# NEW INNOVATIVE PRODUCTS OF HUSK TOMATO FRUIT Nadir, A.; Wafaa. M. Abozeid and G. F. Bareh Food Sci. and Technology Dept., National Res. Centre, Cairo, Egypt.

# ABSTRACT

Two different dried product samples (sheet, raisins) were produced from fresh husk tomato. Chemical analysis, minerals, Hunter color and volatile components were determined for both two dried samples.

Results indicated a highly vit.C content(134.67mg/100g) in fresh fruit while in dried sample were (8.40mg/100g), maximum content of reducing sugars and fat in dried sheet sample (48.45 and 7.46 %) and high values of non-reducing sugars, total sugars, ash and fiber in raisins(31.14,71.81,7.95 and 20.04%) accompanied with higher levels of calcium, magnesium, phosphorus, potassium and sodium.

Drying process caused a reduction in lightness (L) values and increment in (a) values of redness and yellowness (b). Rehydration quality of raisins was improved and gave higher values of organoleptic characteristics than sheet samples. Methyl chavicol and cadinene (Gamma) were the two major volatile compounds found in fresh husk tomato fruits (54.1 and 14.75%). Many new volatile compounds were also found in the two dried products (sheet and raisins) where, Dill apiole compound was the main compound in sheet and raisins samples (64.1 and 66.6%) and reached to 2/3 percent from all the other compounds. Keywords: Husk tomato – dried sheet – raisins– volatile compounds

# INTRODUCTION

Husk tomato (Physaltis Philadelphica) is a solanceous plant cultivated in Mexico and Guatemala and originating from meso America. It is used in the diet since the pre-Columbian time; (Hernandez and Rivera 1994). Individual plants may produce 64 to 200 fruits in season and yielded approximately 9 tons per acre (4046 m<sup>2</sup>) as reported by (Wolff, 1991) and (Quiros, 1984). The fruit is small, spherical and green or green purple. They are surrounded by an enlarged calgx or "husk". As the fruit matures, it fills the husk and can splite it open by time of harvest. Tomatillos are available year round, produced in the U.S. mainlyon small – acreoges in California, with large volumes imported from Mexico (Smith *et al.*, 1999).

Drying of food stuffs is an important method for preservation and it is applicable to a wide range of industrial and agricultural products (Koyuncu, *et al.*, 2007). Also in most fruit, the skin is the solid part that contains the highest percentage of volatile compounds, which gave the fruit its aroma, and phenolic compounds, which are responsible for the color (Torre, *et al* 2010). Dehydrated skins generate interest because of their possible use as well as their addition to musts from grape harvests that are pron in volatile and phenolic metabolites (Torre, *et al.*, 2010).

Fruit can be dried in a food dehydrator, oven or in the sun by using the right combination of warm temperatures, low humidity and air current.

The optimum temperature for drying food is 140°F. Increasing temperatures caused cooking INS tend of drying (Paul 2009).

Color assessment of food is of great interest in the food industry and is made by visual or instrumental evaluation. The chromatic parameters

usually considered are: lightness (L\*), an attribute related to the transmission of light observed in the spectra; hue (hab), the qualitative expression of chromaticity; and Chroma (cab), the quantitative component of chromaticity (Osorio *et al.*, 2007).

The aim of this investigation was the application of drying process for husk tomato fruit to produce dried products (sheet and raisins), increase the consumption period and be available all over the year. Nutritional and sensory evaluations were also studied for both dried products.

# MATERIALS AND METHODS

### Materials:

Husk tomato fruits (physaltis philadelphica) were obtained from The Field Crops Research Institute, Agricultural Research centre, Ministry of Agriculture, Giza, Egypt.

### Methods:

### Preparation of the husk tomato Sheet and Raisins:

Husk tomato fruit (control) was sorted, washed and divided into two samples. The first sample was blended in a juice blender (National Solid state control model MJ. 170N) and the puree sample was treated with sodium meta-bisulphate (0.2%). The second sample was divided into two sample. One was treated with sodium meta- bisulphate(0.5%) for 20 min. and the other one was steamed for20 min. Then all samples ( control, puree and steamed) were placed in a flat trays and transferred to an air dry oven (shell lab 1370fx) at 65 °C for 6hr, then at 50 °C for6hr Fig.(1,2). All dried samples were packed in polyethylene bags and stored at room temperature (25 °C).



Fig. (1): Pretreatment and dehydration techniques of husk tomato sheets.



