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Comparing the foraging behaviour and pollination efficiency of *Apis mellifera* with *Xylocopa olivacea* (Apidae: Hymenoptera) on *Citrullus lanatus* flowers

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Abstract

Comparing the foraging behaviour and the pollination efficiency of honey bee (*Apis mellifera*) with a carpenter bee (*Xylocopa olivacea*) in the farmer garden of watermelon (*Citrullus lanatus*) was conducted in Maroua (Cameroun) in 2016 and 2017. Several foraging parameters were assessed for each bee species during the blooming period of watermelon and comparisons were made between fruit and seed yields from four pollination treatments on female flowers including the no bee visit treatment (T_0), the one bee visit treatment for *A. mellifera* (T_1) and *X. olivacea* (T_2) and the unrestricted treatment (T_3). Results showed an important difference in the rhythm of activity between the two bee species with a peak of floral activity of *X. olivacea* at 07:00-08:00 a.m. time interval and that of *A. mellifera* at 09:00-10:00 a.m. *A. mellifera* was prominent than *X. olivacea* regarding the frequency of floral visits in 2016 (77.74 and 22.26 %) and in 2017 (81.28 and 18.72 %) and the density of individuals per 1000 flowers in 2016 (442/1000 flowers and 97/1000 flowers). Despite the higher foraging intensity of *A. mellifera* compared with that of *X. olivacea*, the carpenter bee was a more efficient pollinator than the honey bee. Indeed, the mean treatments for fruiting rate, mean fruit weight, mean fruit diameter, and mean mature seeds per fruit of watermelon were higher on the one visit basis in T_2 than T_1 . Moreover, the unrestricted treatment T_3 produced fruit with the best marketable value than restricted treatments T_2 , T_1 , and T_0 . *X. olivacea* should be associated with *A. mellifera* in a watermelon field to optimize the pollination of this crop for obtaining improved yields. Considering very high pollinating efficiency of *X. olivacea*, the means of conservation must be developed for this solitary bee which appeared in this work as a synergistic or alternative pollinator of the honey bee within a watermelon farm.

Key words: Foraging behaviour, pollination efficiency, Apis mellifera, Xylocopa olivacea, watermelon, yield.

Introduction

Watermelons (*Citrullus lanatus*) like many other Cucurbit species cannot produce fruits without pollination (Stanghellini *et al.*, 1998). A watermelon plant is monoecious and thus produces both male and female flowers (Adlerz *et al.*, 1966). In order to pollinate the watermelon plant, pollen from the male flowers must be transported to the female flowers by pollen vectors among which insects are the most efficient (Klein *et al.*, 2007). Watermelon in nature is mainly pollinated by bee species that move from flower to flower and distribute pollen (Azo'o *et al.*, 2010). As watermelon farms rely on bees to pollinate the crop, farmers may manage the bee hives to perform this service (Philippe, 1991).

Bee pollination is a vital service for both wild and agricultural systems (Kremen *et al.*, 2004). Without bee pollinators, almost a third of the world's plant species would flower, only to fade and then lie barren (Southwick and Southwick, 1992; Gallai *et al.*, 2009). Bee species are the main service provider along with greater pollinator diversity (Torchio, 1990). Except, bees of the genus *Apis*, all other bee species are known as non-*Apis* bees or wild bees (Aslam *et al.*, 2017).

Utilization of pollinators especially honey bees is considered as one of the cheapest and eco-friendly approach in maximizing the yield of cross-pollinated crops (Free, 1970). Social bee Apis mellifera is ranked first amongst the insect species found to visit and pollinate flowers of several crops such as watermelon (Stanghellini et al., 1998; Klein et al., 2007; Azo'o et al., 2010, 2017). Until recently, honey bees have been so easy to manage and pollination by wild bees has largely been ignored in agricultural systems (Brittain et al., 2013). However, with disease, fire, and competing demands reducing honey bee supplies and increasing the cost of hive rentals, the viability of depending on other bees for pollination has become an important factor (Haubruge et al., 2006). Global decline in honey bee populations, and their limited efficiency in pollinating some crops motivate the search for additional pollinators (Sadeh et al., 2007). Indeed, scientists are discovering more and more that unmanaged wild pollinators could also contribute substantially to crop pollination (Faucon et al., 2002; Potts et al., 2010).

Several unmanaged native bees are widely recognized as important pollinators of a range of wild plants and crop species (Kremen *et al.*, 2004); these include carpenter bees. There are more than 730 species of carpenter bees, *Xylocopa* spp. worldwide. They are robust and large, among the largest bee species known, with some reaching 4.5 cm in length (Hurd, 1978). *X. olivacea* is a species of carpenter bee of the family Apidae and

