

NUTRITIVE VALUE AND FODDER POTENTIAL OF DIFFERENT SWEET SORGHUM GENOTYPES UNDER MEDITERRANEAN CONDITIONS

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ABSTRACT

This study was conducted to determine biomass yields and feed quality parameters of 21 different sweet sorghums (*Sorghum bicolor* var. *saccharatum* (L.) Mohlenbr.) genotypes grown as the second crop after wheat harvest under Cukurova conditions. Field experiments were conducted at the experimental fields of Dogankent-Adana location of Eastern Mediterranean Agricultural Research Institute in randomized blocks design with 4 replications in the years 2016 and 2017 (June-October). According to the results of the variance analyses of the data obtained from the experiment, years, genotypes and year x genotype interactions had significant effects on investigated parameters. Averaged over two years, dry matter yield (DMY), days to 50% flowering, crude protein content, NDF, ADF, and RFV of the genotypes varied between 21.6 and 62.9 t ha⁻¹, 55.0 and 99.1 days, 3.66 and 5.43%, 41.78 and 52.42%, 29.14 and 37.72%, and 111.3 and 148, respectively. It was determined that Ramada, Roma, Topper 76, UNL hybrid-3 and No91 genotypes were identified as late genotypes with DM yields of greater than 51 t ha⁻¹; Ramada, Roma and Topper 76 genotypes had the first places in terms of quality and they were superior than the other standard cultivars and genotypes.

Keywords: Fodder yield, forage quality, genotype, sweet sorghum

INTRODUCTION

World population is now almost 7.8 billion people. Annual average increase in world population is 1.05% and world population is expected to reach 10.9 billion people by the end of 21st century (Cilluffo and Ruiz, 2019). Together with increasing population, food and energy needs are expected to increase 2-3 folds (Fedoroff et al., 2010; Palmgren et al., 2015). As in the world, Turkey has also limited land resources for agriculture and available agricultural lands are continuously depleting because of non-agricultural use of agricultural lands, excessive soil tillage practices and resultant soil loss and erosion, excessive irrigations and resultant salinity problems. Besides, water deficits still posing serious threats on agricultural practices generate serious problems in various parts of the world (Godfray et al., 2010). Recent climate change trends have negative impacts on agriculture. Especially extreme temperatures and severe droughts result in significant yield and quality losses in agricultural products. To mitigate yield losses under adverse climate and soil conditions, appropriate growing techniques and soil management practices should be used and the crops with abiotic stress-tolerance should be grown. Sweet sorghum has quite low fertilizer demand, is quite tolerant to high temperature, drought and salinity, less selective in

soils, quite efficient in water use, thus it can be efficiently grown in arid and semi-arid regions. These characteristics make sorghum more advantageous over the other plant species under the same conditions (Mastrorilli et al., 1999; Gnansounou et al., 2005; Tesso et al., 2005). Although water demand of sorghum is less than the that of maize, sorghum has greater biomass potential than the other C4 crops (Gnansounou et al., 2005). Sweet sorghum is a short-day plant and has quite high temperature demand for optimum growth and development (Reise and Almodares, 2008). Sweet sorghum is a dedicated bioenergy crop, which has a higher biomass and forage yield potential than both forage sorghum and silage maize. Sweetness makes it highly palatable, thus it is highly preferred by livestock. Besides, biofuel generation from fodder crops is getting more popular in each day (Shanti et al., 2017). Insufficient supply of quality fodder is among the most important problems of Turkish livestock industry. To overcome this problem, production of summer grown C4 crops with a high biomass potential per unit area should be increased. Number of studies about potential use of sweet sorghum for forage production is highly limited in Turkey.

In this study, forage production potentials and some forage quality parameters of sweet sorghum genotypes

grown as the second crop after wheat harvest under Mediterranean ecological conditions were investigated.

