

**EFFECT OF SOME COMPONENTS OF VOLATILE OILS, SODIUM CARBONATE AND BICARBONATE FOR CONTROLLING BLUE AND GREEN MOLDS OF Lime (*Citrus aurantifolia*) FRUITS DURING STORAGE**

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**ABSTRACT**

Blue and green molds caused by *Penicillium italicum* Wehmer and *Penicillium digitatum* Sacc. are the most important diseases affecting lime fruits. Some component of volatile oils i.e. limonene, linalool, citral and nerol were extracted from orange peel, coriander seed, lemon grass and petitgrain (Bitter orange) respectively, and were tested for their effect on linear growth of *P. italicum* and *P. digitatum*. Nerol and citral at 0.4% showed 100% reduction of linear growth. Sodium carbonate (SC) and sodium bicarbonate (SBC) showed complete inhibition on linear growth of the tested fungi at the concentration 2.0%.

Effects of citral and SC or SBC singly or in combination for controlling blue and green molds of lime fruits were studied. The most effective treatments were the combined treatments between citral at 0.5% and SC or SBC at 2.0% which completely inhibited disease severity for blue and green molds of lime fruits under artificial infection and during storage for three weeks at 13± 1°C. However under natural infection, the combined treatments with citral at 0.5% and SC or SBC at 2% completely inhibited percentages of infection during storage for three weeks. All treatments decreased the loss of fresh weight of lime fruit and vitamin C content but total soluble solids increased.

**Keywords:** Blue mold, green mold, lime fruits, some component of volatile oils, sodium carbonate and bicarbonate,

**INTRODUCTION**

Green and blue molds, caused by *Penicillium digitatum* Sacc. and *P. italicum* Wehmer are the most damaging postharvest diseases of citrus fruits. Postharvest decay of lime fruits caused by *P. digitatum* and *P. italicum* are important postharvest storage problems (Smilanick and Arruf, 1992). Volatile oils of citrus or their constituents are shown to have a fungicidal activity against postharvest pathogens of citrus (Caccioni *et al.*, 1998). Also he noted that a positive correlation between monoterpenes other than limonene and sesquiterpene content of the oils and the pathogenic fungi inhibition. Volatile oils or their constituents of citrus fruit show complete inhibition of *P. digitatum*, *P. italicum*, (Caccloni *et al.*, 1998). The main constituents of volatile oils are mono-and sesquiterpenes and some of these compounds have shown antibacterial, antifungal and antioxidant activities (Lee and Ahn, 1998). Sweet orange (*Citrus sinensis* L. Rutaceae), contains volatile oil extracted from fruit peels which is characterized by high limonene content, the fruits (seeds) of coriander (*Coriandrum sativum*,L. Apiaceae) contain volatile oil highly constituted with linalool as a main component. Lemongrass (*Cymbopogon citratus* (DC.) Stapf., Poaceae) contains mainly citral as a main component of

volatile oil extracted from leaves (Schaneberg and Khan, 2002) and 1 to 2% essential oil on a dry basis (Carlson *et al.*, 2001). Bitter orange (petitgrain) oil (*Citrus aurantium* var. *amara*, *Rutaceae*) contains nerol. It showed great inhibitory effect on the linear growth of *Alternaria solani* (Abd-El-kareem, 2007). Droby *et al.*, (2008) found that citral, exhibited a stimulatory effect to *P. digitatum* at 0.005 and 0.01 $\mu$ l/plate and completely inhibited both germination and germ tube growth above those concentrations while, in *P. italicum*, germination and growth were stimulated at concentrations up to 0.1 $\mu$ l/plate but were inhibitory at the higher concentrations. Compared to citral, linalool, significantly stimulated germination of *P. digitatum* and *P. italicum* at higher amounts (up to 2 $\mu$ l/plate). Also he found that limonene, myrcene,  $\alpha$  pinene and  $\beta$ . pinene were stimulatory to *P. digitatum* and *P. italicum* but inhibitory to or had no effect on *P. expansum* and *Botrytis cinerea*.

Treatments with sodium carbonate and sodium bicarbonate provided satisfactory control of green mold of lemons (Smilanick *et al.*, 1997) and blue mold of orange (Palou *et al.*, 2001). Sodium carbonate reduced the incidence of citrus green and blue mold (Palou *et al.*, 2001 and Palou *et al.*, 2009). Latifa *et al.* (2011) showed that among 28 tested compounds, sodium carbonate, ammonium carbonate, copper sulfate, EDTA and sodium metabisulfate have completely inhibited the mycelial growth and sporulation of *P. italicum* at only 0.02M.

