

The Olive Leaf Moth *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae) as a Serious Pest in the World: a Review

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Abstract: The olive leaf moth *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae) is an international lepidopteran pest originating in the Mediterranean Basin. The present article was prepared aiming to review several aspects of the distribution, economic importance, morphology, biology, ecology and physiology of this pest. Beside the worldwide distribution, economic losses and morphologic characterization, efforts of insect biologists in several parts of the world had been comprehensively highlighted including developmental biology and reproductive biology. From the ecological point of view, population dynamics, sexual and non-sexual behavioral patterns had been discussed. Also, special attention was paid to the laboratory rearing trials on artificial diets. It may be the first review focusing on these aspects of *P. unionalis* in the world. On the other hand, this pest still needs considerable research work for investigating several aspects such as energy metabolism, homeostasis, enzymatic patterns, hematology, resistance, immunity, reproductive physiology, environmental physiology and the hormonal regulation of these processes. Therefore, the present review enhances the research interests for these important aspects. However, information reviewed herein will support the development of strategies for management of this pest.

Keywords: behaviour, biology, development, distribution, ecology, embryonic, morphology, physiology, population, reproduction.

1. INTRODUCTION

Olive (*Olea europaea* L.) is one of the first fruit trees cultivated by man. It is small evergreen tree in the family Oleaceae, native to the coastal areas of the eastern Mediterranean Region, from Lebanon and the maritime parts of Asia Minor to northern Iran at the south end of the Caspian Sea [1]. Olive is one of the economically important crops in the Mediterranean Basin. Nowadays it is grown in North America, South Africa, China, Japan and Australia [2], although it is considered that about 98% of the world's olive production is located in the Mediterranean area [3, 4].

Olive tree is subjected to attack by several insect pests causing considerable yield losses in quality and quantity. These pests belong to Diptera, Lepidoptera, Hemiptera, Orthoptera, Coleoptera, and Thysanoptera [5]. The most common pests observed in Egypt, as for example, include: *Bactrocera oleae* (Diptera: Tephritidae), *Prays oleae* (Lepidoptera: Yponomeutidae), *Palpita unionalis* (Lepidoptera: Pyralidae), *Zeuzera pyrina* (Lepidoptera: Cossidae), *Saissetia oleae* (Homoptera: Coccidae) and *Parlatoria oleae* (Homoptera: Diaspididae) [6, 7]. However, the olive moth *P. oleae* and olive leaf moth *P. unionalis* are two well known lepidopterous pest species of olives in Egypt and some of other Mediterranean countries [8-10].

The olive leaf moth has several vernacular names, such as olive leaf caterpillar, olive leaf worm, olive buds moth, Jasmine moth, Jasmine bud worm, Jasmine moth and jasmine leaf caterpillar. Its scientific name is *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae) with synonyms *Palpita vitrealis* (Rossi) [11], *Margaronia unionalis* (Hübner) [12] and *Pyrallis unionalis* (Hübner) but Kirti and Rose [13] prepared an identification key for different species of *P. unionalis*, based on the internal and external differences of male genitalia in India, and suggested that this species belongs to the genus *Palpita*.

To develop efficient control measures against a pest and improve Integrated Pest Management (IPM) strategies, knowledge of its agronomic impacts, biology, ecology and other necessary information should be available. Therefore the objective of the present article was to review several aspects of distribution, losses, morphology, biology, ecology and physiology of the economically serious pest, *P. unionalis*.

2. WORLDWIDE DISTRIBUTION OF *P. unionalis*

From the Zoogeographical point of view, the Mediterranean Basin was reported as the original area of *P. unionalis*, where it is found from east to west, and south to the olive-growing regions of northern Africa, the Canary Islands and Madeira. Now it is an international lepidopterous migratory pest in the tropical and mildly subtropical regions of the Old World [14-16]. As reported in the literature, it is distributed in Turkey [10, 17, 18], Greece [19], Slovakia [20], Portugal [21], Israel [22], Italy [23-26], India [13] and Egypt [27]. Different dispersal regions of *P. unionalis* were reported such as Spain, Japan, Australia, North Africa, Tropic Regions of America and South America [28] like Brazil [29]. It is a highly mobile moth dispersing even to northern Europe, such as Sweden and Poland [30-32]. Also, it has been reported to attack wild olives in Southern Africa [33, 34] and jasmine *Jasminum fluminense* in Kenya [35].

For some details, *P. unionalis* was firstly recorded in 1969 in the Aegean region and reported as one of the main pests of olive in Marmara region, Turkey [17, 36, 37]. It was a secondary pest of olive cultivars but now it is considered as a primary pest in olive nurseries, irrigated young olive plantations as well as in mature olive trees [10, 38]. Within a few years later, the pest became an epidemic in olive nurseries throughout the country [39]. In Syria, *P. unionalis* was observed in the last few years in Aleppo and identified as olive buds moth or Jasmine moth [40, 41]. Within a short period, the pest became epidemic in olive nurseries throughout the country [42]. In Malta, *P. unionalis* was reported for the first time to be associated with olive trees [43]. After few years later, its local occurrence was described as an immigrant pest feeding on olive and jasmine [44-46]. Depending on a survey carried out by Haber and Mifsud [1], the moth was recorded during most of the year in many localities of the Maltese Islands. In Egypt, this pest has been known as jasmine moth or olive leaf moth. Rahhal [47] carried out a survey during the first two years of seventh decade of 20th century. Infested samples of olive branches were obtained from different orchards and farms in Alexandria, Mersa Matrouh, Bourq El-Arab, Siewa Oasis, Kaliobia, Cairo, Giza and Fayoum. It is reported as a destructive pest of young olive farms [6, 48] and the old trees [49, 50]. In addition to Mediterranean region, *P. unionalis* was recorded as a serious pest in some Asian countries such as Iran in which the pest was firstly reported in olive orchards of Roudbar City in August 1999 [51]. It took only a few years to spread as a serious pest throughout the country [52-54].

3. HOST PLANTS AND ECONOMIC LOSSES OF *P. unionalis*

P. unionalis is a polyphagous lepidopteran attacking the family Oleaceae, especially the genera of *Ligustrum*, *Oleae*, *Fraxinus*, and *Phyllyrea* [15, 55]. However, a range of host plants had been reported in different parts of the world. In Turkey, *Fragaria ananassa* (Rosaceae) and *Viburnum* spp. (Adoxaceae) are also reported as alternative hosts for this pest [56]. In Greece, the pest has also been reported on a range of other plant hosts including, *Arbutus unedo* (Ericaceae)[57]. In Iran, *Ligustrum vulgare* was found as a desirable host for breeding of the pest in laboratory in comparison with olive cultivars [58]. In Egypt, *P. unionalis* is a pest of olive, as the main host, but it attacks also the jasmine *Jasminum officinale*, *L. vulgare*, *A. unedo* and *Phyllyrea media*, particularly in the new reclaimed lands [6, 49]. Thus, olive is the main host for this pest but Jasmine, *Fraxinus ananassa*, *Viburnum* spp. and *A. unedo* can be considered as alternative hosts [56, 57].

With regard to the economic losses, *P. unionalis* attacks its host plants in different countries causing mild or severe damage, depending on its population density and the host plant. Different losses had been reported in Greece [19], Italy [23, 24, 26], Israel [22] and Egypt [27, 59]. The most important damage of the pest occurs on young trees, nurseries and shoots of old trees [60, 61].

In some detail, *P. unionalis* is considered in Greece a serious pest feeding on young leaves and shoots of *Jasminum* sp. and *Ligustrum* sp. [55, 57]. Young larvae consume entire leaves and buds in the first generation and in second generation, they feed on fruits and seeds if they reach to high population levels [61]. In France, ornamentals, such as jasmine cultivated for perfume production, suffer from larval attacks of *P. unionalis* on leaves and flower buds [62, 63]. In Spain, both leaves and fruits of olive are damaged by *P. unionalis*. In nurseries or young orchards, feeding damage by larvae can reach up to 90% of the leaf area, thereby seriously affecting the development of the plant shoots. During the fruit ripening season, high larvae infestations may also reduce the fruit yield by 30% [64]. In years of high population densities, larvae attack also olive fruits, making them unsuitable for marketing [26, 57]. During heavy *P. unionalis* infestations in Italy, the most important damage occurs

on young trees, nurseries and shoots of old trees [23, 60]. In Sicily, if 90% of olive branches have been damaged, loss rate of yield will not be more than 20% [24]. In Malta, the most damage was observed on the new growth of olive trees, where not only new leaves and buds were eaten but occasionally entire shoots up to 15 cm were completely eroded [1]. In Turkey, *P. unionalis*, was a secondary pest of olive cultivars but now it is considered as a primary pest in olive nurseries or irrigated young olive plantations causing important economical yield losses by fruit fall as well as by damage on leaves, flowers and fruits [10, 17, 36, 38, 65]. Quality of the product may be impaired by larval feeding on all phenological stages of the olives [39]. In Egypt, *P. unionalis* was considered as a destructive pest of young olive groves [63, 66] and new branches of old trees [49]. Its damage increases in the olive groves, particularly in the new reclaimed lands [6]. Significant direct yield loss in olive plantations can be recorded due to fruit fall [67, 68] and destruction of a large part of the crop can be caused by the highest population density of *P. unionalis* [50]. In South Africa, *P. unionalis* seemed to be of negligible potential economic importance to olives in the Eastern Cape [69]. Thus, *P. unionalis* has been considered as a secondary or minor pest in some countries but as a primary, serious or even key pest in others, depending on the attacked plant species, seasonal climatic conditions of the region, densities of the endemic natural enemies, etc. In a similar course, it can be recorded as a minor pest in a country and as a serious one later, depending on the previously mentioned factors.

4. MORPHOLOGICAL CHARACTERIZATION OF *P. unionalis*

Knowledge of insect morphology is essential for the identification of a species without which all efforts exerted in biology, ecology, physiology and pest control can be wasted. Several studies had been carried out on the descriptive morphology of *P. unionalis* in different parts of the world. We have compiled here some of reported works, including the morphological appearance and diagnostic characteristics of all developmental stages (*viz.* adults, eggs, larvae and pupae).

