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Description and Evaluation of Some Newly Introduced Grape Cultivars Under Egyptian Conditions

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ABSTRACT



This research was conducted for two consecutive seasons (2017 and 2018) in a private vineyard located at El-Sadat city, Menofia governorate to describe and evaluate three grape seedless cultivars namely: Early Sweet, Prime and Star light grape cultivars under Egyptian conditions. The results revealed that Early Sweet grape cultivar was the earliest cultivars concerning the phenological dates include bud burst, full bloom, fruit set, veraison and physiological ripening followed by Prime grape cultivar, whereas Star light grape cultivar was the latest in this respect through the two seasons of the study. All the studied cultivars succeeded under Egyptian conditions and characterized by high quality of bunch and berries. Genetic diversity among the three cultivars was assessed using SCoT and ISSR molecular marker techniques. Cluster analysis showed molecular and phenotypic variation among cultivars, which divided into three groups, each group was contained one cultivar. These cultivars were distinguished by 36 molecular markers 27 out of them were from SCoT markers which proved successful to target generic regions across the grape genome. These markers after deciphering their structure may help to evolve reliable molecular markers to select desirable traits in these cultivars. So, we recommend expansion in the cultivation of these cultivars under Egyptian conditions and benefit of our molecular results to elicit reliable molecular markers characterizing desirable traits in these cultivars to use in Egyptian grape cultivars improving programs.

Keywords: Grape cultivars, Phenotypic description, Genetic diversity, SCoT, ISSR, Egyptian conditions

INTRODUCTION

Grape is the one of most important fruit crops in the world and it ranked in Egypt after Citrus. According to the latest Ministry of Agriculture statistics (2018), Egypt's total grape area reached 202655 feddans, with a production of 1892993 tons. Forty years ago, most vineyards were occupied by two major cultivars namely Thompson Seedless and Roumi Ahmer in addition to a small area cultivated with some native cultivars. After the 1981 season, some new table grape cultivars were introduced, which were cultivated in various cultivation areas in the desert and delta regions; various morphological characteristics and bunch performance were recognized in these cultivars.

Ampelography is a well-established scientific method for characterizing grapevine genotypes based on the description of various morphological, phonological, and pomological characteristics. Many scientists have standardized and extended this approach in order to provide a more rational and reliable identification of Vitis material (Alleweldt and Dettweiler, 1986; Dettweiler, 1991; Soylemezoglu *et al.*, 2001 and Santiago *et al.*, 2007).

However, the obtained results often including the environmental effects too (Kumar, 1999). Therefore, in recent years, molecular markers have been used as premium tools for genetic diversity identification. They are neutral, feasible, do not depend on age and tissue type, and also are not influenced by environmental conditions. (Zietkiewicz *et al.*, 1994). Among these molecular markers, Inter Simple Sequence Repeats (ISSRs) and Start Codon Targeted polymorphism (SCoT) are used efficiently for genetic diversity assessment of plants (Etminan *et al.*, 2016). Many studies found that SCoT is might be more effective than other dominant DNA molecular markers like RAPD and ISSR because it is gene-targeted (Gupta *et al.*, 2018). Also, SCoT is superior over these markers in higher polymorphism and better marker resolvability (Gorji *et al.*, 2011). Moreover, SCoT can generate co-dominant markers caused by insertions and deletions, as well as they can generate dominant markers caused by sequence variations like RAPD and ISSR (Aswathy *et al.*, 2017). However, using ISSR and SCoT markers together gives very effective, reliable, and more superior results in genetic diversity study than the use of single markers (Mao *et al.*, 2018).

Previous studies were focused on describing and evaluating grape cultivars (Brooks and Olmo 1972; Watt, 1983; Walker and Boursiquote, 1992; Abd El-Kawi and El-Yam, 1992 a, b and c; Abd El-Fattah and Kastor, 1993 a and b; Morrison, 1994; Tourky *et al.*, 1995; El-Sharkawy 1995; Fawzy 1998; Marwad, 2002 a and b; Gaser, 2006; Girgis 2007; Sabry *et al.*, 2009; Abd El-Rahman, 2016; Mohamed and Tarbia, 2017 and El-Morsy *et al.*, 2017). Recently, some studies have used SCoT and ISSR markers successfully together successfully in the molecular diversity assessment of grape varieties, such as Abdel-Hameed *et al.* (2020) and Bashandy *et al.* (2020). Whereas they explain that both

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methods were effectively efficient for studying the genetic diversity in grape until between closely related cultivars.

Therefore, the main objective of this investigation is to describe and evaluate three grape cultivars namely: Early Sweet, Prime and Star light grape cultivars were studied under Egyptian conditions, with stress on some special morphological characteristics and molecular diversity assessment, which may serve in distinguishing these cultivars.