MATERIALS AND METHODS

Experimental materials

The names and source of the genotypes tested in this study were as follows: **1)** Cowley, Dale, Grassi, M81-E, Mennonita, Nebraska sugarcane, PI579753, Ramada, Roma, Rox Orange, Smith, Sugar Drip, Theis, Topper 76, Tracy, UNL-Hybrid -3 ((26297xM81 E), Williams; **2)** no:2 USDA-China, no91 USDA-Taiwan, no5 USDA South Africa; **3)** Local check Gulseker. The mentioned plant material with the group numbers of 1, 2 and 3 were supplied by UNL (University of Nebraska, Lincoln, USA) and Western Mediterranean Agricultural Research Institute-Antalya/Turkey (supplied from ICRISAT and USDA gene bank) and University of Uludag, Bursa, Turkey, respectively.

Soil and climate characteristics of the experimental site

Experimental soils belong to Arikli soil series. Analyses on soil samples taken from 0-15 and 15-30 cm depths revealed that experimental soils were clay-loam (CL) in texture with pH values of between 7.0-7.50, total salt contents of between 0.22-0.27%, N contents of between 0.10-0.19%, organic carbon (OC) contents of between 0.63-0.90%, phosphorus (P) contents of between 0.63-0.90 mg kg⁻¹, lime (CaCO₃) contents of between 32.5-35.0%, sand contents of between 24-28%, silt contents of between 41-43% and clay contents of between 30-33% (Yucel et al., 2018).

Throughout the experimental period (June-October), average temperature was 25.1°C in 2016 and 24.8°C in 2017; relative humidity was 79.0% in 2016 and 79.6% in 2017; total precipitation was calculated as 46.2 kg m² in 2016 and 48.2 kg m² in 2017. Since the precipitations were not sufficient to meet water demands, irrigations were performed as needed.

Experimental design

Experiments were conducted on the experimental fields of Eastern Mediterranean Agricultural Research Institute in Dogankent (36° 51' 35" N and 35° 20' 43" E) in randomized blocks design with 4 replications during the years of 2016 and 2017. Sowing was performed in mid-June after wheat harvest. Before sowing, 50 kg ha⁻¹ pure nitrogen and phosphorus were applied to the experimental plots as basal fertilizers. Genotypes were sown manually in 4 rows of 5 m long at 70 cm row spacing and 15 cm on-row plant spacing. Dressing fertilizers were applied manually when the plants reached to heights of 40-50 cm as to have 50 kg ha⁻¹ pure nitrogen and irrigations were initiated then. Harvests were performed at the beginning of dough stage. Side rows and 0.5 m top and bottom of the rows were omitted as to consider side effects. Then, plot yields and yield per hectare were determined. Harvest time changed depending on the genotypes, and it was completed in October in the first year and harvest of late

genotypes was completed in the November in the second year because of prevailing precipitations.

Sample preparation and chemical analyses

During the harvest, 3 plants were randomly selected from each plot. Stalks were chopped into 15-20 cm long pieces and 1 kg fresh material was sampled. Samples were then dried in a drying chamber at 60°C for 7-10 days until a constant mass. Dried samples were weighed to determine dry matter contents and dry matter yields. Dried samples were ground to pass 1-2 mm sieves. Kjeldahl method was used to determine nitrogen (N) content of the samples.

Crude protein content was determined according to the following equation (AOAC, 1990); CP content = Nitrogen content (%) x 6.25. (AOAC, 1990). Cell wall components (Neutral Detergen Fiber (NDF) and Acid Detergen Fiber (ADF) contents, %) were determined with the aid of ANKOM fiber analyzer in accordance with the method specified by Van Soest et al. (1991). Digestible dry matter (DDM) content, dry matter intake (DMI) and relative feed value (RFV) were calculated with the equations provided by Schroeder (1994) as: DDM=88.9-(0.779x%ADF); DMI=120/%NDF; RFV=(%DDM X %DMI)/1.29.

Experimental data were analyzed using JMP statistical software in accordance with the randomized complete blocks design (RCBD). Significant means were compared with TUKEY's test at P≤0.05 significance level (Steel et al., 1997).

