METABARCODING IN DIET ASSESSMENT OF ADULT PARASITOID SPECIES, Dolichogenidea metesae (HYMENOPTERA: BRACONIDAE) TOWARDS ITS CONSERVATION AND MANAGEMENT IN THE OIL PALM PLANTATION

FUAT SALBI^{1,2}; AQILAH SAKINAH BADRULISHAM¹; NUR AZURA ADAM²; IZFA RIZA HAZMI¹ and SALMAH YAAKOP^{1*}

ABSTRACT

The abundance of parasitoids in the oil palm plantation is greatly influenced by the host and food availability. However, studies on the main diet or food sources of the adult parasitoid wasp, Dolichogenidea metesae of the oil palm pest, Metisa plana is still lacking. This study investigated the diet of adult D. metesae sampled from Good Agricultural Practices (GAP) and non-GAP plantations through metagenomic analysis by using a trnL marker. A total of 761 145 food plant trnL sequences were generated from 12 samples of the parasitoids collected from four plantations. No significant differences (p=0.683) were observed for diversity (Shannon-Wiener index (H') of 0.12-1.22, p=0.68) among the total 62 OTU's of the food plant species when compared between the GAP and Non-GAP plantations. Polygonaceae was the most abundant plant family (59.95%), followed by Asteraceae (10.15%) and Acanthaceae (7.51%). The Antigonon leptopus, Paraprenanthes oligolepis, Asystasiella neesiana and wild banana (Musa itinerans) were among the most dominant plant species consumed by adult D. metesae. We recommend that these dominant food plant species for the D. metesae should be planted widely in the oil palm plantations to enhance the population abundance of D. metesae for successful Integrated Pest Management (IPM) practices.

Keywords: DNA, food source, genetic, natural enemy, trnL.

Received: 20 December 2021; Accepted: 5 September 2022; Published online: 7 November 2022.

INTRODUCTION

Dolichogenidea metesae (Hymenoptera: Braconidae) is the predominant species of parasitoids that parasitises on a gregarious pest, *Metisa plana* (Psychidae: Lepidoptera) (Halim *et al.*, 2017; 2019). As a primary parasitoid of the *M. plana*, *D. metesae* seeks to infest the third to sixth instar of the host insect, and this predilection has been utilised to

become one of the most effective natural or biological control methods for major pests of agricultural crops under the Integrated Pest Management (IPM) in a sustainable ecosystem (Salmah *et al.*, 2006). The longevity and fecundity of the parasitoids are influenced by the availability and type of their food sources (Liu and Ueno, 2012). Thus, the biological data on the *D. metesae*, especially on its diet, food source and consumption habits in the natural ecosystem would be invaluable for a better understanding of the intricate association between the parasitoids and their hosts, the pests, pollinators and other associates in the ecosystem (Fahimee *et al.*, 2021).

The destruction of ground vegetation deprives food and shelter for adult parasitoids, therefore, suitable agricultural landscape and ecological infrastructure are important for the best habitat

¹ Centre for Insect Systematics, Department of Biological Science and Biotechnology, Faculty of Science and Technology, Universiti Kebangsaan Malaysia, 43600 Bangi, Selangor, Malaysia.

² Department of Plant Protection, Faculty Agriculture, Universiti Putra Malaysia, 43400 Serdang, Selangor, Malaysia.

^{*} Corresponding author e-mail: salmah78@ukm.edu.my

management under the natural biological control system (Landis et al., 2000). Balmer et al. (2013) provided additional evidence regarding the importance of flowering weeds to the parasitoids by highlighting the effectiveness of flowering plants as intercropping plants or as companion plants that supply bountiful and nutritious resources for the natural enemies of the crop pests, including the parasitoids. Studies by Norman and Basri (2010) reported that Asystasia gangetica, a weed species that grew naturally in the oil palm plantations had influenced the population of the parasitoids. However, there is no evidence to prove the efficacy of all available food plants including the flowering weeds to supply their diet or to provide a host or conducive environment for the parasitoids.