MATERIALS AND METHODS

This research was conducted for two consecutive seasons (2017 and 2018) in a private vineyard located at El-Sadat city, Menofia governorate to description and evaluation three grape seedless cultivars namely: Early Sweet, Prime and Star light grape cultivars under Egyptian conditions. The selected vines were five years old, grown in a sandy loam soil, irrigated by the drip system and spaced at 2 X 3 meters apart. The vines were cane pruned during the fourth week of December with maintain a load of 72 buds/vine (10 canes x 6buds) plus (6spurs x 2buds) and trellised by the Spanish Parron system. Four replicates for each cultivar were taken where each replicate consisted of six vines.

Botanical characteristics:

The following characteristics were studied:

1) Phenological dates

Phenological dates include bud burst, full bloom, fruit set, veraison and physiological ripening; the last date was harvested when the TSS reaches 16-17% in accordance with Tourky *et al.* (1995).

2) Descriptive measurements

The morphological studies were carried out according to the International Amelographic Registered Schedule (Dalmasso and Cosmo, 1952 and Cosmo *et al.*, 1958).

The following assessments were classified in accordance with many authors (Bioletti, 1938; Singh & Singh, 1940; Kolenati, 1946; Kolenati 1946; Breider, 1950; Rodrigues, 1959 and Watt 1983) and focused on the growing tip (hairs and colour), the tendril sequence and tip shape), the leaf (size, shape, surface, colour, thickness, pubescence, number of lobes, sinuses, margin and petiole), the bunch (weight, length, shape, density and peduncle) and the berry (weight, size, shape and colour).

3) Bunch physical characteristics

Average bunch weight (g), bunch dimensions (cm) and shot berries (%) were determined.

4) Berry physical characteristics

Average berry weight (g), berry size (cm3), berry dimensions (cm) and berry shape index were determined.

5) Berry chemical characteristics

Total soluble solids in berry juice (T.S.S.) (%) by hand refractometer, total titratable acidity as tartaric acid (%) (A.O.A.C. 1985) and TSS/acid ratio were calculated.

Experimental design and statistical analysis

The completely randomized design was adopted for the experiment. The statistical analysis of the present data was carried out according to Snedecor and Cochran (1980). Averages were compared using the new L.S.D. values at 5% level (Steel and Torrie, 1980).

Phenotypic distance

Based on quantitative data of studied characteristics, Phenotypic distances PD were calculated and agglomerative hierarchical clustering (AHC) dendrogram was drawn by Euclidian method using XLSTAT.7 software statistical approach according to Abd El-Aziz *et al.* (2019).

Molecular diversity assessment: -

Genomic DNA was isolated from juvenile grape leaves using a DNeasy plant mini kit (bio basic). The purity and concentration were measured by UV spectrophotometer. The PCR reaction was carried out using isolated genomic DNA from each cultivar as a template for PCR amplification using seven ISSR primers and nine SCoT primers (Table 1) according to Abd El-Aziz *et al.* (2019). PCR products for each primer were loaded on a 1.3 % agarose gel mixed with ethidium bromide and electrophoresed against a DNA ladder (0.1 to 3.0 kbp). The run was performed at 100 V for about 30 min in BioRad mini-submarine gel.

DNA banding patterns were photographed using a UV light on the Bio-1D Gel Documentation system. All photos were analyzed by GelAnalyzer 3 software. Also, cluster analysis was carried out agglomerative hierarchical clustering (AHC) according to Abd El-Aziz et al. (2019) using XLSTAT.7 software. Polymorphic Information Content (PIC) and DI (Diversity Index) were calculated according (Gorji *et al.*, 2011). Also, from binary data, the Resolving power (Rp) values were calculated as described in Prevost & Wilkinson (1999). Molecular distances MD were calculated by Dice coefficient (Nei and Li, 1979).

To verify the nature of the relationships between molecular distances ($MD_{ISSR, SCoT and All}$) and Phenotypic distances (PD), simple correlation coefficients were estimated using the computational software Minitab 17 (El-Zanaty et al., 2013).

 Table 1. List of all ISSR and SCoT Primers used in the study

SCoT Pri	mers	ISSR-Primers				
Name	Sequence $(5' \rightarrow 3)$	Name	Sequence $(5' \rightarrow 3)$			
SCoT-1	CAACA <u>ATG</u> GCTACCACCA	14A	(CT)8TG			
SCoT-2	CAACAATGGCTACCACCC	44B	(CT)8GC			
SCoT-3	CAACAATGGCTACCACCG	HB-09	(GT)6GC			
SCoT-4	CAACAATGGCTACCACCT	HB-10	(GA) ₆ CC			
SCoT-6	CAACAATGGCTACCACGC	HB-11	(GT)6CC			
SCoT-8	CAACAATGGCTACCACGT	HB-12	(CAC) ₃ GC			
SCoT-10	CAACAATGGCTACCAGCC	HB-15	(GTG) ₃ GC			
SCoT-11	AAGCA <u>ATG</u> GCTACCACCA					
SCoT-12	ACGACATGGCTACCAACG					

The underlines of ATG codon in the primer sequence were fixed.