The present study was performed to evaluate the effect of some components of volatile oils (nerol, citral, linalool and limonene) and sodium carbonate or sodium bicarbonate on linear growth of *P. digitatum* and *P. italicum* *in vivo* and studying *in vitro* control blue and green molds of lime fruits.

## **MATERIALS AND METHODS**

Isolates of both *Penicillium digitatum* and *P. italicum* were isolated from infected lime fruits shown green and blue molds collected from the local market of Sharkia Governorate. For isolation, samples were introduced and carefully washed with running tap water, dried between filter papers and surface sterilized with 1% sodium hypochlorite for 2 min., then the samples were washed thoroughly with sterilized distilled water for several times and dried between sterilized filter paper. Sterilized samples were cut into small pieces and separately transferred to sterilized Petri-dishes containing potato dextrose agar (PDA) media and incubated for 7 days at 27°C. Isolates was identified according to Raper and Thom (1968).

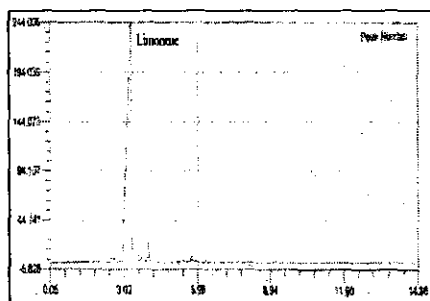
### **Extraction and isolation of volatile oils and its components:**

Four volatile oils which containing the components used in this study i.e. limonene, linalool, citral and nerol were extracted from orange peel, coriander fruits, lemon grass and petitgrain (Bitter orange) respectively using water distillation as described in the British Pharmacopoeia (1963). The Extraction and isolation of volatile oils and its components were carried out at Medicinal and Aromatic Plants Research Department Laboratory, Horticulture Research Institute, Agriculture Research Center, Giza, Egypt. The volatile oils obtained from each sample were analyzed using Ds Chorm 6200 gas

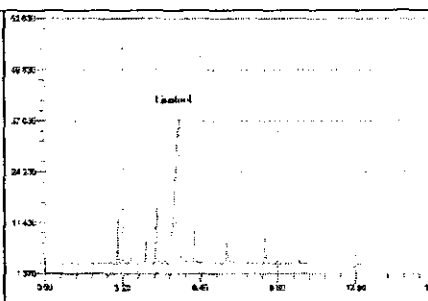
chromatography equipped with a flame ionization detector for separation of volatile oil constituents. The analysis conditions were as follows:

-The chromatograph apparatus was fitted with capillary column BPX-5, 5% phenyl (equiv.) polysilphenylene-siloxane 30 m x 0.25 mm ID x 0.25  $\mu$ m film.

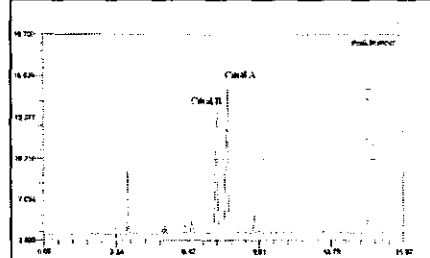
-Temperature program ramp increase with a rate of 10°C/ min from 70 to 200°C. Flow rates of gases was nitrogen at 1 ml/min, hydrogen at 30 ml/min and 330 ml/min for air. Detector and injector temperatures were 300°C and 250°C, respectively. The obtained chromatogram and report of GC analysis for each sample were analyzed to calculate the percentage of main components of volatile oil. Gas chromatography analysis for orange peel, coriander fruits (seeds), lemon grass and petitgrain volatile oils indicating that the content of limonene, linalool, citral and nerol were 91.75%, 71.20%, 86.32 % and 4.09% respectively as shown in Figs. from 1to 4.