The implementation of Good Agriculture Practice (GAP) has had a positive impact towards environmental sustainability in Malaysia (Tan et al., 2009) and in neighbouring countries such as Thailand and Indonesia (Apriani et al., 2020; Saswattecha et al., 2015). The Roundtable on Sustainable Palm Oil (RSPO), a global certification standard for sustainable palm oil, requires the implementation of GAP as part of the criteria in getting the certification (Morley, 2015). GAP principles emphasize on limiting the usage of chemical inputs and minimising the detrimental environmental impacts of farming operations through the implementation IPM (Norman et al., 2019). As a result, IPM practices such as planting beneficial plants and weed management are used in combating the oil palm pest, M. plana (Wood and Norman, 2019a). Failure to follow the GAP code of practice (CoP) had resulted in a high use of chemical pesticides, which had a very significant adverse impact on weed diversity (Aini et al., 2007). According to Möller et al. (2021), the abundance of parasitoids was higher in plots with a good ground cover compared to plots treated with herbicides. Wood and Norman (2019b) and Wood et al. (1974) also reported that the application of chemicals had consequently disrupted the natural enemies of *M. plana* in the plantation habitats.

Despite some previous successful studies demonstrating the relationship between flowering plants and parasitoids, e.g., Cassia cobanenasis, Euphorbia heterophylla, Antigonon leptopus and Turnera subulata (Siti Ramlah et al., 2005; Wood and Norman, 2019a), relatively very few studies had been undertaken to validate that these plants were consumed by the parasitoids. To our knowledge, there is no direct study investigating the various plant species foraged or consumed in the diet of adult D. metesae, the main parasitoid of M. plana in the oil palm plantation. Metabarcoding is now widely used in identifying insect diet. By using *trnL* marker, Fahimee et al. (2021) successfully revealed 262 plant species under 70 families foraged by stingless bee, Heterotrigona itama. This information benefits meliponiculture industry by recommending several food plant species to be planted such as *Momordica charantia, Melastoma* sp. and *Cucumis sativa.* According to Mallott *et al.* (2018), *trnL* yielded more sequences, higher resolution taxonomic identifications and identified a higher number of families in the diet of wild white-faced capuchins, *Cebus capucinus* compared to *rcbl.*

Therefore, based on the economic importance of D. metesae and M. plana in our oil palm industry (Basri et al., 1995; Halim et al., 2018), the main objective of this study was to identify the different food plant species consumed by adult D. metesae by conducting a metabarcoding analysis of the *D. metesae* diet using a *trnL* marker, thereby confirming the various species of plants foraged and utilised by D. metesae in the wild. The study also investigated how D. metesae diets differed between the GAP and non-GAP oil palm plantations. This information could facilitate towards a better understanding on the specific ecosystem components that could influence the introduction of *D. metesae* as an effective biological control of the main pests of oil palm as part of the IPM strategies.

MATERIALS AND METHODS

Sampling of Dolichogenidea metesae

Three *D. metesae* individuals (three replicates) were sampled from each plantation across Peninsular Malaysia, comprising two GAP plantations (certified with RSPO or MPOB CoGAP) and two non-GAP plantations, giving a total of 12 samples (n=12) from four sampling sites. The sampling sites were planted with specific beneficial plant species and had a history of *M. plana* infestation (*Table 1*). Samples of hymenopteran parasitoids were collected using a malaise trap and preserved in 90% ethanol for one year, from March 2020 to February 2021. All the specimens of *D. metesae* were collected and identified based on Norman *et al.* (2017) and Achterberg (1993), while only several individuals were used for the molecular analysis.

DNA Extraction

The DNA was extracted from the adults of *D. metesae* collected from four plantations with three biological replications representing each plantation. The DNA of 12 individuals of *D. metesae* were extracted from the whole body individually, using the NucleoSpin® DNA Insect protocol (Machery-Nagel, Germany) (Halim *et al.*, 2018). Each sample was surface sterilised before molecular analysis was proceeded by submersing the insects in 70% ethanol and distilled water for 2-3 s for 3 times (Avanesyan, 2014; Cooper *et al.*, 2016).