the subfamily Xylocopinae (Pauly et al., 2015). Individuals have particular nesting behaviour which consists to burrow into hard plant material such as dead wood or bamboo (Pauly et al., 2015). Female has mesosoma totally covered by yellow pubescence and tergum 1 yellow with a mite pouch while male is totally covered by yellow pubescence (Pauly et al., 2018). Previous results showed the positive impact of X. olivacea on the pollination and the increase of seed yields of cultivated legumes such as Phaseolus vulgaris, Pisum sativum, Vigna unguiculata and Cajanus cajan in Burundi (Pauly et al., 2015) and in Cameroon (Tchuenguem et al., 2014a,b; Kengni et al., 2015). This wild bee was found among the main flower-visiting insects and pollinators of cucurbit species such as Cucumis melo in Israel (Sadeh et al., 2007), Luffa aegyptiaca in Ghana (Mensah and Kudom, 2011), Momordica charantia in Western Kenya (Oronje et al., 2012), Cucumeropsis mannii (Azo'o and Messi, 2012) and Citrullus lanatus in Cameroon (Azo'o et al., 2017). Moreover, carpenter bee species are known as the primary pollinators of passion fruit in Brazil (Furlaneto et al., 2011; Giannini et al., 2013). The agronomic significance of this wild bee may even be improving.

For a commercial crop as watermelon, a thorough study of the bee species with a neglected pollination potential appears essential and could allow their knowledge, conservation and optimal use in the biological processes of the host plant production. The present study aimed to determine the foraging and pollination activities of *X. olivacea* that may be valuable as synergistic or alternative pollinator of *A. mellifera* in the watermelon in the study area.

Materials and methods

Study site: The study was carried out in Dengui ($10^{\circ}32^{\circ}55^{\circ}$ 'N 14°14'48"E, 442 meters), a neighborhood of Maroua (Far-North Region, Cameroon). The field experiments took place in a farmer's garden of about 5000 m² during the blooming period of *C. lanatus* for two consecutive cultivating seasons from July until September 2016 and 2017. The climate here is of the Sudano-Sahelian type with two seasons; the dry and rainy seasons. These experiments were conducted during the rainy season; the rainfall recorded stood at a mean of 1004.7 mm and temperatures ranged from 27 to 36 °C.

A. mellifera and X. olivacea behavioral parameters: A transect of 4 m^2 was used by the observer along the edge of the garden. Direct observations inside the transect area were done twice a week (Wednesday and Saturday) on bee species targeted at 9 interval periods: 06:00-06:45 a.m., 07:00-07:45 a.m., 08:00-08:45 a.m., 09:00-09:45 a.m., 10:00-10:45 a.m., 11:00-11:45 a.m., 12:00-12:45 p.m., 01:00-01:45 p.m. and 02:00-02:45 p.m. The mean temperature and hygrometry corresponding to each observation interval were recorded using an indoor/outdoor hygro-thermometer HT 9227. The following foraging parameters were registered: the distribution of the two bee species according to the time intervals, the frequency of each bee floral visits, the density of each bee species per 1000 flowers, the foraging preference or floral product harvested, the duration of visit or the time spent by each species on a flower, the foraging speed or number of flowers visited per minute (Jacob-Remacle, 1989) and the percentage of effective visits of each bee species.

Pollination efficiency of A. mellifera and X. olivacea: The study

of the pollination efficiency of A. mellifera and X. olivacea on watermelon flowers was done daily during 3 weeks. We used random samples of twenty experimental plants for each of the four treatments which were: a control treatment (T_{o}) with bagged female flowers benefited from no insect visit; two similar treatments where flowers were previously opened to A. mellifera (T_1) and X. olivacea (T_2) for a single visit. Protection of female flower buds in these treatments was done on the day before their opening using gauze bags; emerged petals were allowed to accurately detect the imminent blossoming of the correspondent flowers the following day (Azo'o et al., 2017). Female flowers here were opened to A. mellifera and X. olivacea between 07:00-09:00 a.m. After the visit of an individual of a given bee species, each flower was re-bagged to avoid any additional visit of insects. The gauze bag was removed the day after and the young fruit, if any, was flagged for continuous survey until maturity; the unrestricted-visit treatment (T_3) , in which plants were tagged at random with open-pollinated flowers that were freely exposed to the foraging activity of anthophilous insects.

To avoid any bias, only the first female flower opened was considered per watermelon flagged plant according to each treatment. All-female flowers that developed after treatments were removed from each test plant using a scissor, insuring that treated flowers were given an optimum chance for development (Stanghellini *et al.*, 1998).

The number of female flowers that set fruit was recorded in each treatment. Fruits were harvested and weighted at physiological maturity. Subsequently, each fruit was cut into the equatorial part with a kitchen knife, allowing the diameter to be measured using a caliper and counting the mature seeds embedded in the fruit flesh. The pollination efficiency of each bee species was estimated in terms of proportion of female flowers which set fruit, the mean weight of fruits, the mean diameter of fruits harvested and the mean number of mature seeds per fruit compared between treatments as a quantitative and qualitative measure of the bees' pollination success.

Statistical analysis: Data collected were keyed into an Excel sheet and analyzed using SPSS software. The data was subjected to the Student's t-test for the comparison of means between two samples. Correlations were established to study the linear relations between two parameters. One-way Analysis of Variance (ANOVA) and post hoc tests (HSD) of Tuckey Kramer with *P* sets to 0.05 were used for multiple comparisons of means. The mean values were followed by the standard error (SE).

