Genetic Diversity, Heritability and Correlation of Quantitative Traits Sweet Corn (Zea Mays L. Var. Saccharata Sthurt) Ms-Unsika Inbred Lines

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Abstract. Sweet corn is an important commodity in Indonesia as the second main carbohydrate source after rice. Along with the growth in population, the demand for corn is increasing. Breeding programs to obtain high-yielding varieties are needed to compensate for the lack of higher demand for corn. Selection is the main activity in plant breeding programs that aims to increase the frequency of characters desired to be used as elders in crosses. Selection based on quantitative characters is difficult because many genes are controlled, but the success of selection can be improved by considering genetic parameters such as means, varians, heritability and genetic correlation. The aim of the study was to select sweet corn lines based on genetic parameters that are directly related to the yield. The study was conducted in July 2018 - December 2018 in the Experimental Station of PT. East West Seed Indonesia, Campaka Village, Purwakarta Regency. The genetic material used was 14 MS-Unsika sweet corn lines and 3 commercial sweet corn. The results showed that there was a high diversity of sweet corn lines tested. Characteristics of plant height, ear height, ear diamater, ear length, cob weight per plant, weight of 100 seeds, number of rows per ear had heritability and a high correlation with yield of tons per hectare. The MS 08 line is the best line as a candidate for hybrid parents.

Keywords: correlation, genetic variability, heritability, MS-Unsika lines, sweet corn

1 Introduction

Corn (Zea mays L.) is an important commodity in Indonesia, considering that this commodity has a multipurpose function, both for food, feed, and industrial raw materials [1]; [2]; [3]. One of the corn cultivars that plays a major role in the economy in Indonesia is sweet corn. Sweet corn is obtained from ordinary corn which has recessive mutations spontaneously, this mutation can control the conversion of sugar into starch in the endosperm. This sweet corn has a wrinkled and transparent seed character, with a high sugar content and low starch content in the endosperm [4]. Sweet corn can double sugar levels (around 12-14 %) and 8-10 times more soluble in water than normal young corn. Sweet corn contains a relatively high sugar content, sweetness is caused by three main genes, namely the sugary (su) gene, sugary enhancer (se), and shrunken (sh2). Sweet corn cultivars which contain the su gene, produce a high amount of

sugar, but changes sugar into starch quickly after harvest if the cob is not in a cold temperature [5]. Sweet corn cultivars containing se genes produce higher amounts of sugar than cultivars containing the su gene. Cultivars containing se genes will also convert sugar into starch such as normal sweet corn, but the process takes longer after harvest because the sugar content is higher. Sweet corn cultivars containing sh2 gene, do not directly convert sugar to starch and therefore after harvest the sweetness lasts for a very long time. At present the productivity of sweet corn is still relatively low so it needs to be improved.

Strategies for increasing sweet corn production can be carried out with plant breeding programs to obtain new varieties. The formation of new superior varieties, disease resistance and high production is the right strategy to solve the productivity problems of sweet corn. Crossing is one step to increase genetic variability and obtain a new genotype. The new genotypes that have been obtained are selected again to obtain better and superior genotypes. Corn breeders use morphological and agronomic markers in selection and crossing, although the expression of morphological and agronomic markers is sometimes influenced by environmental conditions and sometimes also influenced by epistatic and pleiotropic interactions [6], [7]. The success of plant breeding programs, especially sweet corn, is highly dependent on the diversity of germplasm. Germplasm diversity can be measured based on the phenomenon of performance of morphological, agronomic and molecular markers. Selection can be done if germplasm has a high level of genotype and phenotype diversity [8], [9], [10].

Analysis of genetic diversity between prospective hybrid parents, can be seen with molecular markers and agromorphological markers to obtain high heterosis, it is no longer necessary to cross all existing material, but only choose inbreds that have far genetic distances and have high yield potential. Quantitative genetic theory has explained the correlation between pairs between parents and genetic distance with estimates of heterosis [11], [12], [13]. Heterosis is a quadratic function of the difference between the frequency of alleles in the parent, namely genetic differences and also the dominant effects of alleles that control characters. Then, heritability is the relationship between genotype variants and variant phenotypes [14]. This relationship illustrates how far the visible phenotype is a reflection of the genotype. Basically, selection of segregated populations can be done through the value of phenotypic character quantities in the population. In selecting a plant character, attention is given to genetic diversity, heritability, and genetic progress. Selection will be more effective if there is extensive genetic diversity in the population. Heritability is very important in determining the selection method and in which generation the desired character should be selected. Genetic progress illustrates the extent of the effectiveness of the breeding process. Selection will be more effective if the value of high genetic progress is supported by the value of genetic diversity and high heritability. Thus the three genetic parameters greatly determine the success of the breeding program [15], [16].