RESULTS AND DISCUSSION

Botanical description and evaluation:

1) Phenological dates

Early Sweet grape cultivar was the earliest cultivars with regard to the phenological dates represented in bud burst, full bloom, fruit set and grape maturity followed by Prime grape cultivar, whereas Star light grape cultivar was the latest in this respect through the two seasons Table (2).

These findings are consistent with those recorded by Abd El-Fattah and Kasstor (1993a) on Beauty Seedless and

Black Monukka grape cultivars; Abd El-Fattah and Kasstor (1993b) on Black Rose and Ribier grape cultivars; Marwad (2002a) on Black Rose and Ribier grape cultivars; Marwad (2002b) on Beauty Seedless and Black Monukka grape cultivars; Abd El-Rahman, (2016) on Princess and Autumn Royal grape cultivars and Mohamed and Tarbia (2017) on Sable, Midnight Beauty and Desert Red grape cultivars.

Table 2. Dates of bud burst, full bloom, fruit set and maturity of Early Sweet, Prime and Star light grape cultivars in 2017 & 2018 seasons

Carltfanger	50% bud	burst date	70% full t	oloom date	Fruit s	set date	Maturity date		
Culuvar	2017	2018	2017	2018	2017	2018	2017	2018	
Early Sweet	11-Mar	14-Mar	6-Apr	8-Apr	21-Apr	24-Apr	31-May	3-Jun	
Prime	14-Mar	16-Mar	9-Apr	12-Apr	23-Apr	27-Apr	6-Jun	11-Jun	
Star light	19-Mar	22-Mar	17-Apr	19-Apr	29-Apr	1-May	15-Jun	19-Jun	

2) Descriptive measurements

The data related to the evaluation and morphological description of the studied items are shown in Table (3) and illustrated in Figure (1).

• Growing tip:

- Hairs and colour:

All studied grape cultivars were cob-webby hairs with green colour with the exception of the Star light grape cultivar, which was green with purple.

• Tendrils:

- Sequence and tip shape

The sequence of tendrils was intermittent with di-trifid shape in all studied grape cultivars.

• Leaf:

- Leaf size and shape:

All studied grape cultivars were large leaf area (more than 125 cm^2) with Orbicular shape.

- Leaf surface and colour:

All studied grape cultivars were smooth leaf surface with green colour.

- Leaf thickness and pubescence:

All studied grape cultivars were medium leaf thickness with Cob-webby.

- Leaf lobes:

Number of leaf lobes was five in all studied grape cultivars.

- Leaf sinuses:

Regarding depth of leaf sinuses, it was shallow in depth, when folding the lobe, the sinus reached less than one third of the way to petiole in all studied grape cultivars. As for the form of sinuses, it was narrow in all studied grape cultivars.

- Leaf margin:

With respect to the types of margin, it was dentate in all studied grape cultivars. Concerning teeth size, it was medium *i.e.* breadth was equal to length in all studied grape cultivars. As for the apical tooth, it was pointed in Early Sweet grape cultivar, whereas Prime and Star light grape cultivars were convex. Regarding number of teeth, it noticed that Early Sweet, Prime and Star light grape cultivars were medium (58, 54 and 66) respectively.

Table 3. Description and evaluation of Earl	v Sweet, Prime and Star light grape cultivars
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		Early Sweet	Prime	Star light
Growing tin	Hairs	Cob-webby	Cob-webby	Cob-webby
Glowing up	Colour	Green	Green	Green with purple
Tandrila	Sequence	Intermittent	Intermittent	Intermittent
Tenums	Tip shape	Di-tri-fid	Di-tri-fid	Di-tri-fid
	Size	Large(179.4cm ²) cm2)	Large (187.1 cm ²)	Large(174.9cm ²) cm2)
	Shape	Orbicular	Orbicular	Orbicular
	Surface	Smooth	Smooth	Smooth
	Colour	Green	Green	Green
	Thickness	Medium	Medium	Medium
	Pubescence	Cob-webby	Cob-webby	Cob-webby
	Number of lobes	5	5	5
	Leaf sinuses			
	Depth	Shallow	Shallow	Shallow
	Form	Narrow	Narrow	Narrow
Leaf	Leaf margin			
	Туре	Dentate	Dentate	Dentate
	Teeth size	Medium	Medium	Medium
	Apical tooth	Pointed	Convex	Convex
	Number of teeth	Medium (58)	Medium (54)	Medium (66)
	Petiole			
	Shape	U -shaped	U-shaped	U -shaped
	Sinus	Wide	Wide	Wide
	Petiole length (P)	6.6	9.3	8.4
	Leaf length (L)	8.1	9.8	11.1
	Petiole P/L	0.81 (Long)	0.95 (Long)	0.76 (Medium)
	Weight	Medium (440.8g) (440.8g)	Big (557.5g)	Big (601.9g)
	Length	Medium (16.5)	Long (19.7)	Medium (17.0)
Bunch	Shape	Shouldered	Winged	Conical
	Density	Well-filled	Well-filled	Compact
	Peduncle	Medium (3.1cm)	Medium (3.4cm)	Medium (2.6cm)
	Weight	Big (4.87g)	Big (4.90g)	Big (4.95g)
Dorm	Size	Large (4.82cm ³)	Large (4.87 cm ³)	Large (4.91cm ³)
Delly	Shape	Ovoid	Ovoid	Oblate
	Colour	Yellowish green	Yellowish green	Bright red