Results

General activity patterns of *A. mellifera* and *X. olivacea* on watermelon flowers: Table 1a and Table 1b showed a divergence in the rhythm of floral activity of the two bee species in 2016 and 2017. In both cases, the number of bee visits was influenced by the time of the day. The floral activity of *X. olivacea* was prominent at dawn; a visitation peak was reached between 07:00-08:00 a.m., which declined to zero by approximately 10 a.m. Meanwhile, the floral activity of *A. mellifera* was effective throughout the daily opening period of the flowers since the dawn (06:00-07:00 a.m.) until around 02:00 p.m. with an important peak observed at 09:00-10:00 a.m. After this peak, the activity decreased with the flower

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Time frames	Abiotic parameters		Number and frequency of bee floral visits				
	Temperature (°C)	RH (%)	Honey bee	FFV	Carpenter bee	FFV	
06:00-06:45 a.m.	27.72	76.43	94	4.51	133	22.28	
07:00-07:45 a.m.	28.95	71.37	213	10.22	347	58.12	
08:00-08:45 a.m.	31.17	67.46	307	14.72	98	16.42	
09:00-09:45 a.m.	32.09	64.21	538	25.80	16	2.68	
10:00-10:45 a.m.	32.97	62.33	408	19.57	3	0.50	
11:00-11:45 a.m.	33.11	59.52	274	13.14	0	0.00	
12:00-12:45 a.m.	34.43	55.78	193	9.26	0	0.00	
01:00-01:45 p.m.	32.65	51.47	58	2.78	0	0.00	
02:00-02:45 p.m.	31.57	48.26	0	0.00	0	0.00	
Total			2085	100	597	100	

Table 1a. Variation of bee visits according to interval times, temperature and hygrometry in 2016

FFV = Frequency of floral visits

Table 1b. Variation of bee visits according to interval times, temperature and hygrometry in 2017

Time frames	Abiotic parameters		Number and frequency of bee floral visits				
	T (°C)	H (%)	Honey bee	FFV	Carpenter bee	FFV	
06:00-06:45 a.m.	27.31	73.71	104	4.47	176	32.77	
07:00-07:45 a.m.	28.07	69.59	274	11.75	297	55.31	
08:00-08:45 a.m.	30.04	65.28	381	16.34	53	9.87	
09:00-09:45 a.m.	31.23	62.01	603	25.87	11	2.05	
10:00-10:45 a.m.	32.16	60.27	458	19.65	0	0.00	
11:00-11:45 a.m.	33.38	58.13	311	13.34	0	0.00	
12:00-12:45 a.m.	33.92	54.24	137	5.88	0	0.00	
01:00-01:45 p.m.	33.11	50.33	63	2.70	0	0.00	
02:00-02:45 p.m.	32.39	47.54	0	0.00	0	0.00	
Total			2331	100	537	100	

FFV = Frequency of floral visits

wilting and closing. The correlations between the rhythm of floral visits and the temperature corresponding to each daily observation were non-significant for *A. mellifera* (r = 0.25; df = 7; P > 0.05 in 2016; r = 0.07; df = 7; P > 0.05 in 2017) and negative and significant for *X. olivacea* (r = -0.75; df = 7; P < 0.001 in 2016; r = -0.86; df = 7; P < 0.001 in 2017). Moreover, our results revealed non-significant correlations between the rhythm of floral visits and the daily variation of the relative humidity for *A. mellifera* (r = 0.33; df = 7; P > 0.05 in 2016; r = 0.37; df = 7; P > 0.05 in 2017) but a significant correlation is the *X. olivacea* floral visits variation which is a function of the variation of the hygrometry according to time intervals (r = 0.68; df = 7; P < 0.001 in 2016; r = 0.75; df = 7; P < 0.001 in 2016; r = 0.75; df = 7; P < 0.001 in 2016; r = 0.75; df = 7; P < 0.001 in 2016; r = 0.75; df = 7; P < 0.001 in 2017).

Table 2 contains results about the foraging parameters of the honey bee and carpenter bee foragers on watermelon flowers in 2016 and 2017. *A. mellifera* and *X. olivacea* were frequent visitors to watermelon flowers where they mostly foraged nectar than pollen. The frequency of *A. mellifera visits* (77.74 % in 2016 and 81.28 % in 2017) was up than that of *X. olivacea* (22.26 % in 2016 and 18.72 % in 2017). Over 100 floral visits studied, 100 % of *X. olivacea* visits were devoted to nectar harvesting during the two experiments, while 92 % and 90 % correspond to the equivalent values dedicated to *A. mellifera* in 2016 and 2017. During their floral activity, both honey bees and carpenter bees came into contact with stigmas when visiting female flowers and anthers on male flowers; that are why the percentage of effective visits was always 100 % for each bee species.