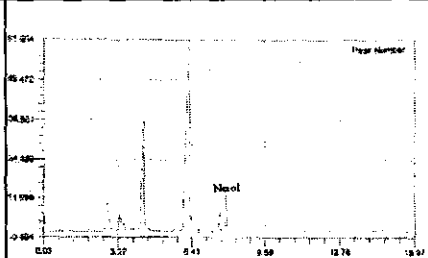
**Fig. (1): Gas chromatography analysis for orange peel volatile oil indicating to limonene content (91.75%)**



**Fig. (2): Gas chromatography analysis for coriander fruits volatile oil indicating to linalool content (71.20%)**



**Fig. (3): Gas chromatography analysis for lemongrass volatile oil indicating to citral (citral b+citral a) content(86.32 %)**



**Fig. (4): Gas chromatography analysis for petitgrain volatile oil indicating to nerol content (4.09%)**

**Isolation of volatile oils component:**

**Limonene isolation:**

Limonene isolated from orange peel volatile oil into a low density water immiscible solvent (ether). The lower layer is the remaining aqueous distillate evaporation of the solvent to finalize the extraction; the ether layer (boiling point, 37°C) was evaporated on a water bath to leave the limonene (boiling point, 176°C). Distillate limonene (150 mg) obtained from 15 g of orange peel skin.

**Linalool isolation:**

Linalool is the major component of the essential oil of coriander fruits, according to Kamel *et al.* (2007) linalool can be isolated directly from the essential oil with 2-methyl-butane (v/v) and dried over anhydrous sodium sulphate. The organic layer was then concentrated, at 30°C using a Vigreux column resulting linalool.

**Citral isolation**

Citral isolated from lemon grass volatile oil by transferring 3 ml of lemon grass oil, 75 ml of water to a 250-ml round-bottom flask. Place a boiling chip into the flask. Connect the flask to a simple distillation apparatus. Heat the mixture to boiling, carefully at first to avoid excessive foaming, which could spill over into the condenser and into the graduated cylinder, using graduated cylinder to collect the distillate. Isolate the citral obtained from mixture of water and citral by extracting it with diethyl ether; evaporate the ether layer to get citral.

**Nerol isolation**

The oil of petitgrain contains primary alcohols and terpene which saponified with the alcoholic potassium phthalic anhydride. The primary alcohols transformed into acid phthalates or other acid esters of dibasic acids. After purification and saponification with alkali, a geraniol-nerol mixture containing 40% nerol and 60% geraniol is obtained. geraniol content is removed by treatment with calcium chloride to geraniol-nerol mixture, in the form of crystalline geraniol compound with calcium chloride, the crystals are removed by filtration. Now, crude nerol as obtained is rectified by dry distillation in vacuum or steam distillation. The yield of pure nerol will be about 2 to 3% weight of the petitgrain oil, (NIIR BOARD, 2003).

**A- In vitro:**

**Effect of component of volatile oils and sodium carbonate or bicarbonate on linear growth of rot fungi of lime fruits:**

Antifungal activity of four components of citrus essential oils i.e., Citral, limonene, linalool and nerol and sodium carbonate (SC) or bicarbonate (SBC) were studied by determining their inhibitory effect on growth of *Penicillium digitatum* and *P. italicum*. Five concentrations i.e 0.1, 0.2, 0.3, 0.4 and 0.5% were tested for each component, Then each concentration was added aseptically to sterile molten PDA medium (45°C) containing Tween 80 (0.5% v/v). The resulting media were immediately dispensed into sterilized Petri Plates (9 cm). In controls, water and Tween 80 instead of volatile oils.

Sodium carbonate and sodium bicarbonate at five concentrations 0.5, 1.0, 2.0 and 3.0% were used. Stock medium of PDA containing the tested concentration was poured into three Petri-Plates after solidification. Three Petri-Plates containing PDA medium without salts were used as control. Discs 3- mm diameter obtained from 7 day- old of each fungus was aseptically transferred to the center of Petri Plates containing volatile oils, salts and plates served as control. All Petri-Plates were incubated at 25°C ± 2°C. The average diameter in mm of each fungus and treated was measured when the fungal growth has covered the check treatment plates.

**B- In vivo:**

Fresh lime fruits cv. Baladi apparently free from physical damage and diseases were washed, disinfected in 0.5% (v/v) sodium hypochlorite, rinsed three times in sterile distilled water and then air dried. The effect of dipping lime fruits in citral oil, sodium carbonate and sodium bicarbonate was studied. This was studied in the naturally infected (uninoculated), fruits, as well as in artificially inoculated fruits with *Penicillium digitatum* and *P. italicum*. lime fruits were wounded with a sterilized needle and inoculation was carried out using a needle, immersed in the spore suspension (10<sup>6</sup> spores/ ml) of each *P.* and *P. italicum*. Wounded fruits inoculated with sterilized water were applied as control. After 24 hours, the inoculated fruits were dipped in sodium carbonate or bicarbonate at 2.0% as a single. Such treatment was also applied as combined with citral as follows:

Treatment	
Single	Combined
NaCO <sub>3</sub> (SC) 2.0%	Citral 0.3% + SC 2.0%
	Citral 0.5% + SC 2.0%
NaHCO <sub>3</sub> (SBC) 2.0%	Citral 0.3% + SBC 2.0%
	Citral 0.5% + SBC 2.0%
Citral 0.3%	
0.5%	
Control (untreated Fruits)	

Control (untreated) was dipped in sterile distilled water. Fruits were dried at room temperature 20 ± 2°C. All treated or untreated (control fruits) were placed in a plastic tray in cardboard boxes and stored at 13± 1°C for three weeks.