4.1. Adult Moths

Adults are characterized mainly by shiny semitransparent or white wings with a brown leading edge of the forewing and two black spots in the middle. Kacar and Ulusoy [70] shed some light on the morphological characteristics of this insect in Turkey. Depending on this study, body length was 13.9 ± 0.17 mm for females while 13.90 ± 0.18 mm for males. The wings bear frenulum, and in resting position stand on the body in gable roof form. The wings appear, also, with two black spots in the middle. It is interesting to refer that front wings are wider than the hind ones. Wing span was in average of 28.93 ± 0.30 mm and 28.27 ± 0.30 mm of females and males, respectively. Dissimilarly, Yilmaz and Genc [10] measured the wingspan in an average of 22.6 ± 5.1 mm and 25.0 ± 3.3 mm for females and males, respectively. The males and females do not differ in length and width albeit females possess a mating pore on the 8th segment of abdomen and an oviposition pore on the 9th one. As seen by naked eye, female moth can be discriminated by her light green abdomen covered with white scales, but abdomen in males has terminal part bearing sets of hair. It should be mentioned that the internal reproductive system in adult male and female was described by Santorini and Vessiliana-Alexopoulou [12]. Also, special attention had been paid to the morphology of antennal sensory receptors as principal organs of intra-specific pheromone-steered communication [9, 71].

4.2. Egg

Depending on the study of Kacar and Ulusoy [70], in Turkey, averages of the egg size and width were determined as 0.95 ± 0.011 mm and 0.72 ± 0.008 mm, respectively. Eggs are white, flattened, with reticulated appearance and 0.5-1.0 mm in length [56]. As described by Yilmaz and Genc [10], eggs were elongated, flattened, and about 0.80 ± 0.10 mm in length, 0.5 ± 0.07 mm in diameter, weigh about 0.1 mg. According to Noori and Shirazi [16], in Iran, eggs are flat oval, light greenish yellow, 1.02 mm long and 0.49 mm wide, exhibiting a mesh appearance.

4.3. Caterpillars

Larvae are pale yellow in 1st and 2nd instars, later becoming gradually green with shading bluish toward the head and the tail. Maximum body length of mature larva ranged from 18 to 20 mm [56] or the average of 22.20 ± 0.15 mm [70]. A detailed study on the larval morphology was carried out in Turkey, also. On the basis of this study, size of the head capsule was 0.18 ± 0.03 mm for the first

instar and up to 1.53 ± 0.04 mm for sixth instar. Different characteristics, such as colour, weight, long, wide, and some body structures had been described for all 6 instars [10]. According to the observation of Noori and Shirazi [16], in Iran, larva is eruciform with three thoracic and five abdominal legs on the 3, 4, 5, 6, and 10th abdominal segments. Crochets are seen in closed and complete spherical form at the end of prolegs. Sometimes a pair of black spots are seen on the body segments close to the pleural part at 3rd and 4th instar instars.

4.4. Pupae

As pointed out by some authors [10, 56, 70], pupae are initially soft and light green in color. Their color turned to brown on the following day. Female pupae measured about 3.07 ± 0.23 mm wide and 14.01 ± 0.90 mm long and weighed 73.61 ± 11.77 mg. Male pupae were about 13.38 ± 0.80 mm long, 2.98 ± 0.21 mm wide and weighed 70.6 ± 12.78 mg. The sex differentiation depending on immature stages was, also, provided [65].

5. DEVELOPMENTAL BIOLOGY OF *P. unionalis*

It is important to know the basic biology of a pest so as to understand factors involved in population fluctuations which is necessary in planning an IPM programme to control this pest. The available literature has been enriched with many reported works on the biological characters of *P. unionalis* [9, 10, 16, 56, 58, 72]. Some of these works focused on its life table on different hosts [18], population dynamics and some of the environmental factors affecting it [17] and the influence of temperature on embryonic development [73]. On the contrary, very few reports on the rearing techniques of *P. unionalis* had been available. Herein, we reviewed both the embryonic and post-embryonic development of this insect.

5.1. Embryonic Development

The egg formation and development in insects had been studied by some researchers [74-76]. The egg shell, or chorion, of an insect is a complex of several layers. It is synthesized within the ovarioles by the follicular epithelium that surrounds the oocytes and begins once vitellogenesis takes place, that is, the uptake of vitellogenins [77, 78]. Following the union of the sperm from male and the egg in female, the newly formed zygote undergoes cleavage within the patterned environment that is present in the egg [79]. Further information about oogenesis and vitellogenesis through the successive stages of embryogenesis can be provided by some authors [80-83].

5.1.1. Embryonic Developmental Rate

Incubation period of the insect egg is usually the interval elapsed between the time of laying a fertilized egg and its hatching. This period may be used as a good indicator of the embryonic developmental rate, i.e., the shorter period indicates a fast rate and *vice versa*. Mean duration of the embryonic development of *P. unionalis* varied depending on the season, in the field studies, or constant rearing temperature in the laboratory studies. It was reported as a range of 15-16 days [23] but as 3 days in summer and about 9 days during winter [27, 47] or a range of 3 days (at 30°C) and 12 days (at 15°C) [84]. Loi [73] evaluated the effect of different temperatures, in a range of 10-35°C on *P. unionalis* embryonic growth and development. The fastest embryonic developmental rate was denoted by the shortest incubation period (3 days) at 30°C and the lowest one was denoted by the longest period (112 days) at 15 °C. On the basis of a study carried out on olive plants under the natural conditions in Adana (Turkey), the embryonic duration was recorded in a mean of 3.45 ± 0.13 (3-5) days and 4.33 ± 0.10 (4-5) days [85]. In Turkey, also, Yilmaz and Genc [10] estimated the mean duration in 4.16 ± 0.09 days at 24 ± 1 °C. Dissimilarly, Noori and Shirazi [16] found the mean embryonic developmental duration as 5.8 ± 1 days. In a detailed study, Yilmaz and Genc [10] followed up the successive developmental stages of the embryo day by day until the formation of mandibles and eyes which could be seen through the chorion just before hatching. Depending on the available literature, this was the first detailed study on embryonic development of this pest all over the world. In conclusion, the embryonic stage of *P. unionalis* has been a temperature-dependent.

5.1.2. Embryonic Survival

It may be reasonable to know the viability or ability of embryos to live as detected by the hatching percentage (hatchability). Although the embryonic survival has been affected by several factors, majority of entomologists paid their attention, until now, to the temperature only. Earlier, the

hatchability of *P. unionalis* was determined as 84% [22], 78-95% in winter [47] or 68 % at 15^oC and 98% at 25 °C [73]. In contrast, the effect of relative humidity (R.H.) seemed to be negligible. In a study, selected diets were provided to adults aiming to investigate the effect of adult diets on different reproductive parameters. Egg hatchability was not affected by adult feeding [38].

5.2. Post-Embryonic Development

Upon hatching from eggs, juvenile insects embark on an excursion of post-embryonic development that will eventually take them to their adult forms. The change that occurs when an insect develops from an immature stage to an adult stage is called “metamorphosis”, literally meaning “change in form”. Insects show three major metamorphic strategies for reaching the adult stage, with the degree of metamorphosis dependent on the degree of divergence between the immature and adults [86]. In the holometabolous insects, like *P. unionalis*, newly hatched larvae feed and grow increasing to critical size and weight, then they moult to change new integument. Moulting has been repeated several times and the mature larvae prepare themselves to pupate. The pupae live certain period, during which the larval structures were destructed (histogenesis) and the adult structures will be constructed (histogenesis). At the end of pupal stage, adult moths can emerge. All vital processes, from the first point until the end have been undergone to hormonal regulation (For details, see: [79, 87-93]). Shortly, this is the dramatic journey of post-embryonic development during the life of *P. unionalis*. However, several aspects are reviewed herein.

5.2.1. Number of Larval Instars

Extensive studies had been conducted on the biology of *P. unionalis* in the laboratory showing the appearance of 5 instars in the larval stage [16, 54, 56, 61]. On the other hand, some investigators reported 6 instars [15, 27, 47, 72, 84]. No author obtained his observation depending on the head capsule measurements. An extensive study on the biology of this pest was carried out in Turkey revealing 6 larval instars in the larval stage, basing on direct quantification of molts and measurements of the head capsule [10].

5.2.2. Larval Development

Larval development can be indicated by the larval duration, i.e. shorter larval duration denotes faster developmental rate. The larval duration, and subsequently the larval development, of *P. unionalis* can be chronologically reviewed as follows. It was determined as 18-48 days [94], 14-19 days [22], ranged between 15.85 and 23.15 days [47], 21-26 days (at 25°C and 65% RH)[19], averaged about 15 days in summer and 23 days in winter [27], 24-30 days [24], 22.28±0.22 days [54], 25 days [95], 21.6±0.3 days [16], as well as 18.50±0.56 days in the first generation and 26.25±0.82 days in the second generation [85]. However, these differences can be due to the pre-pupation (prepupae) period since El-Kifl et al. [27] determined a pre-pupation period in about 1-1.6 days in summer and 2.5 days in winter. In the field, larvae feed on the leaves at the end of the twigs, forming silken webs in which they sheltered and pupated later [54]. In the laboratory, full grown larvae were observed to fold one or more leaves together with white silken webs, inside which pupation took place. Prepupae remained about 1.81±0.40 days and changed within 4-5 minutes into the characteristic pupal appearance [10]. The average prepupal duration was 1.63±0.18 days in the first generation and 1.73±0.35 days in the second generation [85]. Thus, the reported differences in larval duration may be understood because some authors considered the prepupal period within the larval duration but others considered it as a separate phase. Furthermore, durations of the successive larval instars had been determined separately. As for example, Badawi et al. [84] measured the duration of the last (6th) instar as almost double the duration of the first one. Yilmaz and Genc [10] recorded the durations of larval instars, in detail, as follows. First instar: 2.93±0.73 days, second instar: 3.42±1.55 days, third instar: 3.42±1.15 days, fourth instar: 3.00±0.96 days, fifth instar: 3.57±1.28 days and sixth instar: 7.00 ±1.56 days. It should be mentioned that the larval development has been usually affected by the ambient temperature (For some details, see: [9, 16, 54, 73, 85, 95]).

5.2.3. Pupation and Pupal Development

As reported by many authors [16, 58, 85, 96, 97], pupation took place in a silken cocoon under dry fallen leaves, under tree bark or in crevices on the stem. According to Rahhal [47], the pupal period ranged from 8.55 days during summer to 17.06 days during winter. El-Kifl et al. [27] and Noori and

Shirazi [16] determined longer period (9 days during summer and 17-18 days during the winter). Yilmaz [95] studied some biological parameters of this pest in Turkey and determined the pupal duration as about 10 days. In contrast, pupae lasted 7.83 ± 0.112 days under the field and laboratory conditions in Iran [54]. In conclusion, the duration of the pupal stage was much affected by the climatic conditions, especially temperature and relative humidity (R.H.) [72].