The mean density of forager considerably varied and increased from *X. olivacea* to *A. mellifera* (Table 2); the difference was significant between the two means in 2016 (t = 21.76; df = 98; P < 0.001) and in 2017 (t = 27.32; df = 98; P < 0.001). The density

of foragers also varied for a given bee species according to the season, but the difference between the mean values was overall not significant. The foraging speed followed the same evolution as a function of bee species with significant difference between *X*. *olivacea* and *A. mellifera* in 2016 (t=7.52; df=98; P < 0.001) and in 2017 (t=8.11; df=98; P < 0.001). Finally, the difference of the mean duration of visit between foragers on male flowers and for a given bee species according to the season was not significant; meanwhile, the difference of the mean duration of *A. mellifera* visit was significant between the sex of the flower in 2016 (t=5.38; df=98; P < 0.001) and 2017 (t=6.06; df=98; P < 0.001) with the highest value on female flowers than on male flowers; the difference of the mean duration of visit between *A. mellifera* and *X. olivacea* was significant too on female flowers in 2016 (t=8.16; df=98; P < 0.001) and 2017 (t=7.54; df=98; P < 0.001).

Efficiency of different pollination treatments regarding watermelon yields: Table 3 highlights the variation of the fruit set rate, the mean fruit weight, the mean fruit diameter and the mean number of mature seeds per fruit as a function of years and for a given year according to different treatments. No statistical difference was found during both years between all the four studied parameters from the same level treatment. On the contrary, the difference of the fruiting rate ($F_{3,76} = 13.24$; P < 0.001 in 2016 and $F_{3,76} = 11.73$; P < 0.001 in 2017); the fruit weight ($F_{3,76} = 21.92$; P < 0.001 in 2016 and $F_{3,76} = 18.47$; P < 0.001 in 2017); the mean diameter of a fruit ($F_{3,76} = 12.71$; P < 0.001 in 2016 and $F_{3,76} = 14.68$; P < 0.001 in 2017) and the mean number of seeds per fruit ($F_{3,76} = 20.22$; P < 0.001 in 2016 and $F_{3,76} = 16.74$; P < 0.001 in 2017) were significant between the four treatments and for a given parameter yearly. It results from the control or no bee visit treatment (T_0) that all-female flowers are aborted and do not produce edible fruit and then any seeds in both years 2016 and

Parameters	20	016	20	17
	X. olivacea	A. mellifera	X. olivacea	A. mellifera
Foraging preference $(n = 100)$	100Ne; 0Po	91Ne; 9Po	100Ne; 0Po	90Ne; 10Po
Frequency of floral visits (%)	22.26	77.74	18.72	81.28
Density of foragers $(n = 50)$	97 ± 11a	$442\pm27b$	$88 \pm 9a$	$476\pm31b$
Foraging speed $(n = 50)$	$11.17 \pm 0.9a$	$7.62\pm0.73b$	$9.98\pm0.76a$	$7.01\pm0.64b$
Percentage of effective visits ($n = 100$	0) 100 %a	100 %a	100 %a	100 %a
Duration of bee visit $\sqrt[n]{n} (n = 50)$) $2.93 \pm 0.46a$	$4.01\pm0.34a$	$2.78\pm0.33a$	$3.91\pm0.42a$
$\mathcal{Q}(n=50)$) $3.07 \pm 0.73a$	$6.73 \pm 0.62b$	$2.98 \pm 0.37a$	$6.48\pm0.64b$

Table 2. Foraging parameters of *Apis mellifera* and *Xylocopa olivacea* on watermelon flowers

Legend: Means ± SE within a line and/or a column (duration of visit) followed by the same letter are not significantly different. n = sample size; Ne = nectar; Po = pollen; \Diamond = staminate flower; \Diamond = pistillate flower

Treatments	eatments Fruiting rate (%)		Fruit weight (g)		Mean number of mature seeds/fruit		Mean fruit diameter (cm)	
-	2016	2017	2016	2017	2016	2017	2016	2017
T	0a	0a	0a	0a	0a	0a	0a	0a
T ₁	$40\pm11b$	$45\pm13b$	$690\ \pm 480a$	$570\ \pm 459a$	$57.76\pm20.47a$	$63.03\pm21.36a$	$4.37\pm4.03a$	$4.93\pm4.03a$
T,	$85\pm9c$	$85 \pm 11c$	$2420\pm1260c$	$2370 \pm 1420 \text{c}$	$234.10\pm27.60c$	$228.9\pm22.41c$	$14.76\pm5.44b$	$13.08\pm4.03b$
T ₃	$90\pm 3c$	$90\pm4c$	$5360 \pm 1050 d$	$5730 \pm 1310 d$	$503.94 \pm 81.67 d$	$521.17 \pm 97.59d$	$24.81\pm7.58c$	$25.34\pm8.03c$
ANOVA	F = 13.24	F = 11.73	F = 21.92	F = 18.47	F = 20.22	F = 16.74	F = 12.71	F = 14.68
	df = 3, 76	<i>df</i> =3,76	<i>df</i> = 3, 76	df = 3, 76	df = 3, 76	<i>df</i> =3,76	<i>df</i> =3,76	df = 3, 76
	P < 0.001	P < 0.001	P < 0.001	P < 0.001	P < 0.001	P < 0.001	P < 0.001	P < 0.001

