

Determination of Some Physiological Parameters of Some Sweet Maize Inbred Lines

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Abstract: The study was conducted at Selcuk University Agriculture Faculty Crop Science Department in Konya/Turkey during 2013. At the Project during 2013 seeds obtained from 38 sweet maize inbred lines belong to S5 used that have been improved by S.U.Z.F. Crop science Department faculty members. In addition these materials Lümina and Merit hybrid maize varieties were used as standards. The study was conducted according to “Augmented Experimental Design” as four blocks at S.U.Z.F. Crop Science Department Prof. Dr. Abdulkadir AKCİN experimental area. Trials were settled for a year during 2013. When at the blocks all inbred lines were exemplify with an unrepeated line, standard hybrid maize varieties were repeated at all blocks. Stomatal conductance, chlorophyll content, photosynthetic efficiency, proline and MDA level values used to determine physiological parameters of lines. All analyses were done at 5 plants at all lines and standards during tasseling time. While Stomatal conductance values of sweet maize inbred lines varies between 31, 19 mmol.(m²s⁻¹)⁻¹ (4.1.5) to 4, 56 mmol.(m²s⁻¹)⁻¹ (4.26.2), chlorophyll content of lines changes between 57, 28 spad (4.5.2) to 34, 99 spad (4.16.2). Photosynthetic efficiency of lines ranges between 0,835 fv.fm-1 (4.16.2) to 0,732 fv.fm-1 (4.6.4). Proline level of lines changes 0, 70 µmol.g-1 (4.13.1) to 0,17 µmol.g-1 (4.26.2) and MDA levels ranges between 60,56 nmol.g-1 (4.1.3) to 29,34 nmol.g-1 (4.3.2). As a result of the study a lot of morphological and phenological characteristics of lines were determined comparatively with hybrid maize varieties, at this way an important variation is detected with all of those identified parameters. It can be told that those lines can be source of successful hybrid maize varieties in the future.

Keywords: Sweet Corn, Inbred Lines, Physiology, Malondialdehyde, Proline

1. Introduction

Maize is a plant that is very suitable for cultivating agricultural areas intensively, can use solar radiation very efficiently, can produce high level dry matter and grain and can be used multipurposes. Maize culture can be done all parts of our country because of its' wide adaptation capability and high yield potential [1]. Maize that provides benefits of country economy in all respects is a hot climate crop. Maize has an important place between all crop harvesting areas in our country with harvesting areas of 6 586 450 decare, with production of 5 950 000 tones and with yield of 907 kg da⁻¹. Maize cultivating areas of Turkey have high yield and production values [2].

Although maize known well produced wide areas in Turkey, sweet maize which is one of the important maize variety groups has no production and consumption statistical values [3]. Sweet maize (*Zea mays saccharata* Sturt.) that is one of the maize variety groups is cultivated especially USA intensively. When other maize variety groups are used animal feeding and industry with different aims, sweet maize can be used fresh, frozen and as canned food directly for human nutrition. Last years sweet maize consumption of Turkey increases day by day especially touristic areas and cosmopolite cities [4]. Some parts of products of maize, that' s fresh green parts and grains use for human and animal nutrition, use as raw material for producing of starch base sugar industry [5]. Local hybrid maize varieties are developed in our country every year and foreign base maize varieties also take part of planting areas in Turkey. To get own place of maize in Turkey' s planting areas, developed and developing maize varieties adaptation studies should be repeated and the properties of generations that can be transferred to other generations should be detected and recorded. One of the most important aim of maize breeding is to produce inbred parents that can form hybrid genotypes which can be used for trade. Measuring a crop's physiological parameters provides information for interpreting its response to the environment. Remote sensing quickly becoming recognized as a valuable tool that has the potential to assess a wide variety of physiological properties over a large area in a short amount of time [6]. Parameters that will be detected at this study will express values of lines according to physiology and will light the reason of the line becoming at the selection program at this way. To improve sturdy hybrid maize varieties, it is necessary to

improve strong lines. Aim of this study is determination of sturdy lines by identifying lines properties by using physiological parameters.