TABLE 1. LIST OF Doncnogentaea metesae SAMPLES USED FOR METABARCODING ANALISIS						
Sample code	Location	Grid	GAP status	Beneficial plant species		
Plantation A	Peninsular Malaysia. Pahang: Gambang	3.623169771292426, 103.13824123987163	GAP	Turnera sp. Cassia cobanensis		
Plantation B	Peninsular Malaysia. Perak: Slim River	3.6949601778100867, 101.48563129489453	GAP	Antigonon leptopus		
Plantation C	Peninsular Malaysia. Pahang: Kuantan	3.673757257494192, 103.14538126430287	Non-GAP	Antigonon leptopus Turnera sp.		
Plantation D	Peninsular Malaysia. Johor: Segamat	2.4408761993796264, 102.6866661993966	Non-GAP	Antigonon leptopus Turnera sp.		

TABLE 1. LIST OF Dolichogenidea metesae SAMPLES USED FOR METABARCODING ANALYSIS

Library Preparation

Preparation of the library was conducted by using 2-step PCR. The first PCR was performed using locus-specific sequence primers (trnL) with overhang adapters: g-A49425-added overhang (5'-TCGTCGGCAGCGTCAGATGTGTATAA GAGACAG GG CAATCCTGAGCCAA-3') and h-B49466-added overhang (5'-GTCTCGTGGGCTC GGAGATGTGTATAAGAGACAGCCATTGAG TCTCTGCACCTATC-3'). A total of 25.00 µL was prepared for PCR amplification. The mixture consisted of 5.75 µL Q5® High Fidelity DNA Polymerase (New England BioLabs Inc.), 1.25 µL of each forward and reverse primer (10 µM), 15.75 µL of water, nuclease free and 1.00 µL of DNA template. A total of 5.00 uL of DNA template was used for the sample with concentration < 2.5 ng/ μ L. The amplification reactions were performed under the following protocol: initial denaturation of 95°C for 3 min, followed by 30 cycles of denaturation at 98°C for 30 s, annealing at 64°C for 30 s, extension at 72°C for 30 s and a final extension at 72°C for 5 min. The second PCR was conducted to attach dual indices to the amplicon PCR by using Illumina Nextera XT Index Kit v2 following manufacturer's protocols. Even though no qPCR was conducted, however the quality of the libraries was measured using Agilent Bioanalyzer 2100 System by Agilent DNA 1000 Kit and fluorometric quantification by Helixyte Green[™] Quantifying Reagent. The libraries that passed the library QC were subsequently subjected to next-generation sequencing.

Next-Generation Sequencing

The libraries were normalised and pooled according to the protocol recommended by Illumina and then proceeded to sequencing by MiSeq platform using 150 paired-end. The next-generation sequencing was performed by Apical Scientific Sdn. Bhd. The raw data was then processed and analysed according to the bioinformatics analysis. All the read sequences were deposited in the Sequence Read Archive (SRA) of the NCBI (http://www.ncbi.nlm.nih.gov/sra) with the references number SRR20648567 (Plantation A R1), SRR20648580 (Plantation A R2), SRR20648647

(Plantation A R3), SRR20646195 (Plantation B R1), SRR20648581 (Plantation B R2), SRR20648441 (Plantation B R3), SRR20648771 (Plantation C R1), SRR20648869 (Plantation C R2), SRR20648867 (Plantation C R3), SRR20648625 (Plantation D R1), SRR20648622 (Plantation D R2) and SRR20648768 (Plantation D R3).