Nerol did not used because it caused bad effect on lime fruits.

**Disease assessment:**

Percentage of rots disease severity was assessed 1, 2, 3 weeks. Disease severity (D.S) was calculated using the scale from 0 – 4 (0 = healthy, 1= 1-25% infections, 2 = 26 – 50% infection, 3 = 51 – 75% infections and 4 = 76 – 100% infection and formula adopted by Hanounik (1986) as follows:

$$\text{Disease severity (\%)} = \frac{\sum (\text{NCC} \times \text{CR})}{\text{NTC} \times \text{MSC}} \times 100$$

Where: NCC = No. of lime fruits in each class rate.

NTC = No. of total lime fruits examined

CR = Class rate

MSC = Maximum disease severity class rate (4).

$$\text{Infection percentage \%} = \frac{\text{Number of decayed fruits}}{\text{Initial number of stored fruit}} \times 100$$

Efficacy (E) of each treatment was calculated as follows:

$$E = \frac{\text{D.S control} - \text{D.S treated}}{\text{D.S control}} \times 100$$

**Loss in weight:**

Loss in lime fruit fresh weight % was estimated in uninoculated lime (under natural infection) fruits of various treatments according to the following equation:

$$\text{Fruit weight loss \%} = \frac{w_i - w_s}{w_i} \times 100$$

$w_i$  = fruit weight at initial period.

$w_s$  = fruit weight at sampling period.

**Total soluble solids (TSS):**

The percentage of total soluble solids (TSS) was determined by a hand refractometer

**Vitamin C (ascorbic acid):**

Extracts from lime fruits were obtained by blending tissues in a blender for 5 minutes containing oxalic acid 6% equivalent to the tissues (v/w). The tissues were strained through several layers of cheesecloth. The liquid fraction was then centrifuged for 5 min at 3000 rpm at room temperature and the filtrate was used to determine ascorbic acid. Twenty ml of filtrate was brought to volume 100 ml of oxalic acid 3% and titrated with stain (2, 6- dichlorophenol endophenol) until pink color. Amount of ascorbic acid was calculated according to the following formula, (Jemey and Kovacs, 1968).

$$\text{Ascorbic acid with milligrams} = \frac{V \times C \times D}{\text{Weight of sample with gram}} \times 100$$

Where: V = volume of stain

C = concentration of stain

D = dilution of sample.

**Statistical analysis:**

The data were statistically analyzed using MSTAT-C computer program v.2. 10 (1988).

## RESULTS

Four volatile oils which containing the components used in this study i.e. limonene, linalool, citral and nerol were extracted from orange peel, coriander fruits, lemon grass and petitgrain (Bitter orange) respectively.

**A- In vitro:**

Data in Table (1) indicate that all the tested components of citrus essential oil (Citral, limonine, linalool and nerol), sodium carbonate (SC) and sodium bicarbonate (SBC) at different concentrations have significantly reduced linear growth of *Penicillium digitatum*. The reduction in growth was correlated to the increase in component of volatile oil, SC and SBC concentrations. All tested components of volatile oil caused 100% inhibition on linear growth of *P. digitatum* at 0.4% except limonene who caused 94.4 % reduction of linear growth. Nerol and citral at 0.3% gave the highest effect on reducing the growth of *P. digitatum*, but limonene gave the least effect at the same concentrations. Also, the used SC or SBC have completely inhibited growth of *P. digitatum* at 2.0 % and at 0.5% gave least effect on growth of tested fungus.