2. Materials And Methods

2.1. Materials

At the project seeds belong to 38 sweet maize inbred lines obtained from S5 stublines whose inbreeding programs have been going on, developed by Prof. Bayram SADE and Prof. Süleyman SOYLU from Selcuk University Agriculture Faculty Crop Science Department Konya/Turkey were used. In addition to these materials Lümina and Merit hybrid maize varieties used as standards.

2.2. Methods

The study was conducted at S.U.A.F. Crop Science Department Prof. Dr. Abdulkadir AKCIN experimental area as four blocks according to “Augmented Experimental Design”. The trials were settled down for a year during 2013. Also each line were represented unpeatedly as a row, standard hybrid maize varieties took part repeatedly at each block. The soil of experimental area was treated, weeds were removed mechanically and seed bed was prepared. Seeds obtained from inbred lines were sowed at the second week of May. Each inbred line was sowed by hand to 5 m rows. Between of rows was set as 70 cm, between of plants was set 25 cm. The experimental area was fertilized according to soil analyses results as 18 kg/da N and 11 kg/da P₂O₅. 4 kg of N and all of P [DAP (16 N–46 P₂O₅)] were treated with sowing. Rest of N was treated during second weed control and progressive developing stages at the form of ammonium nitrate (%33 N). Blocks were designed with 1 m intervals. Measurements and analysis were done at 5 plant of each row during “Tasseling Time”. The measurements and analysis that were done at inbred lines were summarized below.

3. Measurements And Analysis

3.1. Stomatal conductance

Stomatal conductance was measured with porometer during “Tasseling Time” at five plants of each rows and recorded [7].

3.2. Chlorophyll content

This parameter is measured with Spadmeter Spad-502 during “Tasseling Time” at five plants of each row and results recorded [8].

3.3. Photosynthetic efficiency

This parameter was also determined at five plants of each rows as others. Before measurement surface of treatment area of leaves were closed for 30 minutes. No changeable basal chlorophyll fluorescence (F_o), changeable fluorescence (F_v), maximum fluorescence induction (F_m) and changeable fluorescence/maximum fluorescence (f_v.f_m⁻¹) rate were determined with “Plant Efficiency Analyzer” (PEA) (Hansatech Instruments Ltd.) and results were recorded [7].

3.4. Proline

Free proline content determination was done according to [9]. 520 nm absorbance of toluene fraction that was aspired from liquid phase detected with spectrophotometer, proline concentration was calculated from calibration curve and recorded as fresh weight µmol. proline g⁻¹.

3.5. Lipid peroxidation

This parameter was detected according to the method which was defined by [10], according to this method MDA level that occurs final of TBAR reaction was calculated. Absorbance changes between 532 to 600 nm was measured to determine the activity and recorded.

3.6. Statistical analyses and assessment

Variances obtained from research were analyzed according to “Augmented Experimental Design” with Jump 5.0.1 program.

4. Results And Discussion

4.1. Stomatal conductance

Variance analysis values of control varieties were given Table 1. Merit one of the control varieties used at the study had the highest stomatal conductance value ($7,97 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$), the other one Lümina followed it with value of $\text{mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$.

TABLE I: Variance Analysis Table of Control Varieties Stomatal Conductance Values

Variation Source	Degree of Freedom	Sum of Square	Mean Square	F
Between Blocks	2	2,48	1,241	1,2985
Between Control Varieties	2	12,40	6,198	6,4876
Varieties	36	1046,22	29,062	30,4202
Error	3	2,87	0,955	-
General	43	1063,97	37,455	-

