UV-C Radiation Effect on the Mortality of Fruit Fly Eggs

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ABSTRACT

The current work evaluate death kinetics of *Ceratitis capitata* eggs immersed in water and subjected to different doses of UV-C radiation: 0 (Control); 0.087; 0.261; 0.348; 0.461; 0.692; 0.922; 1.384; 4.152; 5.534 and 8.302 kJ m⁻². After treatments, eggs placed in Petri dishes were incubated at 25°C under light/darkness or in complete darkness. One day after treatments, egg survival was microscopically evaluated. Doses of 1.384 kJm⁻² UV-C or higher led to 100% death of *C. capitata* eggs incubated in a light/darkness cycle.

Keywords: Mediterranean_fruit fly, postharvest, quarantine treatments, Brazil

1. INTRODUCTION

The Mediterranean fruit fly (Diptera - Tephritidae), *Ceratitis capitata* (Weid., 1824), is widespread throughout the five Continents, and it is considered a high invasive colonizing species due to its ability to adjust to diverse climate conditions and hosts, high reproductive performance and quick-spreading faculty (RAGA, 1996). Tephritidae family is one of the biggest within the Diptera order, with 500 genera and approximately 4.000 described species. This family is among those which heavily impacts fruit production worldwide, since it attacks the reproductive plants organs, flowers and fruits. This fact obviously results in economical damage to fruit producers around the world. Currently, about 1 billion dollars/year are lost due to direct damages on fruit production, trade and subsequent sanitary barriers in international trade (SILVA & BATISTA, 2009).

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Since it is a plague subjected to quarantine rules, there are commercial barriers imposed by importing countries, mainly those who are fruit fly-free zones, limiting fresh-fruits exports. Good and sustainable agricultural practices implies a rational and exclusive use of chemicals officially allowed for each fruit, obeying withdrawal periods, environment protection, analysis of residues and proper harvest and post-harvest handling practices (IBRAF, 2007).

Due to trade barriers and a shift in the world wide trend concerning safe and residuefree food production, plague control by physical alternatives instead of agrochemicals applications has been widely investigated during the past decade, since it is safer for the environment and human beings (VIEIRA, 2004; ARRUDA et al., 2004). Several physical techniques like application of heat, cold, ionizing and non ionizing (UV-C) radiation, in association with modified atmospheres, maturity inhibitors and refrigerated storage have been developed and improved as postharvest treatments in order to support quality requirements of fruit and vegetables importing countries (ALLENDE and ARTÉS, 2003; VICENTE et al., 2006; PINHEIRO et al., 2005).

The UV-C radiation is an effective technique for mold disinfection and fruit and vegetables preservation, due to its germicide properties - basically consisting in destroying microbial DNA and protein denaturation - leading to a longer shelf life of fresh intact as well as minimally processed products (AGUAYO et al., 2007). It also has the great advantage of not producing any unwanted co-products, adverse for the environment and the human consumption. Moreover, it does not stimulate the synthesis of unwanted byproducts that might change sensorial characteristics (flavor, odor, and color) of final products (GUERRERO-BELTRÁN et al., 2004). However, little is known about its effects on insects for quarantine purposes.

The UV-C light has been widely tested as an alternative to chlorine for disinfection of minimally processed fruit and vegetables (ARTES et al., 2009). The UV-C light may also be used to inactivate several kinds of dangerous organisms in processed food, increasing its shelf life, and also in pharmaceuticals, electronics, and to disinfect and purify drinking water. However, no references about the use of UV light to destroy fruit flies or any other quarantine plagues have been found. Thus, the objective of this work was to evaluate the effects of several increasing UV-C light doses on *C. capitata* eggs mortality.

2. METHODOLOGY

This experiment was performed at the Pilot Plant for Food Engineering of the Technical University of Cartagena, Murcia (Spain). The *C. capitata* eggs were obtained from mass rearing already carried out at the Instituto Murciano de Investigación y Desarrollo Agrario y Alimentario (IMIDA, Murcia, Spain).

The UV-C equipment consisted of two batches of 15 reflectors with unfiltered germicidal emitting lamps (TUV 36W/G36 T8, Philips, 139 Holland) fixed to a chamber frame. One batch was horizontally suspended on the top of the radiation chamber and the other one was placed below it. The Petri dishes were placed between both lines of UV-C lamps (15 cm of distance) over a polystyrene net. Walls of the experimental chamber were protected with a reflecting inner layer, which enhanced homogeneous distribution of the emitted light and allowed indirect illumination of practically the whole dishes. In order to determine the UV-C radiation intensity of the