The aim of the study was to obtain information on genetic diversity between sweet corn inbred lines based on quantitative traits and to find out genetic distances based on agromorphological markers of yield components to be tested further.

2 Material and Methods

2.1 Genetic Materials and Experimental Site

The fourteen inbred lines and three commercial hybrid of sweet corn were evaluated for yield performance and quantitative traits at the Experimental Station of PT. East West Seed Indonesia, Campaka Village, Purwakarta Regency, Indonesia and the experiments were laid out in a randomized complete block design in three replicates during July untill December, 2018.

2.2 Data Collecting

Quantitative traits parameters data were collected from yield component as follows: ear diameter without husk (EDWoH), ear diameter with husk (EDWH), ear length (EL), ear diameter (ED), number row per ear (NRE), number grain per row (NGR), number grain per ear (NGE), sugar content (SC), ear weight per plot (EWP), ear weight per plant with husk (EWPWH), and ear weight per plant witout husk (EWPWOH).

2.3 Data analysis

Analysis of variance (ANOVA) was carried out to establish the level of significance among genotypes using the STAR 2.0.1 Software from IRRI@2013. The mean values were compared using the least significant difference (LSD) procedure.

2.3.1 Analysis of Phenotypic Variance and Genotypic Variance

The variability of each quantitative trait was estimated by simple statistical measures including the mean, phenotypic and genotypic variances and coefficient of variation. The phenotypic and genotypic variation and coefficient of variation were calculated following the formula suggested by [14] and [6]. Phenotypic and genotypic variances were computed from the respective mean squares as suggested [14].

2.3.2 Genetic Distance

The level of genetic similarity (GS = genetic similarity) is estimated using the Jaccard coefficient [17]. Genetic similarity was analyzed based on *Unweighted Pair Group Method using Arithmetic Averages* (UPGMA) using NTSYS-pc Software version 2.1.0. Analysis of genetic distance matrices is obtained from the results of genetic similarity analysis [18], with a formula: S = 1 - GS, where S = genetic distance, GS = genetic similarity. The cophenetic correlation coefficient (r) is also calculated followed by the Mantel test [19] to see the goodness of fit from the results of cluster analysis.

2.4 Phenotypic Variance (δ2p)

The formula for the components of variance and heritability used are as follows ([14]) at Table 1:

Table 1. Analysis of variance agronomical and morphological traits of sweet corn genotypes

Source	DF	Mean Square	F Value	Pr(> F)
Rep (r)	r-1	MS_r		
Genotipe (g)	g-1	MS_g	$\sigma^2_e + r.\sigma^2_g$	
Galat (e) Total	(g-1) (r- 1) g.r-1	MSe	$\sigma^2 e$	

$$\sigma 2g = \frac{MSg - MSe}{r}$$

$$\sigma 2p = \sigma 2p + \sigma 2e$$

$$GCV = \left(\frac{\sqrt{Vg}}{x}\right). \ 100\%$$

$$PCV = \left(\frac{\sqrt{Vp}}{x}\right). \ 100\%$$

Broad-sense heritability (h2 bs) was calculated as suggested by [6]: $\sigma^2 a$

h2bs=
$$\frac{\sigma^2 g}{\sigma^2 p}$$

Where :

 $\sigma 2g = genotype variance$ mSg = mean square of genotypemSg = mean square of genotypemSg = mean square of errormSg = mean square of errormSg = mean square of errormSg = mean square of genotypemSg = mean square of errormSg = mean squaremSg = mean square

The broad-sense heritability value according to Stanfield (1983) is: 0.50 < h2 < 1.00: high; 0.20 < h2 < 0.50: medium; h2 < 0.20: low.

Phenotypic and genotypic correlations were estimated as:

$$\mathbf{r}_{G.(xy)} = \frac{Cov_{G(xy)}}{\sqrt{(\sigma_{Gx}^2)(\sigma_{Gy}^2)}}$$

3 Results and Discussion

The results of research on genetic diversity, heritability and the genetic distance of the sweet corn inbred lines of the Unsika collection based on quantitative characters can be shown below. The results of the analysis of variance shown in Table 1 show that all the characters observed were significant. The parameters observed were: ear diameter without husk (EDWoH), ear diameter with husk (EDWH), ear length (EL), ear diameter (ED), number row per ear (NRE),

number grain per row (NGR), number grain per ear (NGE), sugar content (SC), ear weight per plot (EWP), ear weight per plant with husk (EWPWH), and ear weight per plant witout husk (EWPWoH).