When lines compared with control varieties it is recorded that 4.1.5 ($31,19 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$), 4.2.3 ($21,19 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$) and 4.14.2 ($18,22 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$) coded lines left behind standards significantly. 4.6.3 ($4,69 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$), 4.6.4 ($4,67 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$) and 4.26.2 ($4,56 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$) coded lines were formed the lowest values group and lagged behind of standards. Values obtained from this parameter changed between $31,19 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$ and $4,56 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$. Stomatas are very sensitive cell structures that provide contact of plant with atmosphere directly are affected too much from environmental conditions. Being stomatal conductance values between normal limits show that plant is compatible with environmental factors and photosynthetic reactions are going on well [11]. When results obtained from this trial were investigated it is detected that stomatal conductance of some lines [4.1.5 ($31,19 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$); 4.2.3 ($21,19 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$); 4.14.2 ($18,22 \text{ mmol} \cdot (\text{m}^2\text{s}^{-1})^{-1}$)] are higher than standards. Based of these results it can be told that those lines adaptation capability to their cultivating areas is better than others. [7] reported that progressive stages of plant growth photosynthesis, transpiration rates and stomatal conductance are decreases significantly. Trial area was irrigated whenever need but Middle Anatolia is a geographical region that water resources are limited, humidity is not enough and this cause water lose by transpiration. So high stomatal conductance level of those lines means that those lines used current water efficiently. Those lines photosynthetic efficiency values became higher than much of other lines because they didn't have to be restrict their stomatal activity because of their efficiently water usage capability [4.1.5 ($0,818 \text{ fv} \cdot \text{fm}^{-1}$); 4.2.3 ($0,808 \text{ fv} \cdot \text{fm}^{-1}$); 4.14.2 ($0,818 \text{ fv} \cdot \text{fm}^{-1}$)]. Some lines profiting by current water more efficiently than others show that they can utilize water enough that is used as electron source at photosynthetic reactions. High stomatal conductance values of lines means plants have enough CO_2 at their structure that is used as C source at photosynthetic reactions. [12] found similar results also [13]. According to their study there is a positive correlation between grain yield, photosynthesis rate and stomatal conductance values. Stomatal conductance values are determined calculating of gas exchange level. High stomatal conductance values provides low leaf temperature, at this way it can be told that high stomatal conductance values provides low temperature of plant vegetation [14][12].

4.2. Chlorophyll content

Variance analysis values of control varieties were given Table 2 below. The control variety Lümina had the highest chlorophyll content value ($51,39 \text{ spad}$), Merit followed it with value of $50,63 \text{ spad}$.

TABLE II: Variance Analysis Table of Control Varieties Chlorophyll Content Values

Variation Source	Degree of Freedom	Sum of Square	Mean Square	F
Between Blocks	2	894,78	447,390	1,2263
Between Control Varieties	2	35,74	17,870	9,1775
Variety	36	930,53	25,848	1,5281
Error	3	35,74	11,915	-
General	43	1896,79	503,023	-

When lines compared with varieties it is observed that 4.1.5 (31,19 spad), 4.2.3 (21,19 spad) and 4.14.2 (18,22 spad) coded lines left standards [Lümina (6,79spad); Merit (7,97spad)] behind clearly. When lines compared with control varieties it is recorded that 4.5.2 (57,28 spad) and 4.3.1(51,58 spad) coded lines left behind standards significantly. 4.27.3 (36,13 spad), 4.27.1 (35,59 spad) and 4.16.2 (34,99 spad) coded lines were formed the lowest values group and lagged behind of standards. Values obtained from this parameter changed between 57,28 spad and 34,99 spad. Photosynthetic reactions are carried out at thylakoid membranes of granas at chloroplasts. Those structures include high level of chlorophyll. Chlorophylls are molecules that photosynthesis carries out, so meaning of having high level chloroplast is high level of photosynthetic activity. At this study some of lines chlorophyll content [4.5.2 (57,28spad); 4.3.1(51,58 spad)] are much more than standard varieties [Lümina (51,39 spad); Merit (50,63 spad)]. At this way it can be told that photosynthetic activity of those lines are higher than others. Chlorophyll fluorescence values of those lines became higher than some other lines also [4.5.2 (0,790 spad); 4.3.1 (0,791 spad)]. [15] reported that net photosynthetic rate values obtained chlorophyll meter have maximum relations with net photosynthetic rate. [12] reported that leaf chlorophyll content shows photosynthetic capacity. Using spadmeter is a cheap, fast and reliable method for detecting chlorophyll content and nitrogen level. Most suitable time using chlorophyll meter is the period after flowering.