Bioinformatic Analysis

Sequence adaptors and low-quality reads were trimmed-off from paired-end reads using BBDuk from the BBTools package. After that, by using USEARCH v11.0.667, the forward and backward reads were merged. All sequences (sequenced on the MiSeq platform) with an average quality score of less than 150 bp or more than 600 bp were eliminated from downstream processing (Edgar, 2010). Following that, the reads were aligned with trnL sequences using the NCBI database. Operational taxonomic units (OTUs) were clustered with 97% similarity cut-off using UPARSE v11.0.66 (Edgar, 2013) and a single representative sequence from each OTU was picked at random. Pynast was used to construct a phylogenetic tree against NCBI database (Caporaso et al., 2010a) and QIIME V1.19.1 was used for taxonomic assignment of OTUs (Caporaso et al., 2010b). The rarefaction curve was plotted to determine the sufficiency of sequency depth by using R V3.6.2 (R Core Team, 2018). The heatmap was also created with R software to display the relative abundances of taxa (phyla) using different shades of colour, with darker representing greater values. colours Alpha diversity (Shannon-Wiener, Simpson, CHAO 1) with ANOVA showed pairwise analysis of variance between groups and its significance was annotated on the box plots with *p*-value set to 0.05. Other statistical analysis was also done in R V3.6.1. (R Core Team 2018).

RESULTS

A total of 761 145 plant *trnL* sequences were generated from 12 samples of *D. metesae*, ranging

from 47 893 to 73 893 (*Table 2*). Except for the Plantation C, all samples from Plantations A, B, and D had more than 60 000 reads, with Plantation A (R3) having the most reads (73 893), while Plantation C (R1) had the lowest reads (47 893). Overall, 62 OTU's were observed, with Plantation A (R1) recording the minimum OTU's (16) while Plantation C (R1) recorded the maximum of OTU's (29). The Shannon-Wiener and Simpson index indicated a higher diversity of OTU's from Plantation D (R2) (H' = 1.22, 1-D = 0.68) and the lowest from Plantation D (R3) (H' = 0.12, 1-D = 0.04). Chao 1 estimator abundance showed that Plantation C (R2) has the highest value (28.75) of OTU's, while conversely, Plantation A (R2) recorded the lowest value (11) (*Table 2*).

The rarefaction curves (*Figure 1*) indicate that all the samples are almost reaching asymptote, implying that sufficient sequencing depth has been achieved. The curve rises steeply at first, and then flattens out gradually at 35 000 sequences. The *D. metesae* sample taken from Plantation C (R2) had the most diverse food plant species, since the number of different species detected was higher than in all the other samples, while the *D. metesae* sample from Plantation A (R2), had the least diverse food plant species (*Figure 1*).

Overall, Polygonaceae (59.95%) was the most common family recorded (Table 3), followed by Asteraceae (21.65%) and Acanthaceae (7.51%), while Arecaceae, Campanulacea, Urticaceae and Phyllanthaceae were the least recorded, accounting for less than 1.00% of the total families observed. The heatmap (Figure 2) showed that Polygonaceae and Asteraceae were present in all the D. metesae samples. High abundance of Polygonaceae was recorded from the samples collected from Plantations B, C and D, while Asteraceae was recorded in high abundance in all samples collected from Plantation A only. Meanwhile, Acanthaceae was found mostly in the samples from Plantation D, while very little of Phyllanthaceae and Malvaceae were recorded and could only be found in one D. metesae sample, i.e., from Plantation C (R2) (Figure 2).

As seen in the bar chart (*Figure 3*), *A. leptopus* (Polygonaceae) was the most dominant species of plant food recorded from Plantations B, C and D, accounting for more than half of the total plant species consumed by *D. metesae*. In contrast, *P. oligolepis* was the most dominant plant food recorded in the *D. metesae* samples collected from Plantation A, although it also occurred in all the other 12 samples. Likewise, Plantations A and D seemed to record a higher number of plant species (seven) compared to Plantations B and C, which recorded only four species among the dominant plant foods of *D. metesae*.