RESULTS AND DISCUSSION

Days to 50% Flowering (day): Variance analysis revealed that genotype and genotype x year interactions had significant effects on days to 50% flowering. Days to 50% flowering was determined as 75.2 and 74.1 days in the first and second years of the study, respectively. Significant differences were observed in days to 50% flowering with the years (Table 1). However, effects of the years on days to 50% flowering varied with the genotypes. Thusly, P1579753, Ramada and Gulseker genotypes reached to 50% flowering in significantly shorter time in the second year than in the first year, but Smith and Williams genotypes reached to 50% flowering in significantly longer time in the second year than in the first year. As the average of two years, days to 50% flowering of the genotypes varied between 55.0 - 99.1 days (Table 1). The Topper 76 genotype with 99.1 days reached to 50% flowering in significantly longer time than the other genotypes. Mennonita, Rox Orange and No2 genotypes reached to 50% flowering in significantly shorter time than the other genotypes. Shukla et al. (2017) classified sweet sorghum genotypes based on number of days to flowering as early (reaching to flowering in ≤70 days), medium (reaching to flowering in between 70-83 days) and late (reaching to flowering in ≥83days). According to this classification, 11 of investigated 21 genotypes were identified as early, 3 as medium and 7 as late genotypes. In other studies, days to 50% flowering of sweet sorghum genotypes were reported as between 54.9-

81.0 days (Mohammed and Mohamed, 2009; Mumtaz et al., 2019).

Table 1. Days to 50% flowering and green dry matter yield of sweet sorghum genotypes

Genotypes	Days to Flowering (Day)			Dry Matter Yield (t ha ⁻¹)		
	2016	2017	Mean	2016	2017	Mean
Cowley	62.3 mno ¹	66.0 j-n	64.1 g*	39.4 f-n ¹	41.2 e-m	40.3 d-h
Dale	74.0 fgh	72.0 f-i	73.0 d	33.4 h-n	35.3 h-n	34.3 f-i
Grassi	71.8 f-i	76.0 f	73.9 d	48.8 c-h	41.1 e-m	44.9 b-f
M81-E	98.5 abc	95.5 bc	97.0 ab	60.8 bc	38.3 g-n	49.6 b-e
Mennonita	56.5 pq	53.5 q	55.0 h	14.5 o	28.7 j-o	21.6 j
N. sugarcane	68.3 ijk	66.3 j-m	67.3 e-g	41.1 e-m	38.2 g-n	39.6 e-h
P1579753	68.8 ijk	61.0 n-p	64.9 fg	28.2 k-o	36.8 h-n	32.5 g-j
Ramada	93.8 cd	87.5 e	90.6 c	57.3 b-f	48.0 c-i	52.6 a-c
Roma	90.0 de	87.8 e	88.9 c	61.5 a-c	49.3 c-h	55.4 ab
Rox Orange	55.0 q	56.3 pq	55.6 h	23.7 m-o	35.0 h-n	29.3 h-j
Smith	62.8 l-o	75.0 fg	68.9 e	27.8 k-o	56.0 b-g	41.9 c-g
Sugar Drip	64.3 k-o	67.8 i-l	66.0 efg	26.9 l-o	31.2 h-o	29.0 h-j
Theis	99.5 ab	95.3 bc	97.4 ab	59.4 b-d	40.6 e-n	50.0 b-e
Topper 76	101.5 a	96.8 a-c	99.1 a	57.7 b-e	45.2 c-k	51.4 a-d
Tracy	67.8 i-l	66.0 j-n	66.9 e-g	31.2 h-o	39.9 e-n	35.6 f-i
UNL-Hyb-3	89.0 de	85.8 e	87.4 c	79.3 a	46.6 c-j	62.9 a
Williams	60.5 op	67.8 i-l	64.1 g	26.7 l-o	35.8 h-n	31.3 g-j
No2	56.5 pq	53.8 q	55.1 h	22.5 no	29.9 i-o	26.2 ij
No91	95.5 bc	96.0 bc	95.8 b	69.0 ab	41.9 d-l	55.5 ab
No5	67.3 i-m	69.0 h-k	68.1 ef	41.3 d-m	31.9 h-o	36.6 f-i
Gulseker	76.0 f	70.0 g-j	73.0 d	28.7 j-o	31.8 h-o	30.3 g-j
Mean	75.2	74.1		41.9 A⁺	39.2 B	
CV (%)		2.45			15.80	
F Genotype (G)		**			**	
F Year (Y)		ÖD			**	
F G x Y Int.		**			**	