**Table 1: Effect of some components of volatile oil, sodium carbonate (SC) and sodium bicarbonate (SBC) on the linear growth of *Penicillium digitatum***

Treatments	Linear growth (mm)									
	Concentration (%)									
	0.1	R%	0.2	R%	0.3	R%	0.4	R%	0.5	R%
Citral	25.5 <sup>d</sup>	71.6	10.5 <sup>f</sup>	88.3	7.0 <sup>e</sup>	92.2	0.0 <sup>c</sup>	100.0	0.0 <sup>b</sup>	100.0
Limonene	45.6 <sup>b</sup>	49.3	27.3 <sup>b</sup>	69.7	10.0 <sup>c</sup>	88.9	5.0 <sup>b</sup>	94.4	0.0 <sup>b</sup>	100.0
Linalool	46.0 <sup>b</sup>	48.9	15.0 <sup>d</sup>	83.3	8.3 <sup>d</sup>	90.6	0.0 <sup>c</sup>	100.0	0.0 <sup>b</sup>	100.0
Nerol	18.3 <sup>s</sup>	79.7	8.6 <sup>d</sup>	90.4	5.0 <sup>f</sup>	94.4	0.0 <sup>c</sup>	100.0	0.0 <sup>b</sup>	100.0
	0.5		1.0		1.5		2.0		3.0	
SC	30.0 <sup>c</sup>	66.7	13.5 <sup>e</sup>	85.0	9.3 <sup>c</sup>	89.7	0.0 <sup>c</sup>	100.0	0.0 <sup>b</sup>	100.0
SBC	45.0 <sup>b</sup>	50.0	20.9 <sup>c</sup>	76.8	12.1 <sup>b</sup>	86.6	0.0 <sup>c</sup>	100.0	0.0 <sup>b</sup>	100.0
Control	90.0 <sup>a</sup>		90.0 <sup>a</sup>		90.0 <sup>a</sup>		90.0 <sup>a</sup>		90.0 <sup>a</sup>	

R= Reduction

Values having the same letters in the same column are not significantly different at P<0.05

Data in Table (2) show that all the tested components of volatile oils, SC and SBC significantly reduced the linear growth of *P. italicum*. The inhibitory effect on growth was increased by increasing the concentrations. Complete inhibition of linear growth of *P. italicum* was obtained with nerol and citral at 0.4 % which reduced growth by 100%, while the least reduction was obtained with limonine. All concentrations of SC or SBC reduced growth of *P. italicum* compared with control. The SC and SBC completely inhibited the linear growth at 2.0 % causing 100% reduction.

**B- In vivo:**

**Effect of single or combined applications of sodium carbonate (SC) or sodium bicarbonate (SBC) and citral on green mold severity of lime fruits**

Sodium carbonate (SC) or sodium bicarbonate (SBC) at 2.0% alone or in combination with citral at 0.3% or 0.5% were tested under storage conditions to study their effect on green mould severity of lime fruits. Data in Table 3 indicate that all treatments have completely inhibited disease severity for one week of storage at 13 ± 1°C.

**Table 2: Effect of some components of volatile oil, sodium carbonate (SC) and sodium bicarbonate (SBC) on linear growth of *Pencilium italicum***

Treatments	Linear growth (mm)									
	Concentration (%)									
	0.1	R%	0.2	R%	0.3	R%	0.4	R%	0.5	R%
Citral	35.0 <sup>b</sup>	61.1	17.5 <sup>a</sup>	80.6	6.5 <sup>d</sup>	92.8	0.0 <sup>d</sup>	100.0	0.0	100.0
Limonine	51.5 <sup>b</sup>	42.8	27.5	69.4	15.3 <sup>b</sup>	83.0	6.7 <sup>b</sup>	92.6	0.0	100.0
Linalool	43.5 <sup>a</sup>	51.7	23.5 <sup>a</sup>	73.9	12.0 <sup>c</sup>	86.7	4.8 <sup>c</sup>	94.7	0.0	100.0
Nerol	25.5 <sup>a</sup>	71.7	13.5 <sup>a</sup>	85.0	5.3 <sup>a</sup>	94.1	0.0 <sup>d</sup>	100.0	0.0	100.0
	0.5		1.0		1.5		2.0		3.0	
SC	40.0 <sup>d</sup>	55.6	15.7 <sup>a</sup>	82.6	4.5	95.0	0.0 <sup>d</sup>	100.0	0.0	100.0
SBC	42.3 <sup>c</sup>	53.0	20.7 <sup>d</sup>	77.0	5.6 <sup>c</sup>	93.8	0.0 <sup>d</sup>	100.0	0.0	100.0
Control	90.0 <sup>a</sup>		90.0 <sup>a</sup>		90.0 <sup>a</sup>		90.0 <sup>a</sup>		90.0 <sup>a</sup>	

R= Reduction

Values having the same letters in the same column are not significantly different at P<0.05

Data in Table 3 also indicated that combined treatment between citral and SC or SBC have significantly reduced the lime rot severity. The most effective treatments were combined treatment between citral at 0.5% and SC or SBC at 2% which completely inhibited the disease severity after two or three weeks storage, while single treatments of each concentration of SC or SBC at 2% and citral at 0.5% reduced disease severity more than 75.1% and 82.1% after two or three weeks, respectively. Generally all treatments decreased disease severity during storage up to three weeks compared to control. High efficacy was obtained with citral at 0.5% combined with SC or SBC at 2% during storage at 13 ± 1 °C for three weeks.