Mean values in the same column (treatments) or in the same line (for a given parameter as function of the year) but with different letters vary significantly (P < 0.05)

2017. The fruiting rate, the fruit weight, the fruit diameter and the mean number of seeds per fruit were substantially increased for female flowers visited by X. olivacea (T_2) than those visited by A. mellifera (T_1) at equal pollination level. The differences were significant between mean treatments from pollination efficiency of the two visitors $(T_1 \text{ and } T_2)$ for these parameters in 2016 and 2017. The low fruiting rate obtained in T_1 justified the insignificant difference observed between this treatment and T₀ with respect to the mean fruit weight, the mean fruit diameter and the average number of mature seeds. Indeed, a single visit of A. mellifera seemed clearly insufficient to provide young fruits which could form a marketable value and a high resistance to the destructive activity of Tephritidae found during the field experiment. Finally, female flowers from T₃ produced numerous and bigger fruits with several mature seeds than female flowers issued from T_0 , T_1 , and T_2 which were under bee visits restriction. Overall, the individual activity of X. olivacea allowed for the best fruit set rate, mean fruit weight, mean fruit diameter, and mean number of mature seeds compared with those from A. mellifera one female flower visit, although the latter is more frequent in watermelon community than the former. Moreover, the results are maximized when these two bee species, the most prominent of the browsers in the study site work synergically without any external influence on their foraging and pollination activity.

Discussion

Honey bee (*A. mellifera*) and Carpenter bee (*X. olivacea*) were prominent visitors of watermelon flowers in our study site; that is why their foraging and pollinating activities were easily comparable. In the field study, the frequency of *A. mellifera* individuals was up than that of *X. olivacea* and all the measured foraging parameters were more significant for the first bee *A. mellifera* which is a social bee species while *X. olivacea* is solitary. The floral resources constitute both bee species which is an important provision for the perfect growth of their offspring. Unlike carpenter bees, honey bees have large population sizes inside their colony and workers are more empowered to recruit several congeners for exploiting an interesting food resource in terms of nectar and/or pollen (Louveaux, 1984). For this last reason, the density of *A. mellifera* was important than that of *X. olivacea* in our study.

The recrudescence of the carpenter bee activity in the morning appears to be a specific trait of nutrition in this insect group. The intense morning activity of A. mellifera on watermelon flowers is synchronized with the higher nectar secretion which occurs 2 to 3 hours after flower opening at dawn (Cervancia and Bergonia, 1991). Indeed, when the watermelon flowers are open, the pollen is dehiscent and the stigma receptive for at least two hours (Philippe, 1991) and a large percentage of fruit results from the deposit of pollen on the stigma between 09:00 and 10:00 a.m. (Adlerz 1966). However, the reduction in honey bee visitation rates after 09:00 and 10:00 a.m. could be explained by the depletion of floral resources and/or adverse abiotic factors. These results are consistent with the findings of Polatto et al. (2014) with regard to the influence of abiotic factors and floral resources available in the daily foraging activity of bees. A. mellifera and X. olivacea were essentially nectarophagous on C. lanatus flowers. According to Delaplane and Mayer (2000), nectar is the primary objective of bees that visit flowers of Cucurbitaceae species.

Observed variations in the mean duration of a bee visit according to the sex of flowers are known with Cucurbitaceae species. For instance in California (U.S.A.), the mean duration of a visit of *A. mellifera* on watermelon varied from 5.7 sec. on male flowers to 8.0 sec. on female flowers (Adlerz, 1966); In Constantine (Algeria), Benachour and Louadi (2011) showed that on the flowers of cucumber (*Cucumis sativus*), the mean duration of visit for nectar harvesting varied from female flowers to male flowers according to bee species: 8.4 and 7.2 sec. for *A. mellifera*; 10.5 and 7.5 sec. for *Ceratina cucurbitina*; 11.1 and 6.8 sec. for *Meliponula piliden*. Kaziev and Seidova (1965) reported that female flowers of Cucurbitaceae secreted a lot of energetic nectar than male flowers. The difference in the average duration of a bee visit on both types of flowers is linked to the optimal foraging principle (Frisch, 1969; Heyneman, 1983), which guarantees a net energy gain to honey bees.

Pollinators play a key role in increasing crop yields (Morandin and Winston, 2006). The knowledge of insect pollinators' diversity for a plant is important so that pollination could be made possible in the absence of a particular insect species (Anoosha *et al.*, 2018; Campbell *et al.*, 2019). Watermelon pollination depends on several insect species. Previous results indicated *A. mellifera* and *X. olivacea* among the pollinators of several Cucurbitaceous species, with honey bee as the most representative (Sadeh *et al.*, 2007; Mensah and Kudom, 2011; Azo'o *et al.*, 2010, 2012, 2017). The pollinator diversity on watermelon reduces the risk of lack of pollination service in absence of one insect species during critical period of crop flowering (Kremen *et al.*, 2004).

It is well known that the populations of native bees and honey bees are generally in decline, threatening food production (Kevan and Phillips, 2001). According to our findings, the absence or lack of pollinators in watermelon cultivation represents a major threat to fruit production of this crop. In the watermelon farm, no pollinators represent no-pollination and no fruit set. However, the floral activity of pollinating insects, especially bees, is essential to obtain improved fruit and seed yields of watermelon.