#### Fig.(2): pretreatment and dehydration techniques of husk tomato raisins

## Preparation and Isolation of volatile compounds: Reparation of husk tomato extract

One gram of husk tomato in fresh fruit or dried fruits were infused with 100 ml freshly boiled water for 5 min followed by filtration.

# Isolation of husk tomato volatile compounds:

The volatiles of husk tomato samples (fresh and dried) were isolated according to (Heath &Reineccius, 1986) using a dynamic husk tomato system, 100gm. Dried and fresh samples were subjected to extraction for four hours using diethyl ether and the extracted volatile compounds were dried over anhydrous sodium sulfate, evaporated and concentrated under gentle stream of nitrogen.

# Separation and Identification of volatile compounds:

The obtained volatile compounds were analyzed according to the method described by (Adams, 1995) using GC-MS apparatus. Separation was performed on thermo gas chromatograph (walnut creek, California, USA) equipped with Finnegan mat SSQ 7000 mass spectrometer and a 30mx. .025mm DB-5 Capillary column. The column temperature was programmed from 40°C (isothermal for min), to 300°C at rate of 5°C/min with 10 min. isothermal hold. The injector temperature was 220°C and the transition line temperature was 300°C. the carrier gas was helium and the column pressure head was 10-15 psi. the mass spectrometer had a delay of 3 min. to avoid

the solvent peak and then scanned from m/z<sup>50</sup> to m/z<sup>600</sup>. Ionization energy was set at 70eV. Identification of compounds was based on the comparison with the MS computer library (NIST and Wiley software package, Thermo Finnegan) and the published spectra. A linear retention index was calculated for each compound using the retention times of a homologous series of C6-C26 n-alkanes [Adams, 1995). Where no reference spectra were available, tentative identifications were made by comparison with spectra of related compounds.

### **Chemical Analysis methods :**

The content of moisture, protein, fat, fiber, ash, total solids, mineral and vitamin C were estimated according to the methods described by A.O.A.C. (2000). Total soluble solids (TSS %) were expressed as Brix value using a refract meter (ATAGO. Japan). Conversion factors were used to calculate protein, fat and total carbohydrate contents using 4, 9 and 4 k cal/g respectively and expressed in Kcal/100g (Pearson, 1991). **Drying Ratio:** 

The drying ratio was determined as reported by Van – Arsdel et al (1973). The drying ratio was calculated as follows:

Drving ratio –	Weight of wet
Drying ratio =	Weight of dried

### **Rehydration ratio:**

The rehydration ratio of dried samples was evaluated using the method of Ranganna (1979).

Ten grams dried sample were boiled in 500ml beaker. Covered by a watch glass for 5 min on an electrical heater, the contents were then dumped into a Buchner funnel for half to one min until the drip from the funnel was almost stopped and there hydration ratio was calculated as follows:

Debudration ratio	Weight of rehydrated sample
	Weight of dehydrated sample

Hydration coefficient = b  $(100-m1) / 100 (a - m_2)$ 

a = weight of the dehydrated sample.

b= the drained weight of the rehydrated sample.

 $m_1$  = moisture content (%) of the fresh sample.

 $m_2$  = moisture content (%) of the dehydrated sample.

## Color measurements:

The color of different samples was measured using a Spectorcolorimeter (tristimulus color machine) with CIE lab color scale (Hunter lab ScanXE, Germany).

Calibrated with a white standard tile of Hunter lab color standarad (LX No. 16379): X = 77.26, y = 81.94 and Z = 88.14, (L<sup>\*</sup> = 92.40, a<sup>\*</sup> = -0.87, b<sup>\*</sup>= -0.17). Color difference ( $\Delta E$ ) Hue angle and saturation index were calculated from a<sup>\*</sup>, b<sup>\*</sup> and L<sup>\*</sup> values (where a<sup>\*</sup> = redness, b<sup>\*</sup> = yellowness and I<sup>\*</sup>= lightness).

Using Hunter – Scotfield's equations (Hunter, 1975)  $\Delta E = (\Delta a^* + \Delta b^2 = \Delta L^2)^{1/2}$  .....(1)

<sup>510</sup> 

Where  $\Delta a = a$ - ao,  $\Delta b = b$ -bo and  $\Delta L = L$ -Lo

(ao, bo and Lo were the values of redness, yellowness and lightness for control sample).

 $\Delta E$  = color difference for a, b, L of sample from the same parameters of control.

Hue angle =  $\tan^{-1}(b/a)$  ......(2)

Saturation index =  $\sqrt{a^2 + b^2}$ .....(3)

# Sensory evaluation:

Products prepared from the different concentrations of the husk tomato fruits were evaluated for their appearance, color, taste and flavor as described by Meligaared *et al.*, (1991) and Fernands and Rodrignes (2007). Organoleptic characteristics results were statically analyzed according to Richard and Gouri (1987).