Early Sweet





Star Light Figure 1. Leaf, bunch and berry of some grape cultivars

- Petiole:

With regard to petiole shape, it was U-shaped in all studied grape cultivars. As for petiole sinus, all studied grape cultivars had wide. Concerning the ratio between petiole length to leaf length P/L, Early Sweet and Prime grape cultivars were long, while it was medium in Star light grape cultivar.

• Bunch:

With regard to bunch weight, it is clear that Early Sweet grape cultivar was medium (251-500g), whereas Prime and Star light grape cultivars were big (501-1000g).

As for bunch length, it was found that in Prime grape cultivar was long (18-24cm), while Early Sweet and Star light grape cultivars were medium (12-18cm).

Concerning the bunch shape, it was noticed that Early Sweet grape cultivar was shouldered, Prime grape cultivar was winged and Star light grape cultivar was conical.

As for bunch density, it was found that Early Sweet and Prime grape cultivars were well-filled, while Star light grape cultivar was compact.

With respect to peduncle of bunches, it was noticed that all studied grape cultivars were medium (2.5-3.5 cm). • Berries:

As for berry weight, all studied grape cultivars were big (3.3-7.0g).

Concerning berry size, all studied grape cultivars were large $(3.3-7.0 \text{ cm}^3)$.

With regard to berry shape, it was noticed that Early Sweet and Prime grape cultivars were ovoid, while Star light grape cultivar was oblate.

With respect to berry colour, it is clear that Early Sweet and Prime grape cultivars were yellowish green, while Star light grape cultivar was bright red.

The findings obtained are in accordance with those of many investigators working on different cultivars (Ismail, 1989, Tourky *et al.*, 1995; Fawzy, 1998; Marwad 2002 a and b; Mohamed and Tarbia, 2017 and El-Morsy *et al.*, 2017).

3) Bunch physical characteristics

As shown in Table (4), data concerning bunch physical characteristics of Early Sweet, Prime and Star light grape cultivars in both seasons was record.

With respect to bunch weight, it was noticed that Star light grape cultivar had significantly the highest values followed by Prime, while Early Sweet grape cultivar resulted in the least values in both seasons.

As for bunch dimensions, it was found that Prime grape cultivar significantly attained the highest values of bunch length and width, whereas Early Sweet and Star light grape cultivars had the lowest values of these ones, which insignificant differences between them in both seasons.

With respect to shot berries percentage, Early Sweet grape cultivar had significantly the highest percentage followed by Prime grape cultivar, whereas Star light grape cultivar did not contain shot berries in both seasons.

Table 4. Bunch physical characteristics of Early Sweet, Prime and Star light grape cultivars in 2017 & 2018 seasons

Cultivars	Bunch weight (g)	Bunch length (cm)	Bunch width (cm)	Shot berries (%)
First season				
Early Sweet	437.9	16.23	19.41	22.61
Prime	553.6	19.57	21.17	13.17
Star light	596.5	16.74	19.54	0
New LSD (5%)	31.9	1.43	1.17	7.54
Second season				
Early Sweet	443.7	16.81	19.76	21.43
Prime	561.3	19.92	21.29	14.68
Star light	607.2	17.36	19.83	0
New LSD (5%)	37.4	1.51	1.24	5.92

4) Berry physical characteristics

Data presented in Table (5) revealed that Early Sweet, Prime and Star light grape cultivars were differing among them concerning berry physical characteristics in both seasons.

As for average berry weight and size, no significant differences were observed among Early Sweet, Prime and Star light grape cultivars of these ones in both seasons.

With regard to berry length, Early Sweet grape cultivar had significantly the highest values, followed by Star light grape cultivar with no significant differences were noticed between them, whereas Prime grape cultivar resulted in the least values in both seasons of the study.