*)The means indicated with the same letter in the same column are not significantly different according to the Tukey test at P≤0.05

+) The means indicated with the same capital letter in the same row are not significantly different at P≤0.05

¹) The means of different year-genotype combinations with the same lower case letters are not significantly different according to the Tukey test at P≤0.05

Dry Matter Yield (t ha⁻¹): As the averaged value over the genotypes, dry matter yield was determined as 41.9 t/ha in the first year and 39.2 t ha⁻¹ in the second year. The averaged dry matter yield of the first year was significantly greater than that of the second year. However, year x genotype interactions had significant effects on dry matter yields, thus effects of years on dry matter yield varied with the genotypes. Thusly, Smith genotype had significantly greater dry matter yield in the second year than in the first year. On the other hand, M81-E, Theis, UNL-Hyb-3 and No91 genotypes significantly lower dry matter yields in the second year than in the first year. Dry matter yields of the other genotypes did not vary significantly with the years. Since Smith genotype reached to 50% flowering in a longer time in the second year than the first year (Table 1), the genotype had longer time to produce dry matter, thus had greater dry matter yield in the second year. On the other hand, days to 50% flowering of M81-E, Theis, UNL-Hyb-3 and No 91 genotypes with significantly lower dry matter yields in the second year than the first year did not significantly vary with the years (Table 1). Low dry matter yields of the genotypes were mostly attributed to lower average temperatures of the second year than the first year throughout the growing

season. As the average of two years, dry matter (DM) yields of the genotypes varied between 21.6 - 62.9 t ha⁻¹ (Table 1) with the greatest dry matter yield from UNL-Hyb-3 genotype and the lowest dry matter yield from Mennonita genotype. The dry matter yields of Ramada, Roma, Topper 76 and No91 genotypes were not significantly different from that of UNL-Hyb-3 genotype. In previous studies conducted with different genotypes and under different ecologies, dry matter yields of sweet sorghum genotypes were reported as between 7.0 - 38.82 t ha⁻¹ (Turgut et al., 2005; Ayub et al., 2012; Cavalaris et al., 2017; Shanti et al., 2017; Vinutha et al., 2017). Except for early genotypes, present dry matter yields were greater than the values reported in earlier studies. Such greater yields were attributed to advantageous soil and climate conditions of the present experimental site.

Crude Protein Content (%): Year, genotype and year x genotype interactions had significant effects on crude protein content of sweet sorghum (Table 2). As the averaged value over genotypes, significantly lower crude protein content (CPC) was observed in the first year of the experiments than the second year. Precipitations in October and lower average temperatures in the second year than in the first year retarded plant aging, thus mean

CPC of the second year was significantly higher than the first year. Thusly, Kruse et al. (2008), Akar et al. (2014) and Ileri et al. (2018) also reported lower CPCs for maize plants in hotter years. Since year x genotype interactions had significant effects on CPCs, effects of the years on CPCs varied with the genotypes. Accordingly, Dale, M81-E, Smith, Sugar Drip, Topper, Williams, No91 and No5 genotypes had significantly greater CPCs in the second year than in the first year, but years did not generate significant differences in CPCs of the other genotypes (Table 2). As the average of two years, crude protein contents of the genotypes varied between 2.59 - 6.63% and differences were found to be significant (Table 2). Rox Orange genotype (5.43%) had significantly greater CPCs than Cowley, M81-E and Theis genotypes. On the other hand, Theis genotype (3.66%) had significantly lower CPC than the other genotypes, except for Cowley, Dale, M81-E, Ramada, Topper, Tracy, UNL-Hyb-3 Williams and No5 genotypes. Mohammed and

Mohammed (2009) reported crude protein contents of sweet sorghum genotypes as between 6.6 - 11.0%, Ayub et al. (2012) as between 6.62 - 8.29%, Mahmood et al. (2013) as between 5.6 - 10.0%, Neto et al. (2017) as between 6.11 - 11.71% and Shanti et al. (2017) as between 9.28 - 10.05%. Aguiar et al. (2006) reported crude protein contents of sorghum above-ground biomass as between 4.2 - 13.3%. CPCs determined in this study were lower than the values reported in previous studies. Such differences were mainly attributed to differences in genotypes, growing conditions, cultural practices, greater plant heights and yields of the present study. Leaf/stalk content decreases with increasing plant heights, stalks contain less crude protein than the leaves, thus greater plant height negatively influence CPCs. Accordingly, Dagtekin (2019) reported highly significant negative correlations between plant height and crude protein content of pearl millet (*Pennisetum glaucum*).