If *A. mellifera* is considered the most important pollinator of watermelon depending on its pollination intensity due to the large size of its colony and the ability of this bee species to recruit several workers during a foraging trip (Frisch, 1969), *X. olivacea* appeared as a more efficient pollinator on an individual basis than *A. mellifera*. Indeed, mean treatments for fruiting rate, fruit weight, fruit diameter and mean of mature seeds per fruit of watermelon in 2016 and 2017 were important for the carpenter bee than the honey bee at equal pollination level. According to Mensah and Kudom (2011), the size of *X. olivacea* has been reported to play a positive role in crop pollination. Furthermore, the low values of the mean duration of a floral visit determine the ability of this carpenter bee species to move swiftly from flower to flower across large areas and suggest that it can cross-pollinate watermelon flowers efficiently.

Apart from the present work-study, other field experiments have shown the importance of other bee species in the pollination of certain crops compared with the honey bee via their pollinating efficiency. For instance, in New-York (eastern U.S.), honey bees (A. mellifera) and two native bee species Bombus impatiens and Peponapis pruinosa were the most abundant species that pollinate Cucurbita pepo. Research has shown that B. impatiens is an efficient pollinator on individual basis, depositing more pollen per visit and needing fewer overall visits to a flower to produce a large pumpkin fruit compared with equivalent visits by either A. mellifera or P. pruinosa (Artz and Nault, 2011). Also, in the U.S.A., managed honey bees A. mellifera were the most common visitor, but numerous other pollinators (e.g., Agapostemon splendens Lepeletier, Campsomeris plumipes var. fossulana Fabricius) were found to be abundant in field experiment (Campbell et al., 2019). In North Carolina, Stanghellini et al. (1998) have shown that watermelon fruit from

bumble bee-visited flowers has consistently higher seed sets than did those visited by honey bees when compared at equal bee visit numbers. In Chania region (Greece), the wild bee genus Lasioglossum was observed to be the main alternative pollinator to honey bees of watermelon crops (Garantonakis et al., 2016). In Yaounde (Centre region, Cameroon), Meliponula erythra was the main pollinator of Dacryodes edulis (Tchuenguem et al., 2001) and wild bees were prominent pollinators on Zea mays (Tchuenguem et al., 2002). Previous findings in Maroua (Far-North region, Cameroon) showed that two wild bee species Eucara macrognatha and Tetralonia fraterna (Hymenoptera: Apidae) have outnumbered honey bee A. mellifera on Okra (Abelmoschus esculentus) pollination (Azo'o et al., 2011) and were more efficient pollinators than this bee species (Azo'o et al., 2012). Hence, maintaining biodiversity in agricultural ecosystems could provide unrecognized benefits (Carvalheiro et al., 2011; Brittain et al., 2013). It is therefore recommended to preserve or restore natural vegetation surrounding watermelon cultivated areas to attract and keep the bees in or near those areas. Such measures may ensure that bees are available to visit the flowers whenever needed.

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References

- Adlerz, W.C. 1966. Honey bee visit numbers and watermelon pollination. *J. Econ. Entomol.*, 59: 28-30.
- Anoosha, V.S., H.D. Saini, Kaushik and S. Yadav, 2018. Diversity of insect pollinators on medicinal tree, Baheda (*Terminalia bellirica* Roxb.) in southern Haryana. *J. Entomol. Zool. Stud.*, 6(4): 429-432.
- Artz, D.R. and B.A. Nault, 2011. Performance of *Apis mellifera*, *Bombus impatiens*, and *Peponapis pruinosa* (Hymenoptera: Apidae) as pollinators of pumpkin. J. Econ. Entomol., 104(4): 1153-1161.
- Aslam, S., M.A. Rafi and A. Zia, 2017. Non-Apis bees of the family Apidae (Hymenoptera) from Potohar region of Pakistan. J. Entomol. Zool. Stud., 5(2): 6-12.
- Azo'o, E.M., J. Messi, F.F-N. Tchuenguem, J.L. Tamesse, S. Kekeunou and J.B. Pando, 2010. Foraging behaviour of *Apis mellifera adansonii* and its impact on pollination, fruit and seed yields of *Citrullus lanatus* at Nkolbisson (Yaoundé, Cameroon). *Cam. J. Exp. Biol.*, 6(1): 41-48.
- Azo'o, E.M., F.F-N. Tchuenguem and J. Messi, 2011. Influence of the foraging entomofauna on okra (*Abelmoschus esculentus*) seed yields. *Int. J. Agric. Biol.*, 13: 761-765.
- Azo'o, E.M. and J. Messi, 2012. Yield responses of *Cucumeropsis mannii* (Cucurbitaceae) to the presence or the absence of the insect foraging activity at Nkolbisson in Cameroon. J. Anim. Plant Sci., 13(3): 1791-1799.
- Azo'o, E.M. A. Madi, F.F-N. Tchuenguem and J. Messi, 2012. The importance of a single floral visit of *Eucara macrognatha* and *Tetralonia fraterna* (Apidae) in the pollination and the yields of *Abelmoschus esculentus* in Maroua, Cameroon. *Afr. J. Agric. Res.*, 7(18): 2853-2857.
- Azo'o, E.M., F.F-N. Tchuenguem and J. Messi, 2017. Biological diversity of the entomofauna associated with *Citrullus lanatus* (Cucurbitaceae) flowers and assessment of its impact on yields. *J. Entomol. Zool. Stud.*, 5(5): 810-815.