5.2.4. Mortality Rate of Immature Stages

As pointed out by many authors [10, 56, 73, 84, 95], the mortality of immature stages was 100% at 10-35°C and less than 50% at 13-30°C. Also, the survival rate was calculated in 60% and 80% for mature larvae and pupae, respectively. In a study, the survival rate of larvae was recorded as 61.6% by rearing in the laboratory on olive leaves (at 24 ± 1 °C, 65% RH and 16:8 h L:D). In addition, the egg stage was the most susceptible stage, whilst the susceptibility of the larvae was decreased as they grew up more so no mortality could be observed amongst the fifth instar larvae. The mortality of eggs depended on the temperature but different host plants, also, affect the insect survival [98-101].

5.2.5. Life Span from Egg to Adult (Total Developmental Period)

Total developmental period for *P. unionalis* was estimated in 26 days (by rearing on olive, at 26°C) but in a range of 21-26 days (at 23.4°C) [19]. It was measured in a range of 29.2-32.0 days by rearing on different host plants including olive, jasmine and ash tree [24, 84, 102]. Fodale and Mule [94] determined a life span of 29-38 days in field but from 24 to 39 days in laboratory. El-Khawas [103] measured the duration of pre-imaginal development (reared on young olive shoots at 27 °C and 65% RH) ranging from 21 to 30 days. Kumral et al. [18] recorded the total developmental period ranging from 27.52 days to 30.00 days. Total developmental period on Zard olive cultivar (at 25 ± 0.5 °C, 65 ± 5 % RH and 12:12 L:D) was found to be ~32 days [16]. By rearing on Gemlik variety of olive plants (at 28.2 °C, 70.1% R.H), average life span was found about 38.4 days or 61.6 days (at 18.4 °C, 64.4% R.H.) [85]. Rearing on Ayvacik olive cultivar resulted in a period of 24 days [10]. As shown by these reported results, the total developmental period of *P. unionalis* is temperature-dependent. In addition, different host plants are known to affect the insect development [98-100]. Moreover, the differences of the mean total developmental period could be due to the environmental conditions under which the experiments performed and the pest biotype [9, 26, 84, 103].

5.2.6. Number of Generations per Year

It is important to point out that *P. unionalis* is a multivoltine species with several overlapping generations per year, ranging from 1 to 10, until now. As reported by Grossley [61], the pest has 2-3 generations in cold to mild regions while more than 5-6 in mid-tropical and tropical regions. It has 6 generations per year in Israel [15, 24, 104, 105], 5 in Spain [24], 1-2 in France [14], 4-5 in Italy [24, 94]. Moreover, this pest has varied number of generations annually in the same country, depending on the differences in environmental conditions. As for example, it has 2 complete generations and 1 partial generation every year [17] but 9 generations at constantly stable [56] in Turkey. In Iran, it completed 4-5 generations [16], 6 generations [72] or 8 generations [54]. In Egypt, it has 10 overlapping generations a year [27, 47, 106] or 9 generations in various regions of different ambient temperature and R.H. (for detail, see [48, 84]). Therefore, the pest has varying number of generations annually depending on the host plant, seasonal temperature and other environmental conditions of over its universal or regional distribution.

5.3. Adult Performance

To shed some light on the most important parameters of its adult performance (emergence, sex ratio, survival and longevity), the available literature can be reviewed herein.

5.3.1. Adult Emergence

When ready to emerge, the moth pushes its head against the pupal skin causing a median dorsal slit which extends longitudinally. Through this slit, the moth finds its way out. The highest% of emergence (about 90%) and the least% of deformities (below 2%) were obtained at 25 and 30 °C. At 15 °C, nearly 35% of the moths failed to emerge. This vital process increased with the increasing R.H. but the most favourable R.H. was found 65%, at which 90% of normal moths emerged [27, 72, 84, 85, 96].

5.3.2. Sex Ratio

Sex ratio of *P. unionalis* is 1:1. in all generations during the year but males tended to be slightly higher during the later generations [10, 16, 19, 47]. In contrast, some field and laboratory studies revealed lower males than females, such as 1:1.16 [24], 1: 1.12 [65] and 1:1.14 [54].

5.3.3. Adult survival

By rearing on leaves of olive, its natural host plant, in the laboratory (24±1 °C, 65% RH and 16:8 h L:D), the survival rate of emerged adults was estimated in 86.9 % in males and 82.8% in females [10]. Selected adult diets had been assessed on the adult performance. Female survival was better among those fed on honey solution, honeybee liquid food, Gatorade and water than males. The adult survival also can be supported by feeding on both flowering plants and honey solutions in the laboratory [38].

5.3.4. Adult longevity

Reported results in different parts of the world revealed variation of the total adult longevity of *P. unionalis*. It was measured in females and males, respectively, as follows: 13.5 days and 15.3 days [73], 9.92-11.4 days and 9.00-11.9 days [9], 9.92-11.64 and 9.00-10.57 days [18], 12.3 days and 14.1 days [56], 14 days and 13.6 days [72], 12.59±1.63 and 14.33±2.4 days [54], 16.0±1.57 days and 16.3±1.21 days [10] and 12.6 and 13.5 days [16]. This variation can be attributed to the host plant, or even its variety, as well as the climatic conditions under which the experiments had been conducted.

Considering the compartments of adult longevity, *P. unionalis* adults mate 2-3 days [27] or 2 days after emergence (pre-oviposition period) and females die immediately after egg laying with no post-oviposition period [65]. On the contrary, three main compartments, (*viz.*, preoviposition period, oviposition period and post-oviposition period) had been recorded under controlled laboratory conditions and feeding on the olive leaves [10]. The pre-oviposition period was recently estimated in 2.3±0.3 days [10] which agreed with earlier result of 2-4 days [47] but disagreed with Shehata et al. [9] who reported a shorter period (1.7 days). Shorter period was also estimated (1-2 days after emergence) by Noori and Shirazi [16]. With regard to the oviposition period (Reproductive life-time), Rahhal [47] reported 4.2 and 8.5 days during summer and winter months, respectively. Longer period (10.5 days, [9] or 8.0±0.7 days, [10]) was reported but shorter period (5.60-9.15 days, [18] or 3-7 days, [16]) was recorded. The unique study determining a postoviposition period for this insect was conducted by Yilmaz and Genc [10] who calculated its mean in 2.4±0.4 days.

In respect of the mating effect on longevity, unmated males tended to live statistically insignificant longer than mated ones [18]. To a great extent, similar result had been earlier reported by Badawi et al. [84]. As pointed out by some authors [18, 84, 38], the adult longevity was much affected by feeding. However, some characteristics such as leaf morphology, chemical composition of the host plant or other interactions were not examined [9, 19].

5.4. Laboratory Rearing on Artificial Diet

The successful insect culturing in the laboratory is necessary for efficient and productive research on virtually every aspect of insect biology [107]. Rearing of *P. unionalis* mainly depends on natural host plants, such as olive leaves. Availability of host leaves, transferring larvae from old leaves to fresh young leaves, and susceptibility of larvae to pathogen infections are important issues to be considered. Such rearing is excessively cost in time and labor [108-110]. Therefore, mass-reared insects tend to be provided with artificial diets that bear little resemblance to their natural host or food source but nonetheless permit satisfactory growth and development of the mass-reared insects [111]. A laboratory rearing method for *P. unionalis* on artificial diet needs to be developed to facilitate the studies of different aspects and responses of this pest which are necessary requirements before planning of pest management strategies.

Since the first attempt by Bottger [112] to rear a phytophagous insect, *Ostrinia nubilalis*, on an artificial diet, a number of insects have been reared on artificial diets [113-117]. As easily appeared in the literature, Çiğdem Yilmaz, singly or with her colleagues, conducted some trials for development an appropriate artificial diet for *P. unionalis*, under the laboratory conditions, in Turkey. Yilmaz [95] compared some biological parameters of this pest after feeding the natural host plant and artificial diets and observed no differences. Almost, similar results were obtained by Sahin and Genc [39]. Along two generations of *P. unionalis* in the laboratory, Yilmaz and Genç [97] assessed the effects of

feeding on different artificial diets on some biological parameters and maintenance of a colony on an artificial diet. They used the artificial diet developed for rearing *Spodoptera* spp. [118] and *Phyciodes phaon* [119] basing on pinto bean, wheat germ and torula yeast which have been previously used to rear *Spodoptera* spp. [118]. It was concluded that this diet seemed to be the most adequate diet to rear *P. unionalis*. On the other hand, there are many factors that affect the larval feeding on artificial diet, such as proportion and balance of nutrients, moisture level and texture of diet [120]. Also, addition of some host plant materials in artificial diets often promotes growth, survival and fecundity, and may act as necessary stimulants for oviposition and successful rearing [119]. However, further studies should be conducted to determine the possibility of rearing larvae on the artificial diet for successive generations.

6. REPRODUCTIVE BIOLOGY OF *P. unionalis*

6.1. Net Reproductive Rate

The net reproductive rate is considered as a parameter of life table (life table will be discussed thereafter in the present review). It is a key statistic that summarizes the physiological capability of an animal relative to its reproductive capacity. Comparison of net reproductive rate often provides considerable insight beyond that available from the independent analysis of individual life history parameters [100]. It is an important indicator of population dynamics [121, 122]. Also, the net reproductive rate may reflect the potential of host plants to contribute to *P. unionalis* populations. It varied among different host plants. For example, the net reproductive rate of this species varied from 129.8 females/female on ash to 298.3 on olive. Thus, ash was relatively less suitable because of the lower reproductive rate of the insects reared on it [18].

6.2. Female Fecundity

As recorded in Greece, the number of laid eggs /female (fecundity) of *P. unionalis* ranged from 86 to 515, with an average of 209 [19]. On the basis of some studies carried out in Iran, various means of fecundity, such as 231, 125±29 and 182±18.1 had been recorded [16, 54, 72]. Several biological and reproductive studies had been conducted on the present pest in Turkey. Fecundity was determined in a range of 194-390 eggs/female at 25°C [18], 385 eggs/female [56], between 4 and 638 eggs during a period from August to September 2009 but between 29 and 643 eggs during a period from September to December 2009 [85] and 352±42.9 eggs/ female, at natural conditions in Adana region [10]. Several biological and reproductive studies had been conducted on the same pest, also, in Egypt. Different values of the female fecundity as 86-515, 141-882 and 414 eggs per female had been documented, depending on the region [27]. As well as Badawi et al. [84] determined the mean fecundity as 414 egg/female under constant conditions of 27.5° C and 65% RH and Shehata et al. [9] estimated a range from 630 to 653 eggs in the first generation but from 425 to 493 eggs in the second one, under the same laboratory conditions.