**Table 3: Effect of sodium carbonate (SC) or sodium bicarbonate (SBC) applied as single or in combination with citral on green mold severity of lime fruits during storage at 13 ± 1°C for 3 weeks**

Treatments	Storage period (weeks)		Disease severity (%)						
			Green mold						
	1	2	1		2		3		
	D.S	E	D.S	E	D.S	E	D.S	E	
Single treatment									
Citral 0.3%	0.0 <sup>b</sup>	100.0	8.9 <sup>b</sup>	75.1	12.7 <sup>b</sup>	83.2			
Citral 0.5%	0.0 <sup>b</sup>	100.0	4.9 <sup>a</sup>	86.2	7.7 <sup>d</sup>	89.8			
SC 2%	0.0 <sup>b</sup>	100.0	7.7 <sup>c</sup>	78.4	10.9 <sup>c</sup>	85.6			
SBC 2%	0.0 <sup>b</sup>	100.0	8.3 <sup>bc</sup>	76.8	13.5 <sup>d</sup>	82.1			
Combined Treatments									
Citral 0.3% + SC 2%	0.0 <sup>b</sup>	100.0	6.2 <sup>d</sup>	82.6	9.5 <sup>c</sup>	87.4			
Citral 0.5% + SC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>d</sup>	100.0	0.0 <sup>f</sup>	100.0			
Citral 0.3% + SBC 2%	0.0 <sup>b</sup>	100.0	3.6 <sup>f</sup>	89.9	5.8 <sup>a</sup>	92.3			
Citral 0.5% + SBC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>d</sup>	100.0	0.0 <sup>f</sup>	100.0			
Control	15.5 <sup>a</sup>		35.7 <sup>a</sup>		75.6 <sup>a</sup>				

Values having the same letters in the same column are not significantly different at P<0.05

D.S = Disease severity

E= Efficacy



**Effect of single or combined application of sodium carbonate (SC) or sodium bicarbonate (SBC) and citral on blue mold severity of lime fruits:**

Data in Table 4 indicated that the most effective treatments were citral at 0.5% combined with SC or SBC at 2%. On the other hand all treatments have completely inhibited disease severity of blue mold after one week, while after two or three weeks, it was completely inhibited with the combined treatment citral at 0.5% with SC or SBC at 2% and efficacy reached 100% during the cold storage at  $13 \pm 1^\circ\text{C}$ . The combined treatment has significantly reduced the disease severity after two weeks.

**Table 4: Effect of sodium carbonate (SC) or sodium bicarbonate (SBC) applied as single or in combination with citral on blue mold severity of lime fruits during storage at  $13 \pm 1^\circ\text{C}$  for 3 weeks**

Storage period (weeks)	Disease severity (%)					
	Blue mold					
	1		2		3	
Treatments	D.S	E	D.S	E	D.S	E
Single treatment						
Citral 0.3%	0.0 <sup>b</sup>	100.0	7.1 <sup>c</sup>	83.6	10.9 <sup>d</sup>	85.9
Citral 0.5%	0.0 <sup>b</sup>	100.0	5.3 <sup>d</sup>	87.7	8.9 <sup>e</sup>	88.6
SC 2%	0.0 <sup>b</sup>	100.0	8.9 <sup>b</sup>	79.4	12.9 <sup>c</sup>	83.4
SBC 2%	0.0 <sup>b</sup>	100.0	9.2 <sup>b</sup>	78.7	15.1 <sup>d</sup>	80.6
Combined Treatments						
Citral 0.3% + SC 2%	0.0 <sup>b</sup>	100.0	4.6 <sup>e</sup>	89.4	7.7 <sup>e</sup>	90.1
Citral 0.5% + SC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>f</sup>	100.0	0.0 <sup>f</sup>	100.0
Citral 0.3% + SBC 2%	0.0 <sup>b</sup>	100.0	2.8 <sup>f</sup>	93.5	6.2 <sup>f</sup>	92.0
Citral 0.5% + SBC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>f</sup>	100.0	0.0 <sup>f</sup>	100.0
Control	20.2 <sup>a</sup>		43.2 <sup>a</sup>		77.8 <sup>a</sup>	