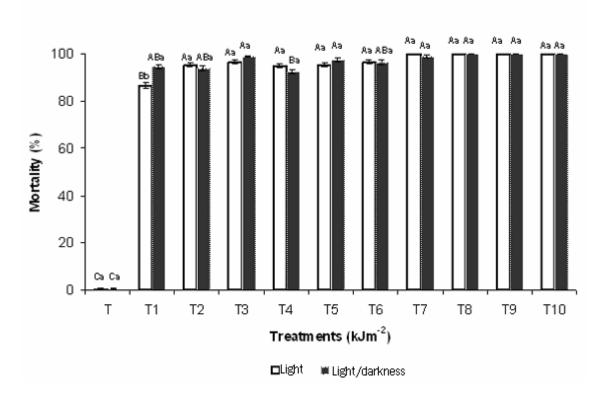
lamps, a VLX 254 radiometer (Vilber 146 Lourmat, Marne la Vallée, France) was used (LOPEZ-RUBIRA et al., 2005). The applied UV-C intensity was calculated from a mean of 18 readings taken at each side of the net. Light intensity was kept constant, and the applied doses varied by modifying the exposure time (ARTES-HERNANDEZ et al., 2009). Eggs were exposed to the following UV-C radiation treatments 0 (control); 0.087; 0.261; 0.348; 0.461; 0.692; 0.922; 1.384; 4.152; 5.534 and 8.302 kJ m⁻². Twenty replicates were evaluated for each treatment, each one containing twenty eggs, selected with an electronic microscope (Olympus SZ40, Japan), with maximum 48 hours of age, being placed in Petri dishes and immersed in 20 ml of distilled water. After each UV-C treatment, half of the Petri dishes (20) were stored in presence of light (12 h light, 600 lux m⁻²-12 h darkness) while the other half (20) where stored in total absence of light (24 h darkness), at 25 °C. Twenty-four hours after treated, samples were evaluated for egg mortality, using a stereoscope microscope (Olympus SZ61, Japan) provided with a photographic camera (Olympus Altra 20, Japan).

The experiment followed a complete randomized factorial design. Data were submitted to analysis of variance (ANOVA), at 5 % significance level (P = 0.05). Means were contrasted using the Tukey's test. The analysis was performed using the statistical software Infostat version 1.0.

3. RESULTS AND DISCUSSION

As main result the application of UV-C light was lethal for the fly fruit eggs, the higher the doses, the higher the level of dead eggs (Fig. 1). In fact, significant higher egg viability in control (T1) than in the rest of treatments was found. All control eggs and some of the lower than 1.384 kJ m⁻² treated ones hatched and became larvae of first stage after 24 h (Fig. 2-B). From the best of our knowledge, results on the effect of UV-C non-ionizing radiation on fruit fly eggs mortality are firstly reported here. For comparison, when the ionizing gamma radiation was applied the increasing destructive effect of increased radiation levels has been previously found (WALDER, 1993; ARTHUR & WIENDL, 1994; RAGA, 1996).

Besides the effect of UV-C dose, an interaction between radiation doses and storage conditions (light or darkness) on the mortality of *C. capitata* eggs was found (Table 1). For control fly eggs, a period of darkness increased their mortality. However, the effect of light was not significant when UV-C light was applied, and eggs died independently of the light conditions.



RADIATION UV-C EFFECT IN THE MORTALITY OF THE FRUIT FLY EGGS

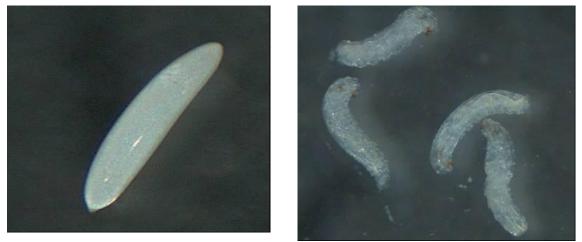
Figure. 1. *Ceratitis capitata* egg mortality when submitted to different UV-C radiation doses and distinct light conditions after 24 h storage. T: 0 (control); T1: 0.087; T2: 0.261; T3: 0.348; T4: 0.461; T5: 0.692; T6: 0.922; T7: 1.384; T8: 4.152; T9: 5.534 and T10: 8.302 kJ m⁻². Values followed by different letter were different significantly by Tukey's test at 5 % significance level (P = 0.05), Capital letters compare radiation levels and low-case letters indicate light condition.

Table 1. Analysis of variance of *C. capitata* eggs mortality subjected to different UV-C treatments and stored during 24 h at 25°C with or without light.

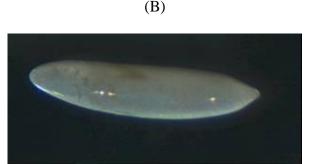
Source	SS	df	MS	F	р
MODEL	6852.48	21	326.31	668.14	< 0.0001
UV-C.	6835.73	10	683.57	1399.66	< 0.0001
LIGHT	1.02	1	1.02	2.09	0.1494
UV-C x LIGHT	15.73	10	1.57	3.22	0.0007
ERROR	96.70	198	0.49	-	
TOTAL	6949.18	219	-	-	

SS – Square sum, df - degrees of freedom, MS - Mean square, F - F value, p - significance level.

Egg mortality by 1.384 kJ m⁻² UV-C (Fig. 2) or higher, followed preferably by 24 h storage under light was fully effective, killing 100% of eggs and reaching the level (99.9968%) required by legal regulations of most importing countries free of fruit flies (MENDONÇA et al., 2000). Based on these findings it could be hypothesized that *C. capitata* eggs are not able to support degenerative damage caused by this UV-C radiation level. Below that dose, there is also high egg mortality, but some of the eggs are able to survive.







(C)
(D)
Figure 2. A - healthy eggs (before UV-C treatments); B - emerged larvae from T1 (control); C and D - dead eggs after 24 h of 1.384 kJm⁻² UV-C radiation in the light storage.

4. CONCLUSIONS

The UV-C radiation at 1.384 kJm⁻² or higher was a very efficient treatment to destroy *Ceratitis capitata* eggs incubated in an aqueous media. The use of UV-C light could be an economically viable promising technique for avoiding at 100% *C capitata* viability, and very probably for other fruit flies. When compared to other quarantine treatments UV-C light could be considered as more efficient and environmentally safer. Further studies should be conducted in order to evaluate mortality of the fly eggs in infested fruits, for demonstrating the *in vivo* efficacy of UV-C radiation at low levels as a quarantine treatment.

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