These various values of fecundity, as previously compiled, can be attributed to different factors. To shed some light on the factors interfering with the female fecundity, Badawi et al. [84] reported the necessity of copulation and fertilization of adult females for producing the production of fertile eggs. This report supported the previous observation of Rahhal [47] who dissected females just after death and observed a number of well developed eggs in their ovaries. Age of the adult females seemed to be another factor, since they laid more eggs in early ages and then fecundity decreased towards the end of their lives [10, 84]. Also, different host plants and the host plant's nutritional value are known to affect the insect reproduction [98-101]. It is interesting to refer the role of chemical stimuli in the oviposition. Kombargi et al. [123] examined the possible role of surface waxes as chemical stimuli. They found that surface waxes vary greatly within and among varieties and also contain compounds that hinder oviposition. Furthermore, according to this hypothesis, when many hosts are simultaneously offered to a female, it is expected that she will follow a hierarchical order of host preference by laying eggs on the best larval diet first, and then on the second best diet, and so forth [124]. According to Kumral et al. [18], the olive Shamy variety discouraged gravid females of *P. unionalis* from oviposition (lower fecundity), compared with Toffahi or Sennara varieties of olive. In addition to these factors, it has been found that feeding of adult females stimulate oviposition and fecundity. The number of eggs laid by fertilized females offered water (134.5 eggs per female) was much lower than those offered honey solution (414 eggs per female) [38]. Other factors interfering with the oviposition and fecundity can be added, such as the pest biotype differences, etc.

7. ECOLOGICAL PARAMETERS AND ETHOLOGICAL PHENOMENA OF *P. unionalis*

Many studies on various ecological aspects and behavioural characterization of *P. unionalis* had been reported in the literature (e.g., [15, 17, 18, 22, 24, 50, 63, 67, 68, 125-128]). We have reviewed here the reported works concerning the life table, population dynamics, sexual behaviour of adults, non-sexual behavioral patterns with special to the feeding and flight behaviors.

7.1. Life-Table Parameters as Affected by Biotic Factors

Leopold was the first to identify the value of life table in study of natural population [129]. Computing life tables become later a fairly current approach used by entomologists to study the insects' population dynamics. In other words, life table is an important analytical tool which provides detailed information of population dynamics to generate simple but more informative statistics. It also gives a comprehensive description of the survivorship, development and expectation of life [130-133]. Life table studies provide an opportunity to assess and evaluate the impact of specific mortality factors acting on insect population [134-136]. In addition, life tables used can make quantitatively and qualitatively evaluation of various host plants [137]. From a pest management standpoint, it is very useful to know when (and why) a pest population suffers high mortality. This is usually the time, when it is the most vulnerable. By knowing such vulnerable stages from life table, we can make time based application of control measure for the management of insect pests, to conserve the biotic and abiotic environmental constituents (for reviews, see [138, 139]).

The quantity and quality of food available to insects have important consequences for growth and development of larvae as well as the adult reproductive performance [86,140-144]. In other words, different host plants are known to play an important role in the life table parameters of phytophagous insects [98-100] such as the population increase and spread of the pest [100].

With regard to *P. unionalis*, few biological observations on the effects of several host plants of Oleaceae, such as olive, privet, jasmine and lilac, had been published [19, 22, 102]. Intrinsic rate of increase and mean generation duration reflect the suitability of the host plant, therefore three host plants (*viz.*, olive, ash and jasmine) were tested on some biological aspects. The insect could complete its life cycle on all plants, but ash was relatively less suitable because of the lower reproductive rate of the insect reared on it [18]. In addition to the role of host plants, life table parameters of *P. unionalis* can be affected by other biotic factors like natural enemies. As for example, parasitoid wasp *Goniozus legneri* affected the life table of this pest at different densities of it in Egypt [49]. Depending on this study, mortality due to parasitism and paralysis by *G. legneri* was density-dependent. This undoubtedly yielded a very low generation survival and population trend in all parasitoid-released treatments comparing to control.

Concerning the overwintering stage under field conditions, *P. unionalis* was observed overwintering in Italy during all developmental stages almost throughout the year, but mostly as the 2nd- and 3rd-instar larvae [23, 25]. In an extensive study on biology of this pest in Iran, the 5th generation provided the overwintering stages that mostly were as 3rd instar-5th instar larvae and pupae [54]. Overwintering larvae had been, also, observed in Turkey [56].

7.2. Population Dynamics

Suffice it to report the important studies including the population dynamics of *P. unionalis* in some being infested countries of Mediterranean region and Middle East. Many factors contribute to the population fluctuations. As for example, white-colored funnel traps captured significantly more males than brown traps, but were only marginally better than yellow or green funnel traps in Central and Northern Greece [55]. In the same country, Athanassious, et al. [145] studied the population dynamics of this pest. In the coastal region and Middle Egypt, El-Kenawy [146] recorded its highest populations in the month of May. Lababidi [42] carried out an ecological study during 2003 and 2004 in two regions in Syria and determined the population fluctuations of *P. unionalis*. In Iran, field observations indicated that the first generation being completed by the end of March and in early April. However, the population reaches its peak during the third and fourth generations [16].

The intrinsic rate of population increase is a basic parameter which an ecologist may wish to establish for an insect population [147]. In consistent with those results of Greenberg et al. [98] on *Spodoptera*

exigua and Hansen et al. [99] on *Sitotroga cerealella*, the intrinsic rate of population increase indicated that *P. unionalis* reared on three host plants exhibited exponential population growth in Turkey [18]. Recently, Kacar and Ulusoy [148] determined the adult and larval population fluctuations of the same pest in the same country by using sexual pheromone capsule, between the years 2009-2010. It was observed that shoot development and climatic factors (temperature and humidity) affected the larval population fluctuation.

7.3. Sexual Behaviour of Adults

Insects are especially suited for research of behavior because they are readily available in large quantities and have a short lifespan. Learning is involved in processes determining sexual selection and incipient speciation [149, 150]. Research in the past few decades has demonstrated that many insect species rely heavily on learning to decide about a variety of behaviors [151-153]. The role of learning in insect sexual behaviour has been either neglected or considered negligible. Quantifying the effects of learning on sexual behaviour in male and female insects can help us understand sexual selection and incipient speciation [154, 155]. In insects, courtship behaviour often includes the extensive use of the antennae, as reported from various insect orders [156-160]. This behavioral pattern had not been investigated for *P. unionalis*. According to the literature available to us, sexual behaviour of *P. unionalis* had not been fully described until now. However, studies on the female calling and male response, for copulation, can be reviewed herein.

7.3.1. Female Calling and Male Response

As reported by some authors [161-164], female calling in many species and pheromone production is synchronous and usually depended on the adult age as well as on other endogenous and exogenous factors. Also, specific pheromone components or their blends can be responsible for several aspects of male copulation in many moth species [165-167]. Few studies had been conducted to investigate the calling behaviour and pheromone production of *P. unionalis* in the world. Its adult females may follow a calling and pheromone biosynthesis pattern of many lepidopterous species in which pheromone production occurs during the period where females are calling and releasing pheromone [163, 168]. Mazomenos et al. [126] achieved a valuable study in this context. According to their results, compounds (E)-11-Hexadecenal and (E)-11-hexadecen-1-yl acetate were found in the abdomen tip extracts from *P. unionalis* females. In laboratory bioassays, both components elicited a low level of upwind flight by males. The two components were inactive when tested separately in the field, but their blend (3:7) was highly attractive to males. Because knowledge of the role of each component is essential for understanding the behavioral mechanisms associated with male mating behaviour, Mazomenos et al. [57] conducted another interesting study on the same insect. Calling activity and pheromone production is periodic and synchronous. Maximal calling and pheromone production was obtained in the fourth day.

7.3.2. Egg-Laying Behaviour

Different patterns of egg-laying behaviour of *P. unionalis* had been reported in different countries as reviewed herein. In Iran, adults were active early in the morning or during sunset while exhibiting a low level of activity, possibly with short flights during the warmer hours of the day [16]. The female mates one day after emergence and deposits her eggs in third day individually or in one row on the lower surfaces of leaves [54, 72]. Almost, similar observations had been reported in Italy [23] while Alford [169] observed the eggs singly or in small groups. In Turkey, adult females deposit their eggs individually or usually in egg-masses (of 6-36 eggs) [10, 56, 85]. In Egypt, Badawi et al. [84] reported that the copulation took place 24 hours after emergence and often after mid-night. It lasted for a period ranging from 45 to 105 minutes. According to the observation of Shehata et al. [9] adults were active at night, laying eggs singly at twilight. However, more than 60 % of eggs were laid singly, 36.3% in small groups (of 2-5 eggs) and 1.24% in groups (of 5 eggs) [47]. The egg mass contains 2-6 eggs or 2-86 eggs [27].

7.4. Non-Sexual Behavioral Patterns

Thoroughly examination of the available literature exhibited no other than the feeding behaviour of larvae and flight behavior of the *P. unionalis* adults as non-sexual behavioral patterns.

7.4.1. Feeding Behaviour

The first instar larvae (caterpillars) of *P. unionalis* aggregated and usually fed on the parenchyma of the olive leaves and on the tender buds. As they grow, they consume entire leaves and buds [10]. In its

second generation, larvae feed on fruits and seeds if they reach high population levels [49, 54, 61].

7.4.2. Flight Behaviour

Stelanesco [170] reported some notes on butterflies and moths recorded at sea off Eivissa and Barcelona (Western Mediterranean) in October 1996. All the species reported display a well-known migratory behaviour. One of them was *P. unionalis*. Before this report, Eitschberger et al. [171] recorded the same insect among seasonal migrants of the first order, and late in the season the offspring of the early migrants are involved in return flights to the southern areas from where their parents originated. Now *P. unionalis* is an international lepidopterous migratory pest in the tropical and mildly subtropical regions of the Old World [14-16]. It is a highly mobile moth dispersing even to northern Europe [32]. Another point of interest is the short and local flight. In Iran, Noori and Shirazi [16] observed active adults in early morning and during sunset while exhibiting a low level of activity, possibly with short flights during the warmer hours of the day. Also, Hegazi et al. [50] determined the seasonal flight trend of the same pest in three large plots of olive varieties during two successive fruiting seasons in Egypt.

8. ENHANCEMENT OF RESEARCH INTERESTS IN PHYSIOLOGY OF *P. unionalis*

Different metabolic and energetic aspects in insects were studied, such as: physiological and environmental considerations in bioenergetics [172, 173], energy metabolism during flight [174, 175], hormonal regulation on the energy metabolism [176-179], regulation of fat metabolism [180, 181], Chitin metabolism [182], reproductive physiology [183-185] and environmental physiology [142, 186, 187]. Unfortunately, the available literature contains no reported works on the metabolism, enzymology, hematology or other physiological aspects of *P. unionalis*. However, Mostafa et al. [188] characterized the proteins in pupal abdominal cuticle during the ecdysial periods of sclerotization in Egypt. In Egypt, also, Solaiman [189] studied the host preference of *P. unionalis* under laboratory conditions. Tophahy variety of olive plant was the preferable variety for the larvae followed by the Agyzy, while the Ballady variety was the least preferable. Also, the food consumption and host preference had been studied on certain leaf olive varieties [190].