Table 2. Crude protein content and neutral detergent fiber of sweet sorghum genotypes

Genotypes	Crude Protein Content (%)			Neutral Detergent Fiber (%)		
	2016	2017	Mean	2016	2017	Mean
Cowley	3.54 hk ¹	4.68 c-j	4.11 cd*	57.14 a-f ¹	41.11 ij	49.12 ab*
Dale	3.41 i-k	5.72 a-g	4.56 a-d	43.68 h-j	40.75 ij	42.22 b
Grassi	4.35 d-k	5.73 a-g	5.05 a-c	56.92 a-g	44.24 f-j	50.58 a
M81-E	2.59 k	5.78 a-g	4.18 b-d	57.59 a-d	46.91 d-j	52.25 a
Menonita	4.74 b-1	5.29 a-h	5.02 a-c	57.24 a-e	35.80 j	46.52 ab
N. sugarcane	4.70 c-j	5.47 a-g	5.08 a-c	47.77 c-j	42.36 h-j	45.06 ab
P1579753	4.48 d-j	5.31 a-h	4.89 a-c	60.46 ab	40.30 ij	50.38 a
Ramada	4.25 e-k	5.28 a-h	4.77 a-d	48.56 b-j	44.25 e-j	46.41 ab
Roma	4.29 d-k	6.07 a-e	5.18 a-c	52.40 a-1	41.00 ij	46.70 ab
Rox Orange	5.02 a-1	5.85 a-f	5.43 a	52.79 a-1	43.04 h-j	47.91 ab
Smith	3.99 g-k	6.63 a	5.31 ab	48.72 b-j	44.09 g-j	46.41 ab
Sugar Drip	3.54 h-k	6.41 a-c	4.98 a-c	54.97 a-h	42.07 h-j	48.52 ab
Theis	2.89 j-k	4.27 d-j	3.66 d	49.26 b-1	47.66 d-j	48.46 ab
Topper 76	3.52 h-k	5.69 a-g	4.60 a-d	45.02 d-j	47.07 d-j	46.04 ab
Tracy	3.54 h-k	5.02 a-1	4.28 a-d	41.62 ij	41.94 ij	41.78 b
UNL-Hyb-3	4.04 f-k	4.79 b-1	4.41 a-d	52.13 a-j	41.54 ij	46.84 ab
Williams	3.53 h-k	6.05 a-e	4.79 a-d	49.24 b-j	41.61 ij	45.42 ab
No2	4.47 d-j	5.76 a-g	5.12 a-c	46.30 d-j	46.51 d-j	46.40 ab
No91	3.24 i-k	6.53 ab	4.89 a-c	60.68 a-c	43.72 h-j	52.20 a
No5	3.31 i-k	5.76 a-g	4.53 a-d	50.24 a-1	44.41 e-j	47.33 ab
Gulseker	4.43 d-j	6.11 a-d	5.27 a-c	62.47 a	42.37 h-j	52.42 a
Mean	3.90 B	5.64 A⁺		52.15 A⁺	42.99 B	
CV (%)		13.49			9.62	
F Genotype (G)		**			**	
F Year (Y)		**			**	
F G x Y Int.		**			**	

*The means indicated with the same letter in the same column are not significantly different according to the Tukey test at P≤0.05

) The means indicated with the same capital letter in the same row are not significantly different at P≤0.05

¹) The means of different year-genotype combinations with the same lower case letters are not significantly different according to the Tukey test at P≤0.05