As for average berry diameter, Star light grape cultivar attained significantly the highest values followed by Prime, grape cultivar, while Early Sweet grape cultivar resulted in the least values in both seasons of the study. Regarding berry shape index; Early Sweet grape cultivar had significantly the highest values (more elongation) followed by Prime grape cultivar, whereas Star light grape cultivar resulted in the least values in both seasons.

Table 5. Berry physical characteristics of Early Sweet, Prime and Star light grape cultivars in 2017 & 2018 seasons

2010 5	casons				
Cultivars	Berry weight (g)	Berry Size (cm ³)	Berry length (cm)	Berry diameter (cm)	Berry Shape index
First season					
Early Sweet	4.84	4.79	2.41	1.91	1.26
Prime	4.88	4.85	2.35	1.94	1.21
Star light	4.93	4.89	2.39	2.05	1.17
New LSD (5%)	N.S.	N.S.	0.03	0.04	0.03
Second season					
Early Sweet	4.89	4.85	2.43	1.93	1.26
Prime	4.91	4.88	2.36	1.95	1.21
Star light	4.97	4.93	2.42	2.08	1.16
New LSD (5%)	N.S.	N.S.	0.02	0.03	0.02

5) Berry chemical characteristics

As shown in Table (6), data concerning berry chemical characteristics of Early Sweet, Prime and Star light grape cultivars in both seasons was record.

With respect to total soluble solids in berry juice, insignificant differences was observed among Early Sweet, Prime and Star light grape cultivars of this parameter in both seasons.

Regarding acidity in berry juice, it was noticed that Star light grape cultivar had significantly the least percentages followed in an ascending order by Prime grape cultivar, whereas Early Sweet grape cultivar resulted in the highest percentages in both seasons. Concerning berry TSS/acid ratio, the highest significant values of this parameter were attained by Star light grape cultivar followed by Prime grape cultivar, while Early Sweet grape cultivar had the least values in both seasons.

The findings obtained are in accordance with those of many investigators working on different cultivars (El-Sharkawy 1995; Fawzy 1998; Marwad, 2002 a and b; Gaser, 2006; Girgis, 2007; Sabry *et al.*, 2009; Mohamed and Tarbia, 2017 and El-Morsy *et al.*, 2017).

Table 6. Berry chemical characteristics of Early Sweet, Prime and Star light grape cultivars in 2017 & 2018 seasons

2010 scas	0115		
Cultivars	TSS (%)	Acidity	TSS/acid ratio
First season	(70)	(70)	Tudo
Early Sweet	16.7	0.49	34.1
Prime	16.5	0.48	34.4
Star light	16.4	0.46	35.7
New LSD (5%)	N.S.	0.02	0.7
Second season			
Early Sweet	16.9	0.48	35.2
Prime	16.6	0.45	36.9
Star light	16.5	0.44	37.5
New LSD (5%)	N.S.	0.01	0.4

Molecular assessment

To assess the genetic diversity among Early Sweet, Prime and Star light grape cultivars, banding patterns and DNA profile (Figures. 2: 4) of seven ISSR and nine SCoT primers were screened to investigate the genetic diversity. Except for HB-12, the other primers revealed polymorphic patterns and illustrated them to be valid with an acceptable degree in discriminating among these cultivars.



Figure 2. ISSR banding patterns of Early Sweet, Prime and Star light grape cultivars, for nine primers. L, ladder (0.1: 3 kbp) and lanes 2 to 4 represent the three cultivars.



Figure 3. SCoT banding patterns of Early Sweet, Prime and Star light grape cultivars, for nine primers. L, ladder (0.1: 3 kbp) and lanes 2 to 4 represent the three cultivars.

			19	SR						SCoT							Ta									
Primers	14A	44B	HB-9	HB-10	HB-11	HB-12	HB-15	s	CoT 1		SCoT 2		so	CoT 3	so	CoT 4		SCoT 6	SCoT 8	SCoT	10	SCoT	11 SCoT 12	ISSR	SCoT	Comb
MS bp	996 652 485 309 259 185	597 284 417 417 484 557 557 362	335 517 347 298 245	738 460 383 311	214 214 262	434 348 310 268 228	952 598 422 417	887 774 544	451 364 321 244 1973	1498 1201 956	836 697 569 527 443	403 332 269 1472	1076 815	767 481 411 367 367 310 1699 1466	991 811 720 657	574 505 413 374 341	317 259 724 578	512 493 461 405	383 359 531 531 468 426 372 353	905 794 697 603 555	507 507 439 398 310	536 481 444	427 382 640 545 545 472 367 367 314	35	75	108
Star light		Ш					II I			I		I			H			11		Ш		Ш		26	49	75
Prime	III <mark>III</mark>							11	IIII	I		П	П	Ш	I	1111		I II			I	П		29	50	79
Early Sweet	I			I II				I			11111			Ш	11			П		Ш	П	III		23	50	73

Figure 4. DNA-profile representation of ISSR & SCoT fingerprints of Early Sweet, Prime and Star light grape cultivars based on 108 amplicons, 36 of them were positive marker loci.