9. CONCLUSIONS

The olive leaf moth *Palpita unionalis* gained a remarkable attention of researchers all over the world for its biological parameters and some of its ecological characteristics, beside the geographic distribution and economic impacts. On the other hand, this pest still needs research attention for investigation of several aspects such as energy metabolism, homeostasis, enzymatic patterns, chitin metabolism, hematology, resistance, immunity, reproductive physiology, environmental physiology the hormonal regulation of these processes, etc. Therefore, the present review enhances the research interests for these important aspects. However, information reviewed in this article will support the development of strategies for management of this pest.

REFERENCES

- [1] Haber G. and Mifsud D., Pests and diseases associated with olive trees in the Maltese Islands (Central Mediterranean). The Central Mediterranean Naturalist 4(3), 143-161(2007).
- [2] Budia P.B., Ecotoxicology of pesticides on natural enemies of olive groves. Ph.D. Thesis, Escuela Tecnica Superior de Ingenieros Agronomos, Madrid, Spain (2012).
- [3] Civantos L., La olivicultura en el mundo y en Espana. In: (Barranco D., Fernandez-Escobar R. and Rallo L., eds). "El cultivo del olivo". Ediciones Mundi-Prensa, Madrid, pp: 17-34(2001).
- [4] Herz A., Hassan S.A., Hegazi E., Nasr F.N., Youssef A.A., Khafagi W.E., Agamy E., Ksantini M., Jardak T., Mazomenos B.E., Konstantopoulou M.A., Torres L., Goncalves F., Bento A. and Pereira J.A., Towards sustainable control of lepidopterous pests in olive cultivation. Gesunde Pflanzen 57, 117-128(2005).
- [5] R. Spooner-Hart, L. Tesorihero, and B. Hall, Field Guide to Olive Pests, Diseases and Disorders in Australia. ISSN 1440-6845 (2007), 65 pp.
- [6] El-Basha N.A., Biological control of some pests infesting olive in Egypt. PhD Thesis. Faculty of Agriculture, Suez Canal University, Egypt (2002).
- [7] Mohamed F.M., Pathogenicity of three commercial products of entomopathogenic fungi

- Beauveria bassiana*, *Metarhizium anisopliae* and *Lecanicillium lecanii* against adult of olive fly *Bactrocera oleae* (Gmelin) (Diptera: Tephritidae) in the laboratory. Plant Prot. Sci. 3, 98-102(2009).
- [8] Broumas T., Haniotakis G., Liaropoulos C., Tomazou T. and Ragoussis N., The efficacy of an improved form of the mass-trapping method, for the control of the olive fruit fly, *Bactrocera oleae* (Gmelin) (Dipt., Tephritidae), pilot-scale feasibility studies. J.Appl. Entomol. 126(5), 217–223(2002).
- [9] Shehata W.A., Abou-Elkhair S.S., Youssef A.A. and Nasr F.N., Biological studies on the olive leaf moth, *Palpita unionalis* Hübner (Lepid., Pyralidae), and the olive moth, *Prays oleae* Bernard (Lepid., Yponomeutidae). J.Pest Sci. 76(6), 155-158(2003).
- [10] Yilmaz Ç. and Genç H., Determination of the life cycle of the olive fruit leaf moth, *Palpita unionalis* (Lepidoptera: Pyralidae) in the laboratory. Florida Entomologist 95(1), 162-170(2012).
- [11] Anonymous, Norfolk Moths. *Palpita vitrealis* (Rossi 1794) <http://www.norfolkmoths.co.uk/micros.php?bf=14080> (2011).
- [12] Santorini A.P. and Vessiliana-Alexopoulou P., Morphology of the internal reproductive system in male and female *Palpita unionalis* Hb. (olive pyralid). Entomologica 7, 29-47(1976).
- [13] Kirti J.S. and Rose H.S., Studies on Indian Species of the genus *Palpita*. J. Entomol. Res. 16(1), 62-77(1992).
- [14] Balachowsky A.S., Entomology applied in agriculture. 3rd ed., Vol. I, Lepidopteres. Masson Paris, 1131-1133 pp. (1972).
- [15] Tzanakakis M.E., Seasonal development and dormancy of insects and mites feeding on olive: a review. Netherlands J. Zool. 52, 87-224(2003).
- [16] Noori H. and Shirazi J., A study on some biological characteristics of olive leaf moth, *Palpita unionalis* Hübner (Lep: Pyralidae) in Iran. J. Agric. Sci. Tech. 14(2), 257-266(2012).
- [17] Kovanci B., Kumral N.A. and Akbudak B., Investigations on the population fluctuation of olive pyralid, *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae) in olive groves in Bursa Province of Turkey. Turkish J.Entomol. 30(1), 23-32(2006).
- [18] Kumral N.A., Kovanci B. and Akbudak B., Life tables of the olive leaf moth, *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae), on different host plants. J. Biol. Environ.Sci. 1(3), 105-110(2007).
- [19] Vassilaina-Alexopoulou P. and Santorini A.P., Some data on the biology of *Palpita unionalis* Hubner (Lepidoptera: Pyralidae) under laboratory conditions. Ann. Institut. Phytopathol. Benaki. 10(4), 320–326(1973).
- [20] Panigaj L.U., New records of Lepidoptera from Slovakian part of the Pieniny Mts. Przyroda i Człowiek 10, 75-83(2008).
- [21] Torres L., Bento A., Pereira J.A. and Mazomenos B., Flight phenology of *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae) in the north-east of Portugal. In: XI Congresso Ibérico de Entomologia. Funchal, Portugal, 13-17 Setembro, p. 179(2004).
- [22] Avidov Z. and Rosen D., Bionomics of the Jasmine moth (*Glyphodes unionalis* Hübner) in the coastal plain of Israel. Bull. Res. Council. Israel B 10, 77-89(1961):.
- [23] Triggiani O., *Margaronia unionalis* Hb. (Olive pyralid). Entomologica 7, 29-47(1971).
- [24] Fodale A.S. and Mule R., Bioethological observations on *Palpita unionalis* Hb. in Sicily and trials of defence. Acta Horticult. 286, 351–353(1990).
- [25] Katsoyannos P., Olive pests and their control in the Near East. FAO Plant Production and Protection Paper 115, 49-50(1992).
- [26] Antonelli R. and Rossi E., La *Palpita unionalis* Hbner (Lepidoptera, Pyraustinae): un fitofago di crescente importanza negli oliveti Toscani. Informatore Fitopatologico 34, 27-32(2004).
- [27] El-Kifl A.H., Abedsalam A.L. and Rahhal A.M., Biological studies on the olive leaf moth *Palpita unionalis* Hbn.. Bull. Soc. Entomol. d'Egypte 58, 337-344(1974).
- [28] Pollini A., Manuale di entomologia applicata. 1st ed., Edagricole - Edizioni agricole della Calderini, Bologna. pp. 14-62(1998).

- [29] Mateus H., Figueiredo E., Luz J.P., Céu Silva M. and Mexia A., The impact of Jasmine moth *Palpita vitrealis* (Rossi) in different olive cultivars in Cova da Beira. *Revista de Ciências Agrárias* 35(2), 205-209(2012).
- [30] Ryrholm N., Migrating moth in Sweden. Department of Animal Ecology Evolutionary Biology Center, Uppsala University, Norpyvogen 18 D, 5, 752-36(1988).
- [31] Sevansson I., Remarkable records on microlepidoptera in Sweden. *Entomologist tidskrift*. 109(2), 65-70(1988).
- [32] Tremewan W.G., Interesting Lepidoptera at m.v. light in West Cornwall (V.C. 1), 2001. *Entomol. Gaz.* 53(2002).
- [33] Kroon D.M., Lepidoptera of Southern Africa host-plants and other associations. A Catalogue, Lepidopterists' Society of Africa, Sasolburg, South Africa: Sasolburg Litho. (1999).
- [34] Vari L., Kroon D.M. and Kruger M., Classification and checklist of the species of Lepidoptera recorded in Southern Africa, Chatswood, Australia: Simple Solutions Australia (2002).
- [35] Copeland R.S., White I.M., Okumu M., Machera P. and Wharton R.A., Insects associated with fruits of the Oleaceae (Asteridae, Lamiales) in Kenya, with special reference to the Tephritidae (Diptera). *Bishop Muse. Bull. Entomol.* 12, 135-164(2004).
- [36] Kovanci B. and Kumral A., Insect pests in olive groves of Bursa (Turkey). 5th International Symposium on Olive Growing, Izmir (Turkey), 27 Sep-2 Oct 2004, p. 68(2004).
- [37] Cobanoglu S., Samyurek B.G., Alzoubi S., Samyurek H. and Gun Y., Important olive pests and their populations on two different olive varieties in southern Turkey (Degirmendere-Mersin). *Revue Ezzaitouna* 11(1), 1-14(2010).
- [38] Yilmaz, Ç. and Genç, H., Egg production and adult longevity of the olive leaf moth, *Palpita unionalis* Hubner (Lepidoptera: Pyralidae) on selected adult diets. *J. Tekirdag Faculty Agric.* 9(1), 1-5(2012).
- [39] Sahin C. and Genc H., An Alien Species in Çanakkale, Turkey: Jasmine Moth (*Palpita unionalis* Hübner, Lepidoptera: Pyralidae). 4th ESENIAS Workshop: International Workshop on IAS in Agricultural and Non-Agricultural Areas in ESENIAS Region 16-17 December 2013, Çanakkale, Turkey (Book of Abstracts) (2013).
- [40] Lababidi M.S., Field investigation on the efficacy of neem Azal®-T/S against olive buds moth *Marganonia unionalis* (Hbn.) (Lepidoptera: Pyralidae) in olive nurseries, Aleppo, Syria. Eighth Arab Congress of Plant Protection, 12-16 October 2003, El-Beida, Libya (2003).
- [41] Lababidi M.S., Research on the parasitoid *Dolichogenidea trachalus* (Nixon) (Hymenoptera: Braconidae), collected from the olive buds moth (Jasmine moth) *Palpita unionalis* Hübner (Lepidoptera: Pyralidae) in Syria. 56. Deutsche Pflanzenschutztagung in Kiel, 22.-25. September 2008(2008).
- [42] Lababidi M.S., Ecological studies on the olive buds moth (Jasmine moth) *Palpita unionalis* Hubner and its associated natural enemies in Syria. Ninth Arab Congress of Plant Protection, 19-23 November 2006, Damascus, Syria (2006).
- [43] Delucca C., The place of the Lepidoptera in the zoogeography of the Maltese islands. *Rapport etprocess-verbaux de reunions de la C.I.E.S.M.M* 18(2), 511-515(1965).
- [44] A. Valletta, The moths of the Maltese islands. Progress Press, 1973, 72 pp.
- [45] Sammut P.M., A systematic and synonymic list of Lepidoptera of the Maltese Islands. *Neue Entomologische Nachrichten*, 13, 124 pp. (1984).
- [46] P.M. Sammut, Il-Lepidoptera. Kullana Kulturali, Pin Publications, Malta, 2000, x + 245 pp.
- [47] Rahhal M.M., Studies on certain olive pests in U.A.R. M.Sc. Thesis, Al-Azhar University, Egypt (1972).
- [48] Foda S.M.; Awadallah, A.M. and Abou-El-Ghar, M.R. (1976): Chemical control of the olive moth *Palpita unionalis* Hb. *Agric. Res. Rev.*, 54 (1): 153-159.
- [49] El-Basha N.A. and Mandour N.S., Effect of *Goniozus legneri* Gordh (Hymenoptera: Bethyilidae) on the life table of *Palpita unionalis* Hb. (Lepidoptera: Pyralidae) in the laboratory. *Egyptian J.Biol. Pest Control* 16(1), 5-11(2006).