Agronomical Traits	DF	Mean Square	F Value	Pr(>F)	Remarks
EDWH	16	0.4154	7.23	0.000*	Significance
EDWoH	16	0.5604	15.00	0.000*	Significance
EL	16	21.1281	7.05	0.000*	Significance
ED	16	0.1288	3.23	0.0023*	Significance
NRE	16	11.863	2.77	0.007*	significance
NGR	16	241.9063	3.11	0.003*	significance
NGE	16	79507.7298	3.58	0.001*	Significance
SC	16	7.7889	5.08	0.000*	Significance
EWP	16	53.6749	15.61	0.000*	Significance
EWPWH	16	0.0218	16.86	0.000*	Significance
EWPWoH	16	0.0173	13.8	0.000*	significance

Table 2. Analysis of variance for agronomical traits of Sweet corn inbred lines MS- Unsika.

Notes: EDWH: ear diameter with husk; EDWoH: ear diameter without husk; EL: ear length; ED; ear diameter; NRE: number row per ear; NGR: number grain per row; NGE: number grain per ear; SC: sugar content; EWP: ear weight per plot; EWPWH: ear weight per plant with husk; EWPWoH: ear weight per plant without husk; *:=p < 0.06 :significance levels

Based on the results of observations and data analysis Table 3 shows that the estimation of genetic variability, the variation of phenotypes and heritability and the coefficient of variance in genotypes and the coefficient of phenotype variability in agronomic parameters shows that the coefficient of genotype diversity is low in ear length; ear diameter; row number per ear, ear weight per plant with husk and sugar content; while the coefficient of genotype diversity that is currently medium in characters ear diameter with husk; ear diameter without husk, number grain per row; number grain per ear; ear weight per plant without husk and ear weight per plot. The results of this analysis indicate that breeding will succeed if the germplasm possessed has high genetic diversity [6], [20].

The heritability value of all quantitative characters observed was high, except for the number row per ear indicating low heritability; then, the ear diameter character, number grain per row and number grain per ear show medium heritability. This shows that quantitative characters that have high heritability values can be used as selection parameters for the selection of new superior hybrid elders in the sweet corn tested such as ear diameter with husk; ear diameter without husk; ear length; sugar content; ear weight per plot; ear weight per plant with husk; and ear weight per plant without husk. This is in accordance with the results of research conducted by [21], [22], [23], [24]

Agronomical Traits	Mean	Mean — Square- Genotype	Varin	nces	CV	7 %	h2 bs (%)	Heritability Criteria
			σ2g	σ2p	GCV	PCV		
EDWH	1.1	0.4154	0.119333	0.176733	31.40423	38.21788	67.52169	High
EDWoH	1.62	0.5604	0.174333	0.211733	25.77361	28.40401	82.33627	High
EL	16.6	21.1281	6.044167	9.039767	14.81017	18.11217	66.86198	High
ED	1.31	0.1288	0.029667	0.069467	13.1481	20.11949	42.70633	Medium
NRE	13.12	11.863	2.525733	6.811533	12.11323	19.89247	37.08025	Low
NGR	26.18	241.9063	54.74623	132.4138	28.2623	43.95386	41.3448	Medium
NGE	364.07	79507.73	19096.45	41314.84	37.95696	55.83006	46.22176	Medium
SC	12.94	7.7889	2.0854	3.6181	11.1599	14.69962	57.63799	High
EWP	16.16	53.6749	16.74557	20.18377	25.32262	27.80096	82.96552	High
EWPWH	0.3312	0.0218	0.006833	0.008133	24.95893	27.22977	84.01639	High
EWPWoH	0.2182	0.0173	0.005333	0.006633	33.46914	37.32598	80.40201	High

Table 3. Means, estimates of genetic variance, phenotypic variance, broad sense heritability, genotypic coefficient of variation (GCV) and phenotypic coefficient of variation (PCV) for agronomical traits in sweet corn

Notes: EDWH: ear diameter with husk; EDWoH: ear diameter without husk; EL: ear length; ED; ear diameter; NRE: number row per ear; NGR: number grain per row; NGE: number grain per ear; SC: sugar content; EWP: ear weight per plot; EWPWH: ear weight per plant with husk; EWPWoH: ear weight per plant without husk; σ^2g : genetic variance; σ^2p : phenotypic variance; GCV: genotypic coefficient of variation; PCV: phenotypic coefficient of variation; h2 bs : broad-sense heritability.