From ISSR banding patterns, the molecular data were estimated (Table 7) detected that the 14A & 44B primers which contain the repetitive motif (CT) showed the highest number of polymorphic amplicons 5 & 4 with 25.7 % from all ISSR amplicons. Also, these primers targeted generating 6 unique amplicons with 66.67% from all ISSR unique amplicons. Indicating that this microsatellite is more

contribution in generating reliable markers in grape. Moreover, these primers showed the highest Rp values were 3.33 and 2.67 with polymorphism % and PIC of (83.33, 0.370) and (57.14, 0.254), respectively. Suggesting the high discriminatory potential of these primers compared to other ISSR primers used in this study according to Prevost & Wilkinson (1999).

Table 7. Molecular data of ISSR molecular marker techn	ique.
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		Amp	licons		Polymorphic				
Primer	Malaanlan		Polymo	rphic		Polymorphism	index	Resolving	
Name	size range	Monomorphic	Without unique	Unique	Total	%	content (PIC)	power Rp	
14A	185:996	1	2	3	6	83.33	0.370	3.333	
44B	335:597	3	1	3	7	57.14	0.254	2.665	
HB-09	245:517	2	2	0	4	50.00	0.222	1.332	
HB-10	245:738	2	2	1	5	60.00	0.267	1.998	
HB-11	262:329	2	1	0	3	33.33	0.148	0.666	
HB-12	228:434	5	0	0	5	0.00	0.000	0.000	
HB-15	309:952	2	1	2	5	60.00	0.267	1.998	

On the other hand, Table 8 illustrated molecular data of SCoT primers used in this study. All these primers were from the Dataset-I type which targets highly expressed genes in plant tissues as described by Sawant *et al.* (1999). The first seven SCoT primers targeted generating of a total of 38 polymorphic amplicons. These primers were similar in the last five nucleotides at the 5'end and different in the last three nucleotides at the 3'end. These primers targeted generating of a total of 25 from 27 unique amplicons with 92.6 %. Where, the SCoT-4, SCoT-10 and SCoT-2 primers which differ only in one out of the last three nucleotides at the 3'end showed the highest Rp values were 5.99, 5.33 and 4.66 with polymorphism % and PIC of (69.23, 0.342), (80.00, 0.355), and (58.33, 0.222) respectively. Indicating

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the high discriminatory potential for this type of SCoT markers and which may be more successful in generating

reliable markers in grape as previously indicated by Prevost & Wilkinson (1999).

_		A	mplicons				Dolymounhio	
Duimon Nomo	Malaanlan		Polyn	orphic	_	Polymorphism	index content	Resolving
r miner manne	size range	Monomorphic	Without unique	Unique	Total	%	(PIC)	power Rp
SCoT-1	224:887	3	2	2	7	57.14	0.254	2.664
SCoT-2	269:1973	5	3	4	12	58.33	0.222	4.662
SCoT-3	310:1472	3	0	5	8	62.50	0.278	3.330
SCoT-4	259:1699	4	3	6	13	69.23	0.342	5.994
SCoT-6	359:724	4	1	3	8	50.00	0.22	2.664
SCoT-8	353:531	4	1	0	5	20.00	0.089	0.666
SCoT-10	310:905	2	3	5	10	80.00	0.355	5.328
SCoT-11	382:536	3	0	2	5	40.00	0.178	1.332
SCoT-12	314:640	2	3	0	5	60.00	0.267	1.998

 Table 8. Molecular data of SCoT molecular marker technique.

In comparison between combined molecular data for both SCoT and ISSR primers, it is evident from Table 9 that the total number of scorable ISSR amplicons was 35 with an average of 5.0 amplicons/primer, with a product size ranged from185 and 996 bp. While the total number of scorable SCoT amplicons was 75 with an average of 8.3 amplicons/primer, with a product size ranged from 224 and 1973 bp. Also, through better discrimination capabilities compared with ISSR, SCoT primers targeted generating 43 polymorphic amplicons with an average of 4.8/primer and 27 unique markers with an average of 3.0/primer. While ISSR primers except HB-12 targeted generating 18 polymorphic amplicons with an average of 2.6 /primer and 9 unique markers with an average of 1.3 /primer. This discrimination capability for the SCoT technique confirmed by P%, UM%, DI, and Rp values which were 55.24, 37.0, 0.25 and 3.18 respectively, compared with the same values for ISSR-technique which were 49.11, 25.7. 0.22 and 1.71, respectively. This indicates the high discriminatory potential of using SCoT primers compared with ISSR primers. Where, SCoT markers were more discriminating, provided more informative data. Also, confirms that it can be relying on the SCoT technique to evaluate the genetic diversity among the grape cultivars better than ISSR markers. More importantly, SCoT marker is generated from the functional region of the genome, so genetic analyses such as genetic diversity, genotype identification, construction of linkage maps and QTL mapping using this marker would be more useful (Hajibarat *et al.*, 2015).