- [50] Hegazi E.M., Konstantopoulou M.A., Khafagi W.E., Schlyter F., Herz A., Raptopoulos D.G., Hassan S. and Atwa A., The population trend of *Palpita unionalis* in different olive varieties in Egypt. *Phytoparasitica* 40(5), 451-459(2012).
- [51] Saieb H., A report of new olive pest in Iran. *Iranian Entomol. Soc. News letters.*, 1(3), 4-5(1999).
- [52] Pazuki A., Olive Leaf Moth. *J. Entomol. Soc. Iran* 7(1), 2-10(2000).
- [53] Ghavami S., Preliminary studies on olive trees infestation rate by olive leaf moth, *Palpita unionalis* in Guilan and Qazvin provinces in 2000. *J. Entomol. Soc. Iran* 8(1), 88-95(2000).
- [54] Fazel M. and Azimi zadeh N., Biology of jasmine moth *Palpita unionalis* (Lep., Pyralidae) in laboratory and field conditions in Zanjan (Tarom-Olia) region. *Appl. Entomol. Phytopathol.* 78(1), 1-24(2010).
- [55] Athanassiou G.C., Nickolas G., Kavallieratos B. and Mazomenos E., Effect of trap type, trap color, trapping location, and pheromone dispenser on captures of male *Palpita unionalis* (Lepidoptera: Pyralidae). *J. Econ. Entomol.* 97(2), 321-329(2004).
- [56] Khaghaninia S. and Pouraband R.F., Investigation on biology of olive leaf worm *Palpita unionalis* Hübner (Lepidoptera: Pyralidae) in constant laboratory conditions. *Munis Entomol. Zool.* 4(2), 320-326(2009).
- [57] Mazomenos B.E., Konstantopoulou M., Stefanou D., Skareas S. and Tzeiranakis L.C., Female calling behaviour and male response to the synthetic sex pheromone components of *Palpita unionalis* (Lepidoptera: Pyralidae). *Bull. OILB/SROP* 25(9), 203-211(2002).
- [58] Azimi Zadeh N., Abdollahi G., Movahhedie F.M. and Shojaiy M., Study on common privet (*Ligustrum vulgare*) desirability for breeding olive leaf moth, Hübner (Lepidoptera: Pyralidae), in the laboratory. *Proc. 16th Iranian Plant Prot. Congress, Tabriz, Iran*, 352 pp. (2003).
- [59] El-Hakim A.M. and El-Helmy, Survey of population studies on olive leaf pest in Egypt. *Bull. Soc. Entomol. Egypt.* 62, 213- 220(1982).
- [60] Pinto M. and Salemo G., The olive pyralid. *Informator, Agrio.* 51 (43), 77-81(1995).
- [61] Grossley S., *Palpita unionalis*. Retrived April, 2001. Available from <http://www.nysaes.cornell.edu/fst/faculty/acree/pheromet/ins/palpiunion.html> (2000).
- [62] Pinto M.Lo and Salerno G., Bioethological observations on *Apanteles syleptae* Ferriere (Hymenoptera: Braconidae), solitary parasitoid of *Palpita unionalis* Hbner (Lepidoptera: Pyraustidae). *Phytophaga Palermo* 5, 3-19(1994).
- [63] Gargani E., *Margaronia unionalis* (Hübner) (Lepidoptera: Pyraustidae) on *Jasminum*: bio-ethological observations and damage. *J. Biol. Environ. Sci.* 28(7), 71-76(1999).
- [64] M.C. Lopez-Villalta, Olive pest and disease management. International Olive Oil Council, Madrid, Spain (1999).
- [65] Yilmaz C. and Genc H., A research on sex differentiation on immature stages of olive leaf moth (*Palpita unionalis* Hubner. The 4th Plant Protection Congress in Turkey (2011).
- [66] Mori P., Minor insect pests of olive. *Vita in campagna* 13(5), 26- 27(1995).
- [67] Hegazi E.M., Konstantopoulou M.A., Milonas P., Herz A., Mazomenos B.E., Khafagi W.E., Zaitun A., Abdel-Rahman S.M., Helal I. and El-Kemny S. Mating disruption of the jasmine moth *Palpita unionalis* (Lepidoptera: Pyralidae) using a two pheromone component blend: A case study over three consecutive olive growing seasons in Egypt. *Crop Protection* 26(6), 837-844(2007).
- [68] Hegazi E., Herz A., Hassan S.A., Khafagi W.E., Agamy E., Zaitun A., Mostafa S., El-Said S. and Khamis N., Field effectiveness of indigenous species of the genus *Trichogramma* (Hymenoptera, Trichogrammatidae) to control the olive moth (Prays oleae, Lepidoptera, Yponomeutidae) and the jasmine moth (*Palpita unionalis*, Lepidoptera, Pyralidae) in an olive plantation in Egypt. In: (Herz A., Zimmermann O. and Hassan S.A., eds.). "Egg Parasitoid News" International Organization for Biological Control IOBC, 18: (Book of Abstracts), Darmstadt, Germany (2007).
- [69] Mkize N., Hoelmer K.A. and Villet M.H., A survey of fruit-feeding insects and their parasitoids occurring on wild olives, *Olea europaea* sp. *cuspidata*, in the Eastern Cape of South Africa. *Biocontrol Sci. Technol.* 18, 991-1004(2008).

- [70] Kacar G. and Ulusoy M.R., Determination of predators and parasitoids of olive leaf moth *Palpita unionalis* (Hüb.) (Lepidoptera: Pyralidae) in olive orchards in the eastern Mediterranean region of Turkey. *Türk. biyo. müc. derg.* 2(1), 39-48(2011).
- [71] Stefanos S., Ultrastructure of antennal sensory receptors of two pests of the olive tree, the olive leaf moth *Palpita unionalis* Hubn. and the olive moth *Prays oleae* Bern.. *Alexandria Sci. Exchange* 24, 257-272(2003).
- [72] Alavi J., Biological study of olive leaf moth, *Palpita unionalis* Hbn. (Lep., Pyralidae) in Golestan Province, Iran. *J. Entomol. Res.* 2(2), 85-95(2010).
- [73] Loi G., The influence of temperature on the embryonic development of *Palpita unionalis* (Lepidoptera: Pyralidae) and data on the fecundity and longevity of the imagines. *Frustula Entomol.* 13(26), 159-168(1990).
- [74] Margaritis L.H. and Mazzini M., Structure of the egg. In: (Harrison F.W. and Locke M., eds) "Microscopic anatomy of invertebrates", vol. 11C, pp. 995-1037, Wiley-Liss, New York (1998).
- [75] Waring G.L., Morphogenesis of the eggshell in *Drosophila*. *Int. Rev. Cytol.* 198, 67-108(2000).
- [76] Woods H.A. and Bonnacaze R.T., Insect eggs at a transition between diffusion and reaction limitation: temperature, oxygen, and water. *J. Theor. Biol.* 243, 483-492(2006).
- [77] Margaritis L.H., Structure and physiology of the eggshell. In: (Kerkut G.A. and Gilbert L.I., eds) "Comprehensive insect physiology, biochemistry, and pharmacology", vol. 1, pp. 153-230, Pergamon Press, Oxford (1985).
- [78] Zissler D., From egg to pole cells: ultrastructural aspects of early cleavage and germ cell determination in insects. *Microsc. Res. Tech.* 22, 49-74(1992).
- [79] M.J. Klowden, *Physiological Systems in Insects*. 2nd edition, 2007, 688 pp. Academic Press Publications, Elsevier Inc., USA.
- [80] Tautz D. and Sommer R.J., Evolution of segmentation genes in insects. *Trends Genet.* 11, 23-27(1995).
- [81] Kawamura N., Fertilization and the first cleavage mitosis in insects. *Dev. Growth Differ.* 43, 343-349(2001).
- [82] Davis G. and Patel N.H., Short, long, and beyond: molecular and embryological approaches to insect segmentation. *Annu. Rev. Entomol.* 47, 669-699(2002).
- [83] Hamdoun A. and Epel D., Embryo stability and vulnerability in an always changing world. *Proc. Natl. Acad. Sci. USA* 104, 1745-1750(2007).
- [84] Badawi A., Awadallah A.M. and Foda S.M., On the biology of the olive leaf moth *Palpita unionalis* Hbn. (Lep., Pyralidae). *Z. Ang. Entomol.* 80, 103-110(1976).
- [85] Kacar G. and Ulusoy M.R., The biology of the Olive leaf moth *Palpita unionalis* (Hüb.) (Lepidoptera: Pyralidae) in natural conditions. *Türk. entomol. derg.* 36(3), 335-344(2012).
- [86] M.R. Speight, M.D. Hunter and A.D. Watt, *Ecology of Insects, Concepts and Applications*. 2nd ed., 2008, A John Wiley and Sons, Ltd., Publication, West Sussex, UK, 628 pp.
- [87] Wigglesworth V.B. The significance of "apolysis" in the moulting of insects. *J. Entomol.* 47, 141-149(1973).
- [88] Cole B.J., Growth ratios in holometabolous and hemimetabolous insects. *Ann. Entomol. Soc. Am.* 73, 489-491(1980).
- [89] Bernays E.A., Evolutionary contrasts in insects: nutritional advantages of holometabolous development. *Physiol. Entomol.* 11, 377-382(1986).
- [90] Reynolds S.E. and Samuels R.I., Physiology and biochemistry of insect molting fluid. *Adv. Insect Physiol.* 26, 157-232(1996).
- [91] Truman J.W. and Riddiford L.M., The origins of insect metamorphosis. *Nature* 401, 447-452(1999).
- [92] Barnes R.S.K., Calow P., Olive P., Golding D. and Spicer J., *The Invertebrates: a Synthesis*. 2001, Blackwell Scientific Publications, Oxford. 512 pp.
- [93] Nijhout H.F., The control of growth. *Development* 130, 5863-5867(2003).