According to the Venn diagram (*Figure 4*), both GAP and non-GAP plantations shared 34 OTU's out of 62 OTU's. A Venn diagram is an illustration

of the relationships between and among groups of samples with their own unique OTUs and those that share OTUs in common. Our study shows that alpha diversity metrics as given in the box plot (*Figure 5*) reveals no significant difference (Chao 1, p = 0.23; Simpson, p = 0.45; Shannon, p = 0.68) between the GAP and non-GAP plantations.

DISCUSSION

The metabarcoding analysis has been conducted to obtain the information on the diet of adult *D. metesae*. The results obtained from *D. metesae* samples collected from the GAP and non-GAP plantations are very significant and comparatively valuable in determining the diversity and abundance of *D. metesae* in oil palm plantations adopting different agricultural practices (Saswattecha *et al.*, 2015).

The rarefaction curve (Figure 1) indicates that the sequencing depth has almost reached asymptote, implying that the maximum number of OTU's have been collected in deriving at the conclusion in this study (Cayuela et al., 2015). The family Polygonaceae, belonging to the flowering plant group of angiosperms (Vandebroek *et al.*, 2018), was recorded as the most abundant family (59.95%)found in the D. metesae diet, whereas several other families such as Asteraceae (21.65%), Acanthaceae (7.51%), Musaceae (6.29%) and Poaceae (2.47%) constitute the top five families consumed by the D. metesae. Both families, i.e., Asteraceae and Acanthaceae are classified as weeds (Ali et al., 2021) and are abundantly found in the diet of *D. metesae*. Notably, the families Musaceae and Arecaceae (fruit plant species) also occurred in the D. metesae diet.

The five most abundant plant food species found in the D. metesae samples were A. leptopus, P. oligolepis, A. neesiana, M. itinerans, and Cocos nucifera (Figure 3). The A. leptopus was introduced into the oil palm plantations as a beneficial plant (Hariyadi and Syahlan, 2021) however, P. oligolepis and A. neesiana existed naturally as weeds in Malaysia (Shi et al., 2011). Both species i.e., M. itinerans and C. nucifera were naturally pollinated by animals such as the Long-tongued Fruit Bat (Macroglossus sobrinus), the nectar-feeding bird, Little Spiderhunter (Arachnothera longirostris) and the Indian Palm Squirrel, Funambulus palmarum (Chakravarthy and Thyagaraj, 2012; Liu *et al.*, 2002). However, our own observations indicated that both species i.e., M. itinerans and C. nucifera were very sparse in all the four plantations. Additionally, the C. nucifera that belongs to the Palmae was not suggested to be planted in the plantation due to its potential to attract pest, Red Palm Weevil (RPW), Rhyncophorus ferrugenius (Aman-Zuki et al., 2021). According to Mathews et al. (2007) from a total of 333

Localities	Sample -	Sequences			Chas 1	Shannon-Wiener	Simpson	Evenness
		Before	After	- 010's	Chao-1	(H′)	(1 - D)	(E^HS)
Plantation A	R1	458 2582	66 511	14	15.5	0.72	0.43	15.55
	R2	496 5272	66 931	11	11.0	0.70	0.49	11.29
	R3	593 4902	73 983	18	18.6	0.68	0.35	20.37
		$Mean \pm SD$		14.33 ± 3.51a	$15.03\pm3.82b$	$0.70\pm0.02a$	$0.42\pm0.07a$	$15.74 \pm 4.55 a$
Plantation B	R1	4 394 041	62 425	13	14.00	0.51	0.32	13.79
	R2	4 649 663	62 582	16	18.00	0.67	0.38	19.95
	R3	5 192 020	68 353	24	24.75	0.98	0.56	25.63
		$Mean \pm SD$		$17.67\pm5.69a$	$18.92\pm5.43a$	$0.72\pm0.24a$	$0.42\pm0.123a$	$19.79\pm5.93a$
Plantation C	R1	3 685 638	47 893	24	25.50	0.49	0.22	26.68
	R2	4 592 931	57 067	25	28.75	0.41	0.20	30.76
	R3	4 153 120	56 846	18	24.00	0.68	0.32	22.08
Mean \pm SD		$22.33\pm3.79a$	$26.08\pm2.43a$	$0.52\pm0.14a$	$0.24\pm0.07a$	$26.51\pm4.34a$		
Plantation D	R1	5 411 881	72 422	15	17.00	0.81	0.47	22.45
	R2	4 778 117	63 958	17	17.00	1.22	0.68	17.00
	R3	4 691 056	62 174	17	18.00	0.12	0.04	18.87
		$Mean \pm SD$		16.33 ± 1.16a	$17.33\pm0.58a$	$0.72\pm0.56a$	$0.40\pm0.33a$	$19.44\pm2.77a$