Table 9. (Comparison	of discriminating	capacity	/ between]	ISSR and S	SCoT N	/Iolecular i	narkers techniques.
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Tashatana	MS	MS SA		PA		UA		D 0/		DI	D
rechnique		Total	Mean	Total	Mean	Total	Mean	P %	UIVI %	DI	кр
ISSR	185:996	35	5.0	18	2.6	9	1.3	49.11	25.7	0.22	1.71
SCoT	224: 1973	73	8.3	43	4.8	27	3.0	55.24	37.0	0.25	3.18
		11 4	11 1		1	TTA T	· ·		D 1 1 1	0/ ID	TO/ TT .

MS: Molecular size; SA: Scorable Amplicons; PA: Polymorphic Amplicons; UA: Unique Amplicons; P%: Polymorphism % ; UM%: Unique Marker %; DI: Diversity Index; Rp: Resolving power.

This result agrees with Gorji *et al.* (2011) in Potato and Etminan *et al.* (2016) in durum wheat, Abdel-Hameed *et al.* (2020), and Bashandy *et al.* (2020) in grape. They found that the SCoT marker was more informative and effective than the ISSR marker to estimate the genetic diversity and perform fingerprinting in these plants. While this result disagrees with Ramadan *et al.* (2019), who found that the ISSR marker is more discriminating and provides more informative data than SCoT in fennel cultivars.

While Baghizadeha and Dehghan (2018), and other researchers, recommend that it is preferable to use these molecular marker techniques in combination with each other for distinctive fingerprinting. Also, indicated that cluster analysis based on ISSR and SCoT data obviously discriminated among the Iranian pistachio cultivars. This was confirmed by Abd El-Aziz *et al.* (2019), who reported that the combined data of ISSR and SCoT molecular marker techniques were suitable and more informative for assessing the genetic relationships and genetic diversity among apricot strains.

Genetic distances estimation and cluster analysis

Genetic distances were estimated as Molecular and Phenotypic distances (MD & PD) based on the combined molecular data and quantitative data of studied characteristics respectively (Table 10). These data exhibited that the highest MD & PD were between Early Sweet and Star light grape cultivars. Whereas the lowest PD was between Prime and Early Sweet strain cultivars, the other molecular distances between Prime with Early Sweet or Star light cultivars were very close.

Table 10.	Dista	nces n	natrix	betwe	en Ea	arly S	weet, Pri	ime
	and	Star	light	grape	e cul	tivars	based	on
	com	bined	molec	ular (data	and	Phenoty	pic
	(nhv	cical a	nd Ch	omico	leha	ractor	istics) de	- ata

(physical and Chemical characteristics) data					
Distances n	natrix	Star light	Prime		
Drimo	Combined molecular data	0.260			
Fille	Phenotypic data	202.5			
Early Sweet	Combined molecular data	0.284	0.263		
Earry Sweet	Phenotypic data	280.0	80.4		

Also, from this matrix, the dendrograms of cluster analysis were performed using molecular and phenotypic distances (Figure 5). These dendrograms are divided into three groups according to the truncated line at a coefficient of dissimilarity= 0.13 & 40.18, respectively. Whereas each group was contained one cultivar in the two AHC dendrograms.



Figure 5. Agglomerative hierarchical clustering (AHC) dendrograms derived by UPGMA method using combined molecular data and phenotypic data of Early Sweet, Prime and Star light grape cultivars. Legend: TL represents truncated lines at a coefficient of dissimilarity or distance for molecular combined data of =0.13 & 40.18, respectively.

Association between molecular markers and distinguishing traits

Finally, by calculating the correlation coefficient between combined MD and PD, positive correlation with coefficient r=0.708 was found. This indicates that the unique markers which were most of them from SCoT markers (27 out of 36) can be associated with the distinguished traits of each cultivar as shown in Table 11 according to Abd El-Aziz *et al.* (2019). This is evidenced by Ibrahim *et al.* (2016), who explained that the SCoT markers were plausibly proved successful to target generic regions across the grape genome, suggesting future studies are needed to decipher the structure of markers which shown in Table 11. These markers after deciphering their structure may help to evolve more reliable molecular markers for the selection of the desirable traits in these cultivars. considering the results of the molecular techniques a distinctive genetic fingerprint for each cultivar. Hence the results of the molecular techniques especially SCoT can be considered a distinctive genetic fingerprint for these cultivars.