Neutral Detergent Fiber (%): Dry matter NDF contents of the genotypes were significantly influenced by the years, genotypes and year x genotype interactions. The averaged value of NDF content over the genotypes in the first year (52.20%) was significantly higher than that in

the second year (42.99) (Table 2). Again, precipitations in October of the second year and resultant cool temperatures retarded plant aging, thus reduced NDF content of dry matter as compared to the first year. Thusly, Lee et al. (2017) reported that grass species grown

in cool regions had 21% less NDF content than in hot regions. Since the year x genotype interactions had significant effects on NDF contents, effects of years on NDF contents varied with the genotypes. Accordingly, Gulseker, No91, P1579753, M 81-E, Mennonita, and Cowley, genotypes had significantly greater NDF contents in the first year than in the second year while years did not generate significant differences in NDF contents of the other genotypes. As the average of two years, NDF contents of the genotypes varied between 41.78 - 52.42% and differences were found to be significant. Gulseker, M81-E, No91, Grassi and P1579753 genotypes had significantly greater NDF contents than Dale and Tracy genotypes. In previous studies, NDF contents of sweet sorghum genotypes were reported as between 32.6 - 62.5% (Mohammed and Mohamed, 2009; Machado et al., 2012; Neto et al., 2017; Vinutha et al., 2017). Our findings were in the range of values given in the mentioned reports.

Acid Detergent Fiber (%): The ADF content of sweet sorghum dry matter was significantly influenced by the years, genotypes and year x genotype interactions. As it was in NDF contents, significantly higher averaged value of ADF content was determined in the first year

than that in the second year (Table 3). NDF is cell membrane component and also includes ADF (Schroeder, 1994; Lema et al., 2000). Therefore, ADF content of dry matter increases with increasing NDF content. The effect of years on NDF content varied with the genotypes. Thusly, Cowley, M81-E, Mennonita, P1579753, no91 and Gulseker genotypes had significantly greater ADF contents in the first year than in the second year, but years did not generate significant differences in ADF content of the other genotypes (Table 3). As the average of two years, ADF contents of the genotypes varied between 29.14 - 37.72% and differences were found to be significant. Gulseker genotype (37.72%) had significantly greater ADF content than Dale, Mennonita, N sugarcane, Ramada, Smith, Topper 76, Tracy, Williams, No2 and No5 genotypes. M81-E genotype had significantly greater ADF content than Tracy and Williams genotypes. In previous studies, ADF contents of sweet sorghum genotypes were reported as between 26.0 - 54.2% (Lema et al., 2000; Madibela et al., 2002; Mohammed and Mohamed, 2009; Mahmood et al., 2013; Vinutha et al., 2017). Present ADF contents were again in range of values given in the mentioned earlier reports.