4.3. Photosynthetic efficiency

Variance analysis table of control varieties were given at Table 5. When photosynthetic efficiency value of Lümina was $0,795 \text{ fv.fm}^{-1}$, photosynthetic efficiency value of the other control variety Merit became $0,801 \text{ fv.fm}^{-1}$.

TABLE III: Variance Analysis Table Belongs To Photosynthetic Efficiency Values of Control Varieties

Source of Variation	Degree of Freedom	Sum of Square	Mean Square	F
Between Blocks	2	0,00013	0,00007	0,1892
Between Control Varieties	2	0,00062	0,00031	0,8832
Variety	36	0,02300	0,00064	1,8095
Error	3	0,00106	0,00035	-
General	43	0,02481	0,00136	-

During comparing of lines with standards, it was observed that 4.16.2 ($0,835 \text{ fv.fm}^{-1}$), 4.15.4 ($0,828 \text{ fv.fm}^{-1}$) and 4.28.2 ($0,821 \text{ fv.fm}^{-1}$) coded lines lagged behind of Merit ($0,801 \text{ fv.fm}^{-1}$) that got the highest photosynthetic efficiency value of standards. Whereas 4.12.1 ($0,752 \text{ fv.fm}^{-1}$), 4.21.1 ($0,749 \text{ fv.fm}^{-1}$) and 4.6.4 ($0,732 \text{ fv.fm}^{-1}$) coded lines lagged behind of Lümina ($0,795 \text{ fv.fm}^{-1}$) whose photosynthetic efficiency value became the lowest of standards. Other values obtained from the parameter changed between them. [8] reported that photosynthetic efficiency is a parameter that is used to determine of plant high temperature tolerance under high temperature stress areas. At the study chlorophyll fluorescence values of some lines [4.16.2 ($0,835 \text{ fv.fm}^{-1}$); 4.15.4 ($0,828 \text{ fv.fm}^{-1}$); 4.14.1 ($0,821 \text{ fv.fm}^{-1}$)] became higher than standards [Lümina ($0,801 \text{ fv.fm}^{-1}$); Merit ($0,795 \text{ fv.fm}^{-1}$)]. Those lines chlorophyll [4.16.2 ($34,99 \text{ fv.fm}^{-1}$); 4.15.4 ($40,29 \text{ fv.fm}^{-1}$); 4.14.1 ($46,43 \text{ fv.fm}^{-1}$)] content also became higher than some other lines similarly. Values obtained from this parameter supported these values also. [16] reported that at the study they conducted Roman, determined some physiological characteristics like leaf occurrence, photosynthetic efficiency, stub occurrence and chemical composition at some hybrid maize groups of FAO 200-300, determined that hybrids whose those characteristics were better had yield more than $7,5 \text{ tones ha}^{-1}$.

4.4. Proline

Variation analysis table of control varieties were given table 4. The highest proline content from standards were obtained from Lümina with value of $0,34 \mu\text{mol.g}^{-1}$, Merit followed it with value of $0,27 \mu\text{mol.g}^{-1}$. While lines compared with standards proline content of 4.13.1($0,70 \mu\text{mol.g}^{-1}$), 4.25.1 ($0,69 \mu\text{mol.g}^{-1}$) and 4.4.3 ($0,65 \mu\text{mol.g}^{-1}$) coded lines became higher than standards, proline content of 4.24.2 ($0,23 \mu\text{mol.g}^{-1}$), 4.1.2 ($0,18 \mu\text{mol.g}^{-1}$) and 4.26.2 ($0,17 \mu\text{mol.g}^{-1}$) coded lines became lower than proline content of Merit ($0,27 \mu\text{mol.g}^{-1}$) whose proline content is the lower of standards. General proline content of values changed between $0,70 \mu\text{mol.g}^{-1}$