D.S=Disease severity

E= Efficacy

Values having the same letters in the same column are not significantly different at  $P<0.05$

**Effect of single or combined application of sodium carbonate (SC) or sodium bicarbonate (SBC) and citral on percentage of infection of lime fruits under natural infection**

Data in Table (5) showed the effect of citral oil and SC or SBC with single or combined treatment on the percentage of infection under natural infection of lime fruits. All treatments have completely inhibited infection of lime fruits for one week of storage at  $13 \pm 1^\circ\text{C}$ . High efficacy 100% was obtained from the single and the combined treatments at same storage period. The combined treatment significantly reduced disease severity after two or three weeks. The treatment combined with citral at 0.5% with SC and SBC at 2% completely inhibited infection percentage after three weeks of storage.

**Table 5: Effect of citral and sodium carbonate (SC) or sodium bicarbonate (SBC) applied as single or in combination on percentage of infection (Inf.%) of lime fruits during 3 weeks  $13 \pm 1^\circ\text{C}$  under natural infection**

Storage period (weeks)	Infection (Inf.%)					
	1		2		3	
	Inf.%	E	Inf.%	E	Inf.%	E
<b>Single treatment</b>						
Citral 0.3%	0.0 <sup>b</sup>	100.0	0.0 <sup>b</sup>	100.0	5.3 <sup>c</sup>	85.2
Citral 0.5%	0.0 <sup>b</sup>	100.0	0.0 <sup>b</sup>	100.0	1.8 <sup>d</sup>	94.9
SC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>b</sup>	100.0	6.4 <sup>b</sup>	82.1
SBC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>b</sup>	100.0	5.8 <sup>bc</sup>	83.8
<b>Combined treatments</b>						
Citral 0.3% + SC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>b</sup>	100.0	1.8 <sup>d</sup>	94.9
Citral 0.5% + SC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>b</sup>	100.0	0.0 <sup>e</sup>	100.0
Citral 0.3% + SBC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>b</sup>	100.0	2.2 <sup>d</sup>	93.8
Citral 0.5% + SBC 2%	0.0 <sup>b</sup>	100.0	0.0 <sup>b</sup>	100.0	0.0 <sup>e</sup>	100.0
Control	7.5 <sup>a</sup>		16.7 <sup>a</sup>		35.7 <sup>a</sup>	

E= Efficacy

Values having the same letters in the same column are not significantly different at  $P < 0.05$

**Effect of sodium carbonate or sodium bicarbonate applied as single or in combination with citral on some lime fruits quality under natural infection**

All treatments lowered the loss in weight in fresh weight of lime fruits compared to control after three weeks of storage as shown in (Table 6). Combined treatments showed the lowest loss in lime fresh weight when compared with single treatment. The loss in weight reached the maximum after three weeks when SCB were used. TSS increased in all treatments compared to control as shown in (Table 6). Ascorbic acid (Vitamin C) was decreased in all treatments compared to the control in naturally infected lime fruits.

**Table 6: Effect of citral and sodium carbonate (S.C) or sodium bicarbonate (SBC) on loss of lime fruits weight, total soluble solids (TSS) and vitamin C (VC) after 3 weeks storage at  $13 \pm 1^\circ\text{C}$  under natural infection**

Treatments	Loss in weight %	TSS (%)	Vitamin C
<b>Single treatment</b>			
Citral 0.3%	3.0	8.5	45.0
Citral 0.5%	2.3	8.5	45.0
SC 2%	3.7	9.0	44.0
SBC 2%	4.0	9.0	43.0
<b>Combined Treatment</b>			
Citral 0.3% + SC 2%	2.7	9.2	45.0
Citral 0.5% + SC 2%	2.2	9.2	44.0
Citral 0.3% + SBC 2%	2.5	8.8	45.0
Citral 0.5% + SBC 2%	2.0	8.5	45.0
Control	6.6	7.5	46.0