- Benachour, K. and K. Louadi, 2011. Comportement de butinage des abeilles (Hymenoptera : Apidae) sur les fleurs mâles et femelles du concombre (*Cucumis sativus* L.) (Cucurbitaceae) en région de Constantine (Algerie). Ann. Soc. Entomol. Fr., 47: 63-70.
- Brittain, C., N. Williams, C. Kremen and A.M. Klein, 2013. Synergistic effects of non-*Apis* bees and honey bees for pollination services. *Proc. Royal Soc.* B., 280: 2012-2767.
- Campbell, J.W., C. Stanley-Stahr, M. Bammer, J.C. Daniels and J.D. Ellis, 2019. Contribution of bees and other pollinators to watermelon (*Citrullus lanatus* Thunb.) pollination. J. Apic. Res., 58(4): 597-603.
- Carvalheiro, L.G., R. Veldtman, A.G. Shenkute, G.B. Tesfay, C.W.W. Pirk, J.S. Donaldson and S.W. Nicolson, 2011. Natural and withinfarmland biodiversity enhances crop productivity. *Ecol. Lett.*, 14: 251-259.
- Cervancia, C.R. and E.A. Bergonia, 1991. Insect pollination of Cucumber (*Cucumis sativus* L.) in the Philippines. *Acta Hort.*, 6: 278-282.
- Delaplane, K.S. and D.F. Mayer, 2000. *Crop pollination by bees*. CABI Edition. London and New York.
- Faucon, J.P., L. Mathieu, M. Ribière, A.C. Martel, P. Drajnudel, S. Zeggane, C. Aurieres and M.F.A. Aubert, 2002. Honey bee winter mortality in France in 1999 and 2000. *Bee World*, 83: 14-23.
- Free, J.B. 1970. *Insect pollination of crops*, 2nd edition, London Academic Press, London.
- Frisch, K.V. 1969. Vie et mœurs des abeilles. A. Michel (ed.), Paris.
- Furlaneto, F.P.B., A.N. Martins, M.S.T. Esperancini, A.A. Vidal and F. Okamoto, 2011. Production cost of yellow passion fruit (*Passiflora* edulis). Rev. Bras. Frutic., 33: 441-446.
- Gallai, N., J.M. Salles, J. Settele and B.E. Vaissière, 2009. Economic valuation of the vulnerability of world agriculture confronted with pollinator decline. *Ecol. Econ.*, 68: 810-821.
- Garantonakisa, N., K. Varikou, A. Birouraki, M. Edwards, V. Kalliakaki and F. Andrinopoulos, 2016. Comparing the pollination services of honey bees and wild bees in a watermelon field. *Scientia Hort.*, 204: 138-144.
- Giannini, T.C., A.L. Acosta, C.I. Silva, P.E.A.M Oliveira, V.L. Imperatriz-Fonseca and A.M. Saraiva, 2013. Identifying the areas to preserve passion fruit pollination service in Brazilian tropical savannas under climate change. *Agric. Econ. & Environ.*, 171: 39-46.
- Haubruge, E., B.K. Nguyen, J. Widart, J.P. Thomé, P. Fickers and E. Depauw, 2006. Le dépérissement de l'abeille domestique, *Apis mellifera* L., 1758 (Hymenoptera : Apidae) : faits et causes probables. *Notes Fauniques de Gembloux*, 59(1): 3-21.
- Heyneman, A. 1983. Optimal sugar concentrations of floral nectars dependence on sugar intake efficiency and foraging costs. *Oecologia* 60: 198-213.
- Hurd, P.D. 1978. An annotated catalog of the carpenter bees (genus *Xylocopa* Latreille) of the Western Hemisphere (Hymenoptera, Anthophoridae).Washington, DC, Smithsonian Institution Press.
- Jacob-Remacle, A. 1989. Comportement de butinage de l'abeille domestique et des abeilles sauvages dans des vergers de pommiers en Belgique. *Apidologie*, 20: 271-285.
- Kaziev, I.P. and S.S. Seidova, 1965. The nectar yield of flowers of some Cucurbitaceae under Azerbaidjan conditions. XXth International Beekeeping Congress, Bucharest, pp. 364-366.
- Kengni, B.S., A. Ngakou and F.F-N Tchuenguem, 2015. Pollination and yield attributes of (cowpea) *Vigna unguiculata* L. Walp. (Fabaceae) as influenced by the foraging activity of *Xylocopa olivacea* Fabricius (Hymenoptera: Apidae) and inoculation with Rhizobium in Ngaoundere, Cameroon. *Int. J. Agron. Agric. Res.*, 6(2): 62-76.
- Kevan, P.G. and T. Phillips, 2001. The economics of pollinator declines: assessing the consequences. *Conserv. Ecol.*, 5: 8.
- Klein, A.M., B.E. Vaissière, J.H. Cane, I. Steffan-Dewenter, S.A. Cunningham, C. Kremen and T. Tscharntke, 2007. Importance of pollinators in changing landscapes for world crops. *Proc. Royal Soc.* B, 274: 303-313.
- Kremen, C., N. Williams and W. Thorp, 2004. Crop pollination from

native bees at risk from agricultural intensifications. Proc. Natl. Acad. Sci., 99: 16812-16816.