¹ and 0,17 $\mu\text{mol.g}^{-1}$. It was thought that proline was an osmolit that protects cell structures during osmotic stress conditions. But nowadays it is known that proline accumulation at plants is effective at different ways' to stress tolerance [17][18]. Proline is a molecular chaperon that protects protein and organizes different enzyme activities. Nowadays some of researchers reported that proline is an antioxidant which prevents ROS (Reactive Oxygen Species) and O_2^1 (Singlet Oxygen) activity [17][18]. Lines rich proline content means that those lines can be used to develop antioxidant rich varieties. Those varieties can be tolerant towards stress factors.

TABLE IV: Variation Analysis Table of Controls' Proline Values

Variation Source	Degree of Freedom	Sum of Square	Mean Square	F
Between Blocks	2	0,0010	0,0005	0,0000
Between Varieties	2	0,0270	0,0135	0,4183
Variety	36	0,8190	0,0228	0,6878
Error	3	0,0993	0,0331	-
General	43	0,9463	0,0699	-

4.5. Lipid peroxidation (MDA)

Variance analysis table of control group was given at Table 5. The highest MDA level was obtained from Merit (45,68 nmol.g^{-1}) and Lümina was followed it with value of 33,03 nmol.g^{-1} .

TABLE V: Variance Analysis Table of Control Varieties MDA (Malondialdehyde) Values

Variation Source	Degree of Freedom	Sum of Square	Mean Square	F
Between Blocks	2	0,0001	0,0001	0,0000
Between Control Varieties	2	336,8171	168,4086	55,7302
Variety	36	1623,3648	45,0935	14,9225
Error	3	9,0656	3,0219	-
General	43	1969,2476	216,5240	-

While lines were examined it was seen that 4.1.3 (60,56 nmol.g^{-1}), 4.3.1 (59,27 nmol.g^{-1}) and 4.27.3 (52,31 nmol.g^{-1}) coded lines got highest values opposite to 4.24.3 (31,66 nmol.g^{-1}), 4.18.1 (31,15 nmol.g^{-1}) and 4.3.2 (29,34 nmol.g^{-1}) coded lines that had lowest values. While comparing lines with standards, it was seen that 4.1.3 (60,56 nmol.g^{-1}), 4.3.1 (59,27 nmol.g^{-1}) and 4.27.3 (52,31 nmol.g^{-1}) coded lines left standards behind for this parameter. 4.24.3 (31,66 nmol.g^{-1}), 4.18.1 (31,15 nmol.g^{-1}) and 4.3.2 (29,34 nmol.g^{-1}) coded lines' MDA level generated lowest values. Other values changed between 60,56 nmol/g and 29,34 nmol.g^{-1} . Lipid peroxidation level that changes by time can be explained as low antioxidative defense system performance [19]. [20] reported that proline (an antioxidant) content increase according to drought at the study of developing drought tolerance bread wheat cultivars. At the same study it was also told that increased lipid peroxidation level effect antioxidative defense system negatively. When datas examined at this way, it can be told that lines whose MDA level is low can be more tolerated towards stress factors [4.1.3 (60,56 nmol.g^{-1}); 4.3.1 (59,27 nmol.g^{-1}); 4.27.3 (52,31 nmol.g^{-1})]. Drought stress whose effects increase by time can cause protein denaturation and lipid peroxidation and can trigger and accelerate cell corruption [21]. ROS damage cell components by inactivating enzymes. Increased ROS level is the most known indicator of stress. Plants and other organisms developed different mechanisms to cope with this problem. Free radicals cause lipid esterification by starting lipid peroxidation. Membrane damage and ethylene production are also free radicals effects. The capacity of antioxidative defense system is very important of preventing damage of oxidative stress and to detoxify ROS which belong to normal metabolism reactions. Reactive oxygen species are also produced at normal metabolism of alive cells [22] [23] [24].

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6. References

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