- [94] Martelli G.M., Intorno a due specie di Lepidoteri dei generi Zelleria e Glyphodesviventis sull'olivo. Bull. Lab. Zool. Gen. Agrar. R. Scuola Sup. Agric. 10, 89-102(1916).
- [95] Yilmaz C., Study of feeding physiology of olive leaf moth (*Palpita unionalis* Hübner) on its natural host and artificial diets. Türkiye IV. Bitki Koruma Kongresi Bildirileri, 28-30(2011).
- [96] Azimi Zadeh N., Fazel M.M. and Parvar A., Comparison of some biological parameters of jasmine moth, *Palpita unionalis* Hubner (Lep., Pyralidae), on olive and privet leaves in laboratory conditions. J. Entomol. Res. 4(1), 33-41(2011).
- [97] Yilmaz Ç. and Genç H., The first attempt to rear olive leaf moth [(*Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae)] on artificial diet. J. Biol. Environ. Sci. 7(19), 25-32(2013).
- [98] Greenberg S.M., Sappington T.W., Legaspi B.C., Liu T.X. and Setamou M., Feeding and life history of *Spodoptera exigua* (Lepidoptera: Noctuidae) on different host plants. Ann. Entomol. Soc. America 94(4), 566-575(2001).
- [99] Hansen L.S., Skovgard H. and Hell K., Life table study of *Sitotroga cerealella* (Lepidoptera: Gelechiidae), a strain from West Africa. J. Econ. Entomol. 97(4), 1484-1490(2004).
- [100] Liu Z.D., Li D.M., Gong P.Y. and Wu K., Life table studies of the cotton bollworm, *Helicoverpa armigera* (Hubner) (Lepidoptera: Noctuidae), on different host plants. Environ. Entomol. 33(6), 1570-1576(2004).
- [101] Pereyra P.C. and Sanchez N.E., Effects of two Solanaceous plants on developmental and population parameters of the Tomato leaf miner, *Tuta absoluta* (Meyrick) (Lepidoptera: Gelechiidae). Neotropical Entomol. 35(5), 671-676(2006).
- [102] Arambourg Y., Pyralidae. *Margaronia unionalis* Hbn. In: (Arambourg Y., ed) "Traite d'entomologie oleicole". International Olive Oil Council, Madrid, Spain, pp: 75-80(1986).
- [103] El-Khawas M.A., Integrated control of insect pests on olive trees in Egypt with emphasis on biological control. Ph.D. Thesis, Fac. of Sci., Cairo Univ. Egypt (2000).
- [104] Z. Avidov and I. Harpaz, Plant Pests of Israel. Israel Universities Press, Jerusalem (1969).
- [105] Zervas G.A., Katranis N., Kazanas I. and Skotaras N., Problems in olive culture from the lepidopteran *Palpita unionalis* (Hbn.) (Pyralidae). In: Proc. 2nd Panhellenic Entomol. Conf, Athens, 11-13 Nov. 1989, pp: 46-56 (1989).
- [106] El-Sherif L.S. and Kaschef A.H., Morphological and biological studies on *Apanteles syleptae* F. (Hymenoptera: Braconidae) recovered from the Jasmine moth, *Palpita unionalis* Hb. Zeitschrift für Angewandte Entomologie 84(4), 419-424(1977).
- [107] Knipling, E.F. (1966): Introduction. In: (Smith, C.N. ed.) "Insect Colonization and Mass Production". pp.: 1-12, Academic Press, New York.
- [108] Herz A. and Hassan S.A., Are indigenous strains of *Trichogramma* sp. (Hym., Trichogrammatidae) better candidates for biological control of lepidopterous pests of the olive tree? Biocontrol Sci. Technol.16(8), 841-857(2006).
- [109] Mansour A., Biocontrol studies on using *Bracon* sp. (Hymenoptera: Braconidae) to control lepidopterous pests infesting Egyptian olive. Ph.D. Thesis, Faculty of Science, Al-Azhar University, Egypt (2011).
- [110] Kleespies R.G., Wang Y., El Salamouny S., Awad M., Agamy E., Salama R. and Jehle J.A., Identification of a new multiple nucleopolyhedrovirus isolated from the Jasmine moth, *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae) in Egypt. 47th Annual Meeting of the Society for Invertebrate Pathology and International Congress on Invertebrate Pathology and Microbial Control [3-7 August 2014, University of Mainz, Germany] , vol. 47, pp.: 129-130(2014).
- [111] E.S. Vanderzant, Defined diets for phytophagous insects. Academic Press, New York, 1966, 277 pp.
- [112] Bottger G.T., Development of synthetic food media for use in nutrition studies of the European corn borer. J. Agric. Res. 65(10), 493-500(1942).
- [113] Shorey H.H. and Hale R.L., Mass rearing of the larvae of nine noctuid species on a simple artificial medium. J. Econ. Entomol. 58, 522-524(1965).
- [114] C.N. Smith, Insect Colonization and Mass Production. Academic Press, New York, 1966, 618 pp.

- [115] Dimetry N.Z., A successful technique for mass rearing of the cotton leaf worm, *Spodoptera littoralis* (Boisd.) on an artificial diet. Z. Angew. Entomol. 66, 408-4011(1970).
- [116] Klein M., Navon A., Andreadou M. and Keren S., Development of an artificial diet for mass rearing of the spiny bollworm, *Earias insulana*. Phytoparasitica 9, 145-148(1981).
- [117] P. Singh and R.F. Moore, Handbook of Insect Rearing Vol. II. Elsevier Science Publishers, Amsterdam, 1985, 514 pp.
- [118] Guy R., Leppla N.C., Rye J.R., Gren C.W., Barretteand S.L. and Hollien K.A., *Trichoplusia ni*. In: (Singh P. and Moore R.F., eds). "Handbook of Insect Rearing", Vol. II, Elsevier Science Publishers. Netherland, pp. 487-493(1985).
- [119] Genc H. and Nation J.L., An Artificial Diet for the Butterfly *Phyciodes phaon* (Lepidoptera: Nymphalidae). Florida Entomologist 87, 194-198(2004).
- [120] J.L. Nation, Insect Physiology and Biochemistry. CRC Press, 2002, pp. 485.
- [121] Richard O.W., The theoretical and practical study of natural insect populations. Annu. Rev. Entomol. 6, 147-162(1961).
- [122] Varley G.C. and Gradwell G.R., Recent advances in insect population dynamics. Annu. Rev. Entomol. 15, 1-24(1970).
- [123] Kombargi W.S., Michelakis, S. E. and Petrakis, C.A. (1998): Effect of olive surface waxes on oviposition by *Bactrocera oleae* (Diptera: Tephritidae). J.Econ. Entomol., 91: 993-998.
- [124] Thompson J.N., Evolutionary ecology of the relationship between oviposition preference and performance of offspring in phytophagous insects. Entomol. Exp. Appl. 47, 3-14(1988).
- [125] Fodale A.S., Mule R. and Tucci A., Bioethiological observation on *Margaronia unionalis* Hb. in Sicily and trails on its control. Annalidell. Instituto sperimental per Io livicotura 10, 31-44(1990).
- [126] Mazomenos B.E., Raptopoulos D., Lefkidou I. and Mazomenos- Pantazi A., Female sex pheromone components of the jasmine moth *Palpita unionalis* (Lepidoptera: Pyralidae). J. Chem. Ecol. 20(3), 745-751(1994).
- [127] Sparks T.H., Dennis R.L.H., Croxton P.J. and Ccade M., Increased migration of Lepidoptera linked to climate change. Eur. J. Entomol. 104, 139-143(2007).
- [128] Umbers K.D.L., Symonds M.R.E. and Kokko H., The mathematics of female pheromone signaling: Strategies for aging virgins. The American Naturalist 185(3), 417-432(2015).
- [129] Harcourt D.G., The development and use of life tables in the study of natural insect populations. Annu. Rev. Entomol. 14, 175-196(1969).
- [130] Gabre R.M., Adham F.K. and Chi H., Life table of *Chrysomya megacephala* (Fabricius) (Diptera: Calliphoridae). Acta Oecologica 27, 179-183(2004).
- [131] Ali A. and Rizvi P.Q., Age specific survival and fecundity table of *Coccinella septempunctata* L. (Coleoptera: Coccinellidae) on different aphid species, Ann. Plant Protec. Sci. 15, 329-334(2007).
- [132] Afrane Y.A., Zhou G., Lawson B.W., Githeko A.K. and Yan G., Life table analysis of *Anopheles arabiensis* in Western Kenya Highlands: Effects of land covers on larval and adult survivorship. Am. J. Trop. Med. Hyg. 77, 660-666(2007).
- [133] Yazdani M. and Samih M.A., Life table attributes of *Clitostethus arcuatus* (Coleoptera: Coccinellidae) feeding on *Trialeurodes vaporariorum* and *Siphoninus phillyreae* (Hemiptera: Aleyrodidae). J. Asia-Pacific Entomol. 15, 295-298(2012).
- [134] A.S. Atwal and S.S. Bain, Applied Animal Ecology (Ludhiana: Kalyani Publishers), 1974, New Delhi, India.
- [135] Bellows Jr. TS., Van Driesche R.G. and Elkinton J.S., Life-table construction and analysis in the evaluation of natural enemies. Annu. Rev. Entomol. 37, 587-612(1992).
- [136] Mohapatra L.N., Life table studies of spotted bollworm *Earias vitella* (Fab.) on cotton. Indian J. Agric. Res. 41, 63-66(2007).
- [137] Ambegaonkar J.K. and Bilapate G.G., Life fecundity tables of *Earias vitella* (Fabricius) on cotton and okra. Proced. Ind. Sci. Acad. B 48(2), 207-213(1982).

