 18. The replications and blocks were treated as fixed factors whereas genotypes, environment and their interactions were treated as random factors. The linear model used for data analysis was:

 $Yijkl = \mu + Gi + Ej + Rk(j) + Bl(jk) + GEij + Eijkl$ Where, Yijkl is the observed value of genotype I in block I and replication k of environment j,  $\mu$  is the grand mean, Gi is the effect of genotype i, Ej is the environment or location effect, Rk(j) is the effect of replication k in environment j, Bl(jk) is the effect of block I in environment j and replication k, GEij is the interaction effect of genotype I with environment j, and eijkl is the error (residual) effect of genotype I in block I and replication k of environment j. A dendrogram was constructed using RStudio in R statistical package.

The genetic parameters including genotypic and phenotypic variance components and coefficient of phenotypic and genotypic variability were estimated according to Burton (1951) and Singh and Chaudhary (1985). The formulae are as follows:

 $\delta 2g = MSe - MSg/r$ Where: MSg = mean square due to genotypes, MSe = error mean square r = the number of replication

Environmental variance  $(\delta 2e) = error mean square$ = MSe

Phenotypic variance  $(\delta^2 g) = \delta^2 g + \delta^2 e$ Phenotypic coefficient of variance (PCV)  $= \frac{ (\sqrt[6]{\sqrt{\delta 2p}} x \mathbf{100})}{xmean} x \mathbf{100}$ Genotypic coeffcient of variation (GCV)  $= \frac{\sqrt{\delta 2g}}{xmean} x \mathbf{100}$  Heritability (H<sup>2</sup>): heritability in broad sense for all characters was computed using the formula given by (Falconer et al., 1996).

Heritability (H2) = 
$$\frac{\delta 2g}{\delta 2p} x100$$

Where:  $H^2$  = heritability in broad sense  $\sigma^2 =$  Phenotypic variance  $\sigma^2 =$  Genotypic variance

Genetic advance in absolute unit (GA) and percent of the mean (GAM), assuming selection of superior 5% of the genotypes was estimated in accordance with the methods illustrated by Johnson et al. (1955) as:

#### $GA = K\delta PH2$

Where: K=the standardized selection differential at 5% selection intensity (k=2.06)

 $s_p$  =phenotypic standard deviation on mean basis  $H^2$ =heritability in broad sense

Genetic advance as percent of mean was calculated to compare the extent of predicted advance of different traits under selection, using the following formula:

$$GAM = \frac{GA}{X} \times 100$$

Where: GAM= genetic advance as percent mean GA= genetic advance under selection

= Mean of the population in which selection employed

### Conclusions

Analysis of variance revealed enormous genetic variability among the soybean genotypes studied. The phenotypic and genotypic coefficients of variations were high for grain yield, medium for days to flowering and lowest for hundred seed weight. This was attributable to the preponderance of additive gene action and possession of high selective value and thus, selection pressure could profitably be applied on these characters for further improvement. The phenotypic coefficient of variations (PCV) were higher than genotypic coefficient of variation (GCV) for all characters indicating the influence of environmental factors. High estimates of heritability coupled with high genetic advance and genetic advance as a percent of mean were recorded for days to flowering, hundred seed weight, and seed number per pod. The high estimate of broad sense heritability revealed that the characters could be effectively used for cultivar development and hybridization programmes. The expected genetic advance was high for most of the traits, medium for seed number per pod and hundred seed weight and low for pod number per plant. The expected genetic advance as percent of mean was high for all traits. There was vast variation observed among the genotypes.

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Where: xmean = grand mean of a character

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