## DISCUSSION

Postharvest green and blue molds caused by *Penicillium digitatum* and *P. italicum* are important postharvest storage problems of lime fruits (Smilanick and Arruf, 1992). Component of volatile oil i.e limonene, linalool, citral and nerol were extracted from orange peel, coriander fruits, lemon grass and petitgrain (Bitter orange) respectively, and used in this study. *In vitro* nerol and citral caused complete inhibition on linear growth of tested fungi at 0.4 and 0.5%. Linalool gave the lowest effect on linear growth of the two fungi. The differences between component of each other and the antimicrobial characteristics seemed to be correlated with their structure, solubility and capacity to penetrate cell wall and to interfere with the enzymatic reaction of the cytoplasmic membrane (Knobloch *et al.*, 1989). Sodium carbonate has completely inhibited the mycelial growth at concentrations 2.0% and 3.0%. These results are similar to those reported by Wassana (2004) who found that Sodium bicarbonate at 1% appear to inhibit growth of *P. digitatum* and their concentration at 2% and above could completely inhibit spore germination. Inhibition of microorganisms by salts might be caused by reducing of the cell turgor pressure with collapse and shrinkage of hypha and spore (Fallik *et al.*, 1997) or by alteration of cell transport function and inhibition of enzymes involved in the glycol improved lytic pathway (Sofos *et al.*, 1986).

Citral, sodium carbonate (SC) and sodium bicarbonate (SBC) were used in the present study against green and blue molds *in vivo*. Citral oil, SC and SBC were effective in controlling green and blue molds. The effectiveness is quiet good when applied as single, but improved when used in combination. Citral at 0.5% inhibited percentage disease severity of lime fruits under artificial infection condition with *P. digitatum* and *P. italicum* and percentage of infection under natural infection condition. Using citral for controlling postharvest disease of oranges as reported by Abd-El-kareem and Abd-Alla (2002). The mechanism of citral in reducing postharvest disease appears to be related to fungistatic property (Rodove *et al.*, 1985). Treatment of lime fruits with SC and SBC when applied as single treatment reduced fungal infection and delayed disease development under artificial infection during storage up to three weeks. Effective control of green and blue molds of lime fruits by combined treatment between citral at 0.5% and SC or SBC at 2.0%. High efficacy 100% was obtained at the same treatment. The inhibitory effect of SC or SBC depends on the presence of salt residues within the wound infection courts occupied by the fungus and on interactions between this residue and constituents of the rind. These interactions presumably alter the original *in vitro* toxicity of the salts to spores. All treatments lowered the loss in fresh weight of lime fruits during storage in naturally infected fruits compared to control. Total soluble solids and vitamin C decreased at all treatment compared to control. Sami *et al.* (2007) reported that the loss was decreased in fresh weight of lime fruit and increased in TSS under natural infection during storage at 13 °C for four weeks.

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تأثير بعض مكونات الزيوت الطيارة وكربونات أو بيكربونات الصوديوم على  
مقاومة العفن الأزرق والأخضر لثمار الليمون أثناء التخزين  
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تم إستخلاص بعض مكونات الزيوت الطيارة وهي السيترال والليمونين واللينالول  
والنيروول من البرتقال والكزبرة وحشيشة الليمون والنانج علي التوالي وتم إختبارهم علي النمو  
الميسليومي للفطرين بنسليوم إيتالكم وديجيتاتم اللذان يسببان العفن الأزرق والأخضر لثمار  
الليمون.

أوضحت الدراسة أن النيروول والسيترال عند ٠,٤ % أدوا إلي تثبيط النمو الميسليومي  
للفطرين تحت الدراسة بنسبة ١٠٠% وكربونات أو بيكربونات الصوديوم عند ٢% أدت إلي  
التثبيط الكامل لنمو الفطرين.

تمت دراسة تأثير السيترال وكربونات أو بيكربونات الصوديوم بصورة فردية أو مختلطة علي  
مقاومة العفن الأزرق والأخضر لثمار الليمون وأوضحت النتائج أن تكامل المعاملات بين السيترال  
بتركيز ٠,٥% وكربونات أو بيكربونات الصوديوم بتركيز ٢% أدوا إلي التثبيط الكامل لإصابة  
ثمار الليمون تحت ظروف العدوى الصناعية بالفطرين بنسليوم إيتالكم وديجيتاتم وأدت نفس  
المعاملات إلي منع الإصابة تماما تحت ظروف العدوى الطبيعية عند تخزينها عند درجة ١٣ م  
لمدة ثلاث أسابيع.

ويُتقدّر الجودة للثمار أثناء التخزين وجد أنه قد قل الوزن الطازج لثمار الليمون  
ومحتواها من فيتامين C بينما زادت نسبة المواد الصلبة الكلية.  
وتشير النتائج الي إمكانية استخدام تكامل المعاملات بين السيترال وكربونات أو بيكربونات  
الصوديوم لمقاومة أمراض ما بعد الحصاد لثمار الليمون.

قام بتحكيم البحث

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