# **RESULTS AND DISCUSSION**

#### Chemical analysis of fresh and dried husk tomato products

Chemical analysis of fresh husk tomato fruit and its dried products was given in table (1). Results showed a higher content of total sugar (68.56%) in husk tomato sheet sample than fresh and raisin samples. While non-reducing sugar and reducing sugar content were high in raisins and husk tomato sheet samples (31.14,20.11,40.67and48.45%) respectively compared to all other samples (fresh, raisins) by Abozeid *et al.* (2007).

Table (1): Chemical composition of fresh husk tomato fruits and their dried products:

Constituents	Fresh hu: contr	sk tomato ol (%)	Dried sh	eets (%)	Dried raisins(%)		
(70)	Wet weight	Dry weight	Wet weight	Dry weight	Wet weight	Dry weight	
Moisture	91.32±0.87*	-	20.16±0.75	-	9.67±0.31	-	
Red. Sugars	3.12±0.18	35.94±0.31	38.69±0.63	48.45±0.36	36.74±0.61	40.67±0.51	
Non. Red. Sugars	2.38	27.42	16.09	20.11	28.13	31.14	
Total sugars	5.50±0.21	63.36±0.52	54.78±0.48	68.56±0.41	64.87±0.81	71.81±0.61	
Protein	0.49±0.15	5.64±0.12	5.44±0.12	6.81±0.32	6.12±0.13	6.78±0.11	
Fat	0.53±0.08	6.11±0.33	5.96±0.14	7.46±0.36	5.91±0.17	6.54±0.19	
Ash	0.64±0.03	7.37±0.22	6.21±0.20	7.66±0.38	7.18±0.14	7.95±0.17	
Fiber	1.52±0.09	17.51±0.45	8.11±0.81	10.15±0.48	18.11±0.20	20.04±0.22	
Vit. C (mg/100g)	11.69±0.87	134.67±1.18	6.71±0.43	8.40±0.29	7.50±0.19	8.30±0.14	

\*The obtained data expressed as a mean value ± standard Deviation.

There were no variation between protein content in dried sheet and raisins samples but, their values were higher than that of fresh samples. Higher values of fat content (7.46 and 6.54%) were observed in husk tomato sheet and raisins samples respectively and ash content was also high in raisins samples compared to other one.

The results also showed that, raisins contained higher levels of fiber (20.04%) than in fresh fruit (17.15%) and in dried sheets (10.15%) which

could be related to the losses during preparation method through cheese cloth. The reduction in vitamin C content was observed in dried sheet and raisins samples compared with both fresh husk tomato, and raisins samples. This may be due to exposure to heat treatment during processing. Results in table (2), showed that the mineral contents of raisins and dried sheets were relatively higher than in fresh husk tomato, potassium and phosphors appeared to be the higher amount than all other minerals content either raisins (753.48 and 46.58mg/100g) or in dried sheet samples (741.34 and 44.13 mg/100gm). The lowest components of copper and manganese were in raisins (0.28 and 0.27 mg/100gm) while it was (0.15 and 0.25 mg/100gm) in dried sheets respectively.

Elements	Fresh husk tomato	Dried husk tomato	Dried husk tomato
(mg/100g)	control	sheets	rasinis
Calcium	6.89	9.97	10.56
Iron	0.56	2.53	2.55
Magnesium	18.97	27.69	28.15
Phosphorus	37.85	44.13	46.58
Potassium	259.64	741.34	753.48
Sodium	0.92	1.89	2.87
Zinc	0.23	0.40	0.48
Copper	0.09	0.15	0.28
Manganese	0.18	0.25	0.27

Table (2): The minerals content of fresh husk tomato fruits and their dried products

Table (3) indicates the changes that occurred in the color values of all samples. It was clearly observed that the values of a and b of dried sheet samples were increased than that in fresh samples.

The darkness values could be related to the larger surface area exposure to heat during drying in an air oven  $(65^{\circ}C / 6 \text{ hr})$ . Which explain the more lightness than dried sheets. These results were is in agreement with Paul, (2009) and Osorio et al (2007).