Table 11. The relationship between molecular markers and disting	guished traits for each cultivar.
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C14		Unique	Distinguished traits				
Culuvars	Technique	Primer	Molecular size bp	Total	Trait	Mean performance	
	ISSR	HB-15	417	1			
		SCoT 1	541		Bunch shape Berry shape Berry color	Conical	
		SCoT 2	956				
Stor light		SCoT 3	767,1472	12		Oblata	
Star fight	SCoT	SCoT 4	317, 720, 991	11 12		Bright red	
		SCoT 6	493,724				
		SCoT 10	517				
		SCoT 11	427				
	ISSR	14A	485,652,996	5			
		44B	362, 567	5			
		SCoT 1	451				
Prime	SCoT	SCoT 3	815,1076	13	Bunch shape	Winged	
		SCoT 4	574,1466	8			
		SCoT 6	512				
		SCoT 10	310, 555				
		44B	335		Maturity date Shot berries Bunch shape	Early (1 st week of June) High (22.61%) Shouldered	
	ISSR	HB-10	738	3			
		HB-15	422				
Farly Sweet	t	SCoT 2	403,1201,1973	11			
Larry Sweet		SCoT 3	481	11			
	SCoT	SCoT 4	811	8			
		SCoT 10	507,517				
		SCoT 11	444				

In conclusion, it is clear from the results that the Early Sweet, Prime and Star light grape cultivars succeeded under Egyptian conditions, which characterized by the high quality of bunch and berries. SCoT markers were more discriminating, provided more informative data, and evaluated the genetic diversity among the grape cultivars better than ISSR markers. So, we recommend expansion in the cultivation of these cultivars under Egyptian conditions and benefit from the results of the molecular techniques applied in this study to elicit reliable molecular markers characterizing desirable traits in these cultivars to use in Egyptian grape cultivars improving programs.

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وصف وتقييم بعض أصناف العنب المستوردة حديثاً تحت الظروف المصرية علا عبد الرحيم أحمد¹ و محمد حسن عبد العزيز² أقسم بحوث العنب – معهد بحوث البساتين – مركز البحوث الزراعية بالجيزة – مصر 2قسم الوراثة – كلية الزراعة – جامعة المنصورة- مصر

أجرى هذا البحث لمدة موسمين متثاليين (2017، 2018) بأحد المزارع الخاصة بمدينة السادات التابعة لمحافظة المنوفية لدراسة وصف وتقييم ثلاثة أصناف من العنب اللابذرية وهي: الإيرلي سويت، البرايم، الإستار لايت تحت الظروف المصرية. أظهرت النتائج أن صنف عنب الإيرلي سويت من أبكر الأصناف فيما يتعلق بالمواعيد الفينولوجية متمثلة في (تُقتّح البراعم، اكتمال التزهير، عقد الثمار ، نضّج الحبات)، يليها صنف عنب البرايم، بينما صنف عنب إستار لايت كان من الأصناف المتأخرة في هذا الصدد خلال موسمي الدراسة، كما أظهرت الدراسة نجاح هذه الأصناف تحت الظروف المصرية، حيث تميزت بصفات جودة عالية من العناقيد والحبات تم تقييم التنوع الوراثي بين الأصناف الثلاثة باستخدام تقنيات الواسمات الجزّيئية ScoT و ISSR. حيث قدمت علامات SCoT بيانات أكثر إفادة وكانت أكثر تمييزًا مع قيم عالية لنسبةً تباين حزم ونسبة العلامات جزيئية متفرده و ودليل تنوع و القدرة التميزية مقارنةً بعلامات ISSR. أظهر التحليل العنقودي تبايئًا جزيئيًا وظاهريًا واضحاً بين الأصناف الثلاثة، حيث تم تقسيمها إلى ثلاث مجموعات ، كل مجموعة تحتوي على صنف واحد أيضاً تميزت هذه الأصناف بـ 36 علامة جزيئية متفرده ، 27 منها كانت من علامات SCoT والتي أنبنت نجاحها في استهداف مناطق جينية منتوعة من جينوم العنب. قد تساعد هذه العلامات الجزيئية المتفر ده بعد فك تشفير تتابعاتها في تطوير علامات جزيئية موثوقة للإنتخاب للصفات المرّغوبة في هذه الأصناف. لذلك ، نوصى بالتوسع في زراعة هذه الأصناف تحت الظروف المصرية والاستفادة من نتائجنا الجزيئية لاستنباط علامات جزيئية موثوقة تُميز الصفات المرغوبة في هذه الأصناف لاستخدامها في برامج تحسين أصناف العنب المصرية.