- [138] Aziz M.A., Iftkhar A. and Hanif M., Life table studies of *Trilochoa virescence* (Bombycidae: Lepidoptera) on *Ficus nitida*. Asian J. Agric. Biol. 1(1), 2-7(2013).
- [139] Kakde A.M., Patel K.G. and Tayade Sh., Role of Life Table in Insect Pest Management--A Review. IOSR J.Agric. Veter. Sci. 7(1), 40-43(2014).
- [140] N.E. Stamp and T.M. Casey, Caterpillars: ecological and evolutionary constraints on foraging. 1sted., 1993, Chapman and Hall, New York, 548 pp.
- [141] L.M. Schoonhoven, T. Jermy, and J.J.A. van Loon, Insect-Plant Biology. 1st ed., 1998, Chapman and Hall, London. 409 pp.
- [143] S.L. Chown and S.W. Nicolson, Insect physiological ecology: mechanisms and patterns. 1st ed., 2004, Oxford University Press, New York 243 pp. Nespolo R.F., Castaneda L.E. and Roff D.A., Dissecting the variance-covariance structure in insect physiology: the multivariate association between metabolism and morphology in the nymphs of the sand cricket (*Gryllus firmus*). J. Insect Physiol. 51(8), 913-21(2005).
- [144] Chown S.L. and Terblanche J.S., Physiological diversity in insects: ecological and evolutionary contexts. Adv. Insect Physiol. 33, 50-152(2007).
- [145] Athanassios C.G., Kavallieratos N.G. and Mazomenos B.E., Population dynamics of *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae) in Central and Northern Greece. Bulletin OILB/SROP 28(9), 117-120(2005).
- [146] El-Kenawy A., Management of some economic olive insect pests with emphasis on their biological control in two agro-ecosystems in Egypt. M.Sc. Thesis, Istituto Agronomico Mediterraneo di Bari (IAMB)-Centre international de hautes études agronomiques méditerranéennes (CIHEAM), Italy (2012).
- [147] Birch L.C., The Intrinsic rate of natural increase of an insect population. J. Animal Ecol. 17(1), 15-26(1948).
- [148] Kacar G. and Ulusoy M.R., Olive leaf moth, *Palpita unionalis* (Hübner) (Lepidoptera: Pyralidae) population fluctuation in olive orchards and damage rate on fruits in the eastern Mediterranean region. Bitki Koruma Bulteni 53(1), 7-31(2013).
- [149] Gibson R.M. and Langen T.A., How do animals choose their mates? Trends Ecol. Evol. 11, 468-470(1996).
- [150] Beltman J.B., Haccou P. and Tencate C., Learning and colonization of new niches: a first step toward speciation. Evolution 58, 35-46(2004).
- [151] D.R. Papaj and A.C. Lewis (eds), Insect learning: ecological and evolutionary perspectives. The 1st ed., 1993, 240 pp., Chapman & Hall, New York.
- [152] Dukas R., Effects of perceived danger on flower choice by bees. Ecol. Lett. 4, 327-333(2001).
- [153] Greenspan R.J. and van Swinderen B., Cognitive consonance: complex brain functions in the fruit fly and its relatives. Trends Neurosci. 27, 707-711(2004).
- [154] Alexander R.D., Marshall D.C. and Cooley J.R., Evolutionary perspectives on insect mating. In: (Choe J.C. and Crespi B.J., eds.) "The Evolution of Mating Systems in Insects and Arachnids", pp. 4-31, Cambridge, New York: Cambridge University Press (1997).
- [155] Dukas R., Learning in the context of sexual behaviour in insects. Animal Biol. 56(2), 125-141(2006).
- [156] Ho H.Y. and Millar, J.G., Identification and synthesis of a male-produced sex pheromone from the stink bug *Chlorochroa sayi*. J. Chem. Ecol. 27, 1177-1201(2001).
- [157] Cheng L., Howard R.W., Campell J.F., Charlton R.E., Nechols J.R. and Ramaswamy S.B., Mating behavior of *Cephalonomia tarsalis* (Ashmead) (Hymenoptera: Bethyilidae) and the effect of female mating frequency on offspring production. J. Insect Behav. 17, 227-245(2004).
- [158] Rowe L., Westlake K.P. and Currie D.C., Functional significance of elaborate secondary sexual traits and their evolution in the water strider genus *Rheumatobates*. Can. Entomol. 138, 568-577(2006).
- [159] Friberg M., Vongvanich N., Borg-Karlson A.K., Kemp D.J., Merilaita, S. and Wiklund C., Female mate choice determines reproductive isolation between sympatric butterflies. Behav. Ecol. Sociobiol. 62, 873-886(2008).

- [160] Steiner S.M., Kropf C., Graber W., Nentwig W. and Klopstein S., Antennal courtship and functional morphology of tyloids in the parasitoid wasp *Syrphoctonus tarsatorius* (Hymenoptera: Ichneumonidae: Diplazontinae). *Arthropod Struc. Devel.* 39, 33-40(2010).
- [161] Dunkelblum E., Kehat M., Harel M. and Gordon D., Sexual behaviour and pheromone titre of the *Spodoptera littoralis* female moth. *Entomol. Exper. Appl.* 44, 241-247(1987).
- [162] Noldus L.P. and Potting J.J., Calling behaviour of *Mamestra brassicae*: effect of age and photoperiod. *Entomol. Exp. Appl.* 56, 23-30(1990).
- [163] Babilis N.A. and Mazomenos B.E., Pheromone production in *Sesamia nonagrioides*: Diel periodicity and effect of age and mating. *J. Insect. Physiol.* 38, 561-564(1992).
- [164] Kakimura M. and Tatsuki S., Diel rhythms of calling behavior and pheromone production of oriental tobacco budworm moth *Helicoverpa assulta* (Lepidoptera: Noctuidae). *J. Chem. Ecol.* 12, 2953-2963(1993).
- [165] Linn C.H. and Gaston L.K., Behavioral responses of male *Trichoplusia ni* in a Sustained flight tunnel to the two sex pheromone components. *Environ. Entomol.* 10, 379-385(1981).
- [166] Linn C.H. and Gaston L.K., Behavioral function of the components and the blend of the sex pheromone of the cabbage looper, *Trichoplusia ni*. *Environ. Entomol.* 10, 751-755(1981).
- [167] Quartey G.K. and Coaker T.H., Role of the sex components in the orientation of *Ephestia cautella*. *Entomol. Exp. Appl.* 66, 237-245(1993).
- [168] Raina A.K., Klun J.A. and Stadelbacher E.A., Diel periodicity and effect of age and mating on female sex pheromone titer in *Heliothis zea* (Lepidoptera: Noctuidae). *Ann. Entomol. Soc. Am.* 79, 128-131(1986).
- [169] D.V. Alford, A color atlas of pest of ornamental trees, shrubs and flowers. The 1st ed., 1998, Halsted Press: An imprint of John Wiley & Sons New York Toronto. 448 pp.
- [170] Stelanesco C., Bunerflies and moths (Insecta, Lepidoptera) recorded at sea off Eivissa and Barcelona (Western Mediterranean) in October 1996. *Boll. Soc. Hist. Nat. Balears* 40, 51-56(1997).
- [171] Eitschberger U., Reinhardt R. and Steiniger H., Wanderfalter in Europa (Lepidoptera). Appeal for international cooperation in the research of the migration of insects. *Atalanta* 22(1), 1-67(1991).
- [172] Watanabe M., Kikawada T., Minagawa N., Yukuhiro F. and Okuda T., Mechanism allowing an insect to survive complete dehydration and extreme temperatures. *J. Exp. Biol.* 205, 2799-802(2002).
- [173] Gade G., Regulation of intermediary metabolism and water balance of insects by neuropeptides. *Annu. Rev. Entomol.* 49, 93-113(2004).
- [174] Beenackers A.M.T., Vanderhorst D.J. and Vanmarrewijk W.J.A., Insect flight metabolism. *Insect Biochem.* 14, 243-60(1984).
- [175] Suarez R.K., Energy metabolism during insect flight: biochemical design and physiological performance. *Physiol. Biochem. Zool.* 73(6), 765-771(2000).
- [176] Gade G. and Auerswald L., Mode of action of neuropeptides from the adipokinetic hormone family. *Gen. Comp. Endocrinol.* 132, 10-20(2003).
- [177] Socha R., Kodrok D., Simek P. and Patockova M., The kind of AKH-mobilized energy substrates in insects can be predicted without a knowledge of the hormone structure. *Eur. J. Entomol.* 101, 29-35(2004).
- [178] Schooley D.A., Horodyski F.M. and Coast G.M., Hormones controlling homeostasis in insects: endocrinology. In: (Gilbert LI, Iatrou K. and Gill S., eds). "Comprehensive Molecular Insect Science". Vol. 3. Amsterdam: Elsevier. pp.: 493-550(2005).
- [179] Lorenz M.W. and Gäde G., Hormonal regulation of energy metabolism in insects as a driving force for performance. *Integ. Comp. Biol.* 49(4), 380-392(2009).
- [180] Arrese E.L., Canavoso L.E., Jouni Z.E., Pennington J.E., Tsuchida K. and Wells M.A., Lipid storage and mobilization in insects: current status and future directions. *Insect Biochem. Mol. Biol.* 31, 7-17(2001).

- [181] Athenstaedt K. and Daum G., The life cycle of neutral lipids: synthesis, storage and degradation. *Cell Mol. Life Sci.* 63, 1355-1369(2006).
- [182] Merzendorfer H. and Zimoch L., Chitin metabolism in insects: structure, function and regulation of chitin synthases and chitinases. *J. Exp. Biol.* 206, 4393-4412(2003).
- [183] F. Engelmann, The physiology of insect reproduction. The 1st ed., 1970, 307 pp., Pergamon Press,
- [184] Hayes E. and Wall R., Age-grading adult insects: a review. *Physiol. Entomol.* 24, 1-10(1999).
- [185] Simonet G., Poels J., Claeys I., Van Loy T., Franssens V., De Loof A. and Broeck J.V., Neuroendocrinological and molecular aspects of insect reproduction. *J. Neuroendocrinol.* 16(8), 649-659(2004).
- [186] Brown J.H., Gillooly J.F., Allen A.P., Savage V.M. and West G.B., Toward a metabolic theory of ecology. *Ecology* 85, 1771-1789(2004).
- [187] Zhao L. and Jones W.A., Expression of heat shock protein genes in insect stress responses. *Invertebrate Survival J.* 9, 93-101(2012).
- [188] Mostafa Z.K., Rashad E.M. and Ramadan Kh.S., Pupal cuticle proteins of *Palpita unionalis* Hb (Lepidoptera: Pyralidae) characterization and profiles during sclerotization. *J. Egypt. Ger. Soc. Zool.* 42(E), 101-118(2003).
- [189] Solaiman R.H.A., Ecological, Biological, and Sex Pheromone Studies with the olive leafhopper, *Docotettix cornutus*; the olive whitefly, *Aleurolobus olivinus* and the olive leaf moth, *Palpita unionalis*. Ph.D. Thesis, Faculty of Agric. (Fayoum), Cairo Univ., Egypt (2005).
- [190] Salama R.A.K., Swailam S.M., Agamy E.A. and Awad M.A., Food consumption and host preference of jasmine moth, *Palpita unionalis* (Hubner) (Lepidoptera: Pyralidae) on certain leaf olive varieties. *Egypt. J. Appl. Sci.* 22(12), 252-260(2007).