Table (3): Hunter color values of fresh husk tomato fruits and their dried products

Sample	L	а	b	a/b	Saturation	Hue
Fresh Husk tomato (control)	45.64	12.48	49.78	0.25	51.32	75.93
Dried sheets	41.26	16.77	52.35	0.32	54.97	72.24
Dried raisins	43.74	15.28	50.38	0.30	52.65	73.13

Table (4) illustrated the significant differences in rehydration between raisins and dried sheet samples. It could be also noticed from table (4) that, moisture content of rehydration raisins (81.10%) exhibited lower values than the dried sheet samples (88.50%). This means that, the production of 1kg raisins from 4.2 kg fresh husk tomato was economically than that for production of 1kg dried sheets from 7.2 kg fresh sample.

Table (	(4):	Re	ehy	dratio	n qualit	y of	dried	d	husk	to	mat	0 9	shee	ets	and	rais	sins:	1
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Rehydration quality	Husk tomato sheets	Husk tomato raisins
Rehydration ratio	1:7.2 <sup>a</sup> ± 0.071	1 : 4.2 <sup>b</sup> ± 0.563
Moisture content of rehydrate sample	88.56 <sup>a</sup> ±0.112	81.10 <sup>b</sup> ±0.385
Hydration coefficient	1.56±0.040	0.95±0.063

- The obtained data expressed as a mean value ± standard Deviation.

- Alpha level of L.S.D = 0.01

Sensory evaluation of the dried sheet and raisins samples produced from fresh husk tomato fruit is shown in table (5). Fernands and Rodrigues (2007).

Results showed that raisins sample received the highest score value (96.43). There was a significant difference in flavor and color between dried sheet and raisins samples. There was also no significant difference in taste and general appearance between the two dried products produced from fresh husk tomato fruit.

Table (5): Organoleptic characteristics of husk tomato sheets and raisins

Characteristics	Score	Husk tomato sheets	Husk tomato raisins	L.S.D
Flavor	25	22.60 <sup>B</sup> ± 0.71	23.11 <sup>A</sup> ±0.64	0.47
Color	25	23.33 <sup>B</sup> ±0.53	24.12 <sup>A</sup> ±0.41	0.68
Taste	25	24.20±0.61	24.40±0.38	n.s
General appearance	25	24.60±0.48	24.80±0.52	n.s
Total	100	94.73	96.43	

n.s. = not significant

- L.S.D. = less significant diffewrence at 0.05 alpha level.

Results obtained in table (6) showed a variation in volatile compounds of fresh husk tomato. The first main compound was methyl chavicol (54.1%) while; the other components were cadinene (Gamma) and farnesene, (alpha) compounds were (14.75% and 8.1%) respectively. The other volatile compounds in fresh sample ranged from 3.5%-6.56% in which Eudesmol< $\gamma$ -Gpi-Alpha > was the lowest and Bisabolene< z-Gamma> was the highest. While Cubebol and Eudesmol ( $\gamma$ -Gpi-Alpha > compounds had relatively the same percent. Similar result was noticed by Torres et al (2007). For tetradecane and Bisabolene (Beta) compounds.

No	Compounds	RT	%
1	Methyl chavicol	22:30	54.1
2	Tetraldecane	22:76	4.58
3	Farnesene, (alpha)	23:21	8.10
4	Bisabolene (Beta)	23:53	4.89
5	Cadinene (Gamma)	24:85	14.75
6	Cubebol	25:53	3.6
7	Bisabolene< Z-Gamma>	28:26	6.56
8	Eudesmol<γ-EPi-Alpha>	28:62	3.5

Table (6): Volatile compounds in fresh husk tomato fruit.

Tables (7) and (8) showed the volatile compounds for both dried husk tomato and sheet and raisins. The percent values of several compounds such as furopelargone A, and Dill apiole were higher (7.4 and 66.6%) in dried husk tomato fruits raisin than in dried sheet sample. The opposite higher results were noticed for opolopenane (beta), unknown, hexadecane, 2, ethyl and Amyl ciannamldyde (E) in dried sheet of husk tomato. These results may be due to the effect of dehydration process which may break the cells or organelles that contained these compounds and thus, causing effective extraction, as the water decreased, the acidity would have increased, and hydrolysis of these volatile compounds, increased causing the release of these compounds (Torre et al., 2010).

Also, the volatile compound Furopelargone A, Oplopenane (beta), Hexadecane, 2, ethyl and Amyl cinnamaldhyde (E) had the same concentration (7.4)% in dried husk tomato with different retention times (RT). The results also indicated that, the volatile component dill apiole was the main compound in both dried and husk tomato sheets being (66.6 and 64.1%) respectively.

No	Compounds	RT	%
1	Furopelargone (A)	32:16	6.5
2	Opolopenane (beta)	35:18	7.6
3	Dill apiole	37:22	64.1
4	Un Known	40:11	3.9
5	Hexadecane, 2, ethyl	44:17	8.3
6	Amyl ciannamldhyde (E)	49:12	8.6

Table (7): Volatile compounds of dried husk tomato sheet.