In table 4 shows that the correlation coefficient between characters shows a high value between ear weight per plot with number grain per ear, ear length, ear diameter with husk; and ear diameter without husk; while ear diameter shows a high correlation between number row per ear; number grain per row; and number grain per ear. For sugar content there is no positive correlation with all other characters; ear weight per plant with husk.

Traits	EDWo H	EDW H	EL	ED	NR E	NG R	NG E	SC	EW P	EWP WH	EWPW oH
EDWo H	1	0.89*	0.63 *	0.46	0.48	0.53 *	0.57 *	0.1 4	0.95 *	0.95*	0.91*
EDWH		1	0.57 *	0.51 *	0.53	0.60 *	0.62 *	$\begin{array}{c} 0.0 \\ 7 \end{array}$	0.83 *	0.83*	0.92*
EL			1	0.67 *	0.70 *	0.58	0.64 *	0.3 0	0.60 *	0.60*	0.60*
ED				1	0.73 *	0.60 *	0.65 *	0.2	0.44	0.42	0.49
NRE					1	0.73 *	0.83 *	0.1 3	0.47	0.47	0.55*
NGR						1	0.98 *	0.1 3	0.48	0.49	0.57*
NGE							1	0.1 7	0.53 *	0.53*	0.61*
SC								1	0.16	0.15	0.08
EWP									1	1.00*	0.91*
EWPW H EWPW										1	0.91*
oH											1

Table 4. Correlation coefficients among agronomical traits in sweet corn

Notes: EDWH: ear diameter with husk; EDWoH: ear diameter without husk; EL: ear length; ED; ear diameter; NRE: number row per ear; NGR: number grain per row; NGE: number grain per ear; SC: sugar content; EWP: ear weight per plot; EWPWH: ear weight per plant with husk; EWPWoH: ear weight per plant without husk

Furthermore, based on the analysis of genetic diversity using NTSYS version 2.1.0 Software shows high genetic diversity with Euclidian distance 0.02-0.26 on quantitative characters (Figure.ure 1). In Figure.ure 1 the dendogram is divided into 2 clusters, namely cluster IA, IB and cluster II A, II B, subclass IIIA, III B. The distribution pattern among the 17 genotypes of the Sweet corn Unsika line based on qualitative characters is shown in Figure.ure 1. Spread pattern among 17 genotypes Sweet corn based on quantitative characters is shown in Figure.ure 1.

The graph is divided into 3 quadrants. Quadrant I consists of genotypes AB1 and AB2 with contributions of sugar content and ear weight per plot that influence the presence variation. Quadrant II contains genotypes AB2 and AB4, In quadrant III consists of genotype AB3 and AB5 with contributions character of ear diameter and ear number per row that affect variation. Figure.ure 2. Dendogram of clustering for genetic diversity based on agronomical traits in sweet corn inbred lines.

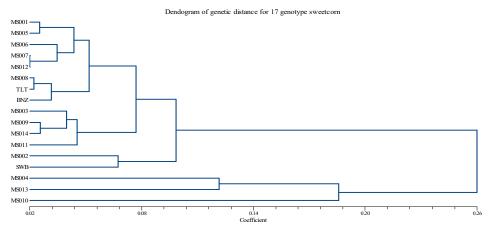


Figure. 1. Sweet corn Unsika line based on qualitative characters

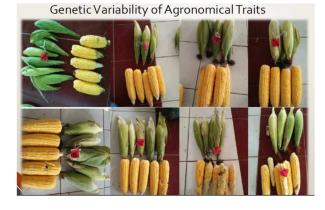


Figure. 2. Genetic diversity based on agronomical traits in sweet corn inbred lines

4 Conclusion

The results of this study concluded that the level of diversity of sweet corn tested based on quantitative characters was high, the genetic distance relationship between genotypes was also high so that it could be used for selection of prospective sweet corn parents; high heritability for all characters except sugar content; the best genotypes based on quantitative traits are MS 008 and MS 007; while the best hybrid is Bonanza F1.

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