Table 3. Acid detergent fiber and relative feed value of sweet sorghum genotypes

Genotypes	Acid Detergent Fiber (%)			Relative Feed Value		
	2016	2017	Mean	2016	2017	Mean
Cowley	38.97 a-e ¹	25.96 ij	32.46 a-c*	96.50 g-l ¹	157.2 ab	126.9 a-c*
Dale	30.10 c-j	29.59 e-j	29.84 bc	141.6 b-f	150.4 a-d	146.0 ab
Grassi	36.63 a-g	30.41 c-j	33.52 a-c	103.5 e-l	137.4 b-g	120.4 bc
M81-E	39.57 a-c	31.35 c-j	35.46 ab	94.4 i-l	128.3 b-j	111.3 c
Mennonita	39.15 a-e	22.78 j	30.96 bc	95.1 h-l	188.1 a	141.6 ab
N. sugarcane	31.42 c-j	31.07 c-j	31.25 bc	130.0 b-j	143.1 b-e	136.5 a-c
P1579753	41.02 ab	26.84 h-j	33.96 a-c	88.2 kl	157.9 ab	123.1 a-c
Ramada	30.35 c-j	30.34 c-j	30.34 bc	125.5 b-k	137.7 b-g	131.6 a-c
Roma	35.48 a-i	28.22 f-j	31.85 a-c	109.6 d-l	151.9 a-c	130.8 a-c
Rox Orange	35.70 a-i	29.10 f-j	32.35 a-c	107.9 c-l	144.7 b-e	126.3 a-c
Smith	34.55 a-i	28.52 f-j	31.54 bc	118.5 b-l	141.4 b-f	130.0 a-c
Sugar Drip	37.47 a-f	29.36 f-j	33.41 a-c	101.2 f-l	146.2 b-d	123.7 a-c
Theis	35.24 a-i	32.30 b-j	33.77 a-c	116.3 b-l	124.5 b-k	120.4 bc
Topper 76	29.91 d-j	29.09 f-j	29.50 bc	136.3 b-h	131.2 b-j	133.8 a-c
Tracy	29.90 e-j	28.38 f-j	29.14 c	146.9 a-d	149.3 a-d	148.1 a
UNL-Hyb-3	36.05 a-h	31.67 c-j	33.86 a-c	108.8 d-l	144.6 b-e	126.7a-c
Williams	31.11 c-j	27.26 g-j	29.18 c	124.7 b-k	151.5 a-c	138.1 a-c
No2	30.58 c-j	29.21 f-j	29.90 bc	132.2 b-h	134.4 b-i	133.3 a-c
No91	39.50 a-d	30.96 c-j	35.23 a-c	90.2 j-l	138.0 b-g	114.1 c
No5	33.14 b-i	27.38 g-i	30.26 bc	118.1 b-l	142.6 b-f	130.3 a-c
Gulseker	43.94 a	31.51 c-j	37.72 a	81.5 l	141.6 b-f	111.6 c
Mean	35.23 A	29.11 B⁺		112.7 B	144.9 A⁺	
CV (%)		10.50			11.42	
F Genotype (G)		**			**	
F Year (Y)		**			**	
F G x Y Int.		**			**	

*)The means indicated with the same letter in the same column are not significantly different according to the Tukey test at P≤0.05

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¹) The means of different year-genotype combinations with the same lower case letters are not significantly different according to the Tukey test at P≤0.05

Relative Feed Value: Relative feed value (RFV) of the sweet sorghum genotypes were significantly influenced by the years, genotypes and year x genotype interactions. Averaged value of RFV (144.9) in the second year was significantly higher than that in the first year (112.7) (Table 5). RFV is closely related to DDM and DMI (Linn and Martin, 1999). Effects of the years on RFV of genotypes varied with the genotypes (Table 3). Accordingly, Cowley, Mennonita, P1579753, Roma, Sugar Drip, No91 and Gulseker genotypes had significantly greater RFVs in the second year than in the first year, but RFVs of the other genotypes did not vary significantly with the years (Table 3). As the average of two years, RFVs of the sorghum genotypes varied between 111.3 -148.1 and differences were found to be significant. Tracy genotype had significantly greater RFV than Grassi, M81-E, Theis, No91 and Gulseker genotypes, but the other genotypes did not significantly different from each other in RFV. Also, Dale and Mennonita genotypes had greater RFV than M81-E, No91 and Gulseker genotypes. RFV is used as an indicator of feed quality and assumed to be 100 for alfalfa hay at flowering stage. (Moore and Undersander, 2002; Hackmann et al., 2008). RFVs determined in this study were all greater than 100. Atis et al. (2012) reported RFVs of sorghum genotypes harvested at different development stages of plants as between 85.2 - 129.2.

CONCLUSION

In this study, field experiments were conducted with 21 different sweet sorghum genotypes grown as the second crop after wheat harvest under Eastern Mediterranean (Adana) ecological conditions for two years. Dry matter (DM) yields varied between 21.6 - 62.9 t/ha. Ramada, Roma, Topper 76, UNL hybrid-3 and No91 genotypes were identified as late genotypes with DM yields of greater than 51 t/ha; Ramada, Roma and Topper 76 genotypes had the first places in terms of quality and they were superior than the other standard cultivars and genotypes. Sweet sorghum yields were 4-5 times greater than yields of silage maize grown as second crop under Cukurova conditions (Korkmaz et al., 2005; Yucel et al., 2005) and feed quality values were equivalent or greater than silage maize. From the results of this study, it was concluded that sweet sorghum with greater tolerance to drought and high temperatures, less fertilizer needs and selectivity in soils than maize could reliably be used as an alternative of silage maize to meet roughage deficits of livestock industry.

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