Table (8)	: Volatile co	mpounds of	f dried husk	tomato fruits	s raisins
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No	Compounds	RT	%
1	Furopelargone (A)	28:36	7.4
2	Opolopenane (beta)	30:54	7.4
3	Dill apiole	31:67	66.6
4	Un Known	34:75	3.7
5	Hexadecane, 2, ethyl	39:66	7.4
6	Amyl ciannamldhyde (E)	40:61	7.4

### Conclusion

The present study indicated that husk tomato fruit could be successfully dried to raisins and sheet products having the desired quality of color, taste and rehydration ratio.

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منتجات جديدة مبتكرة من ثمار الحرنكش عبد العزيز ندير ، وفاء محمد ابو زيد و جميل فايز بارح. قسم الصناعات الغذائية - المركز القومي للبحوث – القاهرة – مصر

تم في هذا البحث أنتاج ٢ منتج مختلف (اللفائف و الزبيب) بالتجفيف من الثمار الطازجة للحرنكش و قد تم تقدير التركيب الكيماوي و المعدني و القيم اللونية و كذلك المركبات المتطايرة في كل من العينات المجففة للمنتجين. و قد اوضحت النتائج ارتفاع محتوي فيتامين ج (١٣٤,٦٧) كل من العينات المجففة للمنتجين. و قد اوضحت النتائج ارتفاع محتوي فيتامين ج (١٣٤,٦٧) كل من العينات المحففة للمنتجين. و قد اوضحت النتائج ارتفاع محتوي فيتامين ج (١٣٤,٦٧) للفائف الطازجة و أعلي محتوي للسكريات المختزلة و كذلك الدهن في عينة اللفائف الجافة (٧,٤٦،٤٨٤) هو كذلك قيم عالية للسكريات المختزلة و للسكريات الكلية و اللفائف الجافة (٧,٤٦،٤٨٤٥) هو كذلك قيم عالية للسكريات الغير مختزلة و السكريات الكلية و اللماد و الألياف في الزبيب (٧,٤٦،٤١٦،٢١٢٤) و كناك الصوديوم . عملية التجفيف احدثت أنخفاض الكالسيوم , الماغنسيوم , الفسفور , البوتاسيوم و كذلك الصوديوم . عملية التجفيف احدثت أنخفاض الكالسيوم , الماغسيوم , الماغسور , البوتاسيوم و كذلك الصوديوم . عملية التجفيف درثات الخلية و مركبي لقيم (لما دو الألياف في الزبيب تحسنت و اعطت اعلي القيم للصفات الحسية عن عينةاللفائف. و مركبي و تعمر الربيب (١٤,٢٥،٥٤) هو كذلك الصوديوم . عملية التجفيف احدثت أنخفاض الكالسيوم , الماغسيوم , الفسفور , البوتاسيوم و كذلك الصوديوم . عملية التجفيف احدثت أنخفاض الميثل شافيكول الوالات في الوري الذي العمر المودينين (جاما) الحرنكش الطازجة (١٤,٧٥،٥٤) هو قد وجد العديد من مركبات النكهة الجدينين(جاما) الحرنكش الطازجة (١٤,٦٥،٥٤) هو الميثل شافيكول الفائف و الكادينين(جاما) الحرنكش الطازجة (١٤,٦٥،٥٤) هو قد وجد العديد من مركبات النكهة الجديني في كلا المنتجين (اللفائف و الزبيب) مركب الذي يصل و قد وجد العديد من مركبات النكهة الجدينية في كلا المنتجين (اللفائف و الزبيب) وي الزبيب) مركب المركبات النكهة الجدينة في كلا المنتجين (اللفائف و مركبي و قد وجد العديد من مركبات النكهة الجدينين(جاما) الحرنكش الطازجة (٦٤,٦٥،٦٤) هو الميثل الميثلي الفيكول الفائف و الزبيب (٢٤,٦٩) مركب كان مركب الميثيب في عينات اللفائف و الزبيب (١٤,٦٠،٢٦) مركب الى نوى مركبي ال الغوني مركبات النكهة الجديني والكان و مركبي الحرنكش الطازجة (٢٦,٦٦،٦٤) مركب كان مركب الميثيب في عرك المنتجي (اللفائف و الزبيب) مركب الذي ي مركب المي مالغيب (١٤,٢٥،٢٤،٢٥) مرك

قام بتحكيم البحث

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