- Louveaux, J. 1984. L'abeille domestique dans ses relations avec les plantes cultivées. In: *Pollinisation et production végétales*, Pesson P. & Louveaux J. (ed.), INRA (eds.), Paris. p. 527-555.
- Mensah, B.A. and A.A. Kudom, 2011. Foraging dynamics and pollination efficiency of *Apis mellifera* and *Xylocopa olivacea* on *Luffa aegyptiaca* (Cucurbitaceae) in Southern Ghana. J. Pollinat. Ecol., 4(5): 34-38.
- Morandin, L.A. and M.L. Winston, 2006. Pollinators provide economic incentive to preserve natural land in agroecosystems. Agr. Ecosyst. Environ., 116(8): 289-292.
- Oronje, M.L., M. Hagen, M. Gikungu, M. Kasina and M. Kraemer, 2012. Pollinator diversity, behaviour and limitation on yield of karela (*Momordica charantia* L., Cucurbitaceae) in Western Kenya. *Afr. J. Agric. Res.*, 7(11): 1629-1638.
- Pauly, A., B. Nzighidahera, C. Eardley, L. Ndayikeza, A. Mpawenimana and B. Habonimana, 2015. Les abeilles du genre *Xylocopa* Latreille (Hymenoptera: Apoidea: Apidae) au Burundi, de bons pollinisateurs de légumineuses. *Belg. J. Entomol.*, 25: 1-27.
- Pauly, A., Z.A. Hora, D.N. Wayessa and K. Amberbir, 2018. The *Xylocopa* Latreille, 1802, of Ethiopia (Hymenoptera: Apidae). *Belg. J. Entomol.*, 62: 1-30.
- Philippe, J.M. 1991. La pollinisation par les abeilles: Pose des colonies dans les cultures en floraison en vue d'accroître les rendements des productions végétales. EDISUD, Aix-en-Provence.
- Polatto, L., J. Chaud-Netto and V. Alves-Junior, 2014. Influence of abiotic factors and floral resource availability on daily foraging activity of bees. J. Insect Behav., 1-20.
- Potts, S.G., S.P.M. Roberts, R. Dean, G. Marris, M. Brown, R. Jones and J. Settele, 2010. Declines of managed honeybees and beekeepers in Europe. J. Apic. Res., 49: 15-22.
- Sadeh, A., A. Shmida and T. Keasar, 2007. The carpenter bee *Xylocopa pubescens* as an agricultural pollinator in greenhouses. *Apidologie*, 38: 508-517.
- Southwick, E.E. and L. Southwick, 1992. Estimating the economic value of honey bee (Hymenoptera, Apidae) as agricultural pollinators in the United States. *J. Econ. Entomol.*, 85: 621-633.
- Stanghellini, M.S., J.T. Ambrose and F.R. Schultheis, 1998. Seed production in watermelon: a comparison between two commercially available pollinators. *Horti. Sci.*, 33: 28-30.
- Tchuenguem, F.F.N., J. Messi and A. Pauly, 2001. Activité de *Meliponula* erythra sur les fleurs de *Dacryodes edulis* et son impact sur la fructification. *Fruits*, 56(3): 179-788.
- Tchuenguem, F.F.N., J. Messi and A. Pauly, 2002. L'activité de butinage des Apoïdes sauvages (Hymenoptera: Apoidea) sur les fleurs de maïs à Yaoundé (Cameroun) et réflexion sur la pollinisation des graminées tropicales. *Biotech. Agron. Soc. Environ.*, 6(2): 87-98.
- Tchuenguem, F.F-N., T.S. Fameni and D. Bruckner, 2014a. Foraging and pollination behaviour of *Xylocopa olivacea* (Hymenoptera: Apidae) on *Phaseolus coccineus* (Fabaceae) flowers at Ngaoundéré (Cameroon). *Int. J. Trop. Insect Sci.*, 34(2): 127-137.
- Tchuenguem, F.F-N., J.B. Pando and J.L. Tamesse, 2014b. Pollination efficiency of *Xylocopa olivacea* (Hymenoptera: Apidae) on *Cajanus cajan* (Fabaceae) flowers at Yaoundé, Cameroon. *Int. J. Trop. Insect Sci.*, 34(2): 138-148.
- Torchio, P.F. 1990. Diversification of pollination strategies for U.S. crops. *Environ. Entomol.*, 19: 1649-1656.

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