TABLE 2. NUMBER OF OBSERVED OTU'S, ALPHA DIVERSITY INDICES FOR THE PLANT DNA FROM 12 SAMPLES OF Dolichogenidea metesae

Note: Mean \pm SD value with a different letter indicates a significant difference (ANOVA, *p*<0.05).



Figure 1. The rarefaction curve of trnL gene sequence for 12 samples of Dolichogenidea metesae.

plant species in 83 families collected in an oil palm plantation in Malaysia, 279 of them were angiosperms. The abundant flowers of the angiosperms would attract beneficial insects by providing nectar and sugar as a source of food and energy (Heimpel and Jervis, 2005; Tompkins *et al.*, 2010). Our results have also proven that the angiosperm species serve as the main food source for *D. metesae* (Patt *et al.*, 1997) providing for adult longevity and fecundity of the parasitoids (Olson *et al.*, 2005).



Figure 2. Heat analysis at the family level. The darker colour represents a correspondingly greater value of abundance.



Figure 3. Relative abundancy bar chart at the species level. The ten most abundant food plant species recorded in all the 12 samples of Dolichogenidea metesae.

This study provides the first evidence that *A. leptopus* (Polygonaceae) is highly consumed by *D. metesae* as a source of diet (*Figure 3*). Yusdayati *et al.* (2014) reported that parasitoid wasps *Brachymeria carinata* and *D. metesae* could be found in higher abundance on the plant foods *C. cobanensis* and *Turnera* sp. compared to *A. leptopus*. However, our findings revealed that *D. metesae* consumed more *A. leptopus*, while there was no record of *C. cobanensis* and *Turnera* sp. in the diets despite their presence

in the plantation. The use of metabarcoding here is novel since the protocol has not been used previously to confirm the diet of parasitoids. This information is important since nectar, pollen, and water are all necessary for the adult parasitoids to survive and flourish (Damien *et al.*, 2020).

Although previous research works had shown that the common weed locally known as Chinese violet (*A. gangetica*: Achantaceae) could increase parasitoid diversity in the oil palm plantations

TABLE 3. RELATIVE ABUNDANCE OF THE TOP 10 FAMILIES OF PLANTS RECORDED IN THE 12 SAMPLES OF Dolichogenidea metesae

er zennegennen merecue				
Family	Relative abundance (%)			
Polygonaceae	59.95			
Asteraceae	21.65			
Acanthaceae	7.51			
Musaceae	6.29			
Poaceae	2.47			
Lejeuneaceae	1.62			
Arecaceae	0.35			
Campanulaceae	0.07			
Urticaceae	0.05			
Phyllanthaceae	0.04			



Figure 4. Comparison of the Venn diagrams between the GAP and non-GAP plantations.

(Norman and Basri, 2010; Tiong, 1982) our study data had shown that this weed species was not consumed or foraged by *D. metesae* despite its abundant presence in the sampling sites. In contrast, the weedy herbs *P. oligolepis* (Asteraceae) and *A. neesiana* (Acanthaceae) were detected in almost all *D. metesae* samples in this study (*Figure 3*). Based on literature information, the herbaceous species *P. oligolepis*, which is a synonym to *Melanoseris oligolepis* (Shi *et al.*, 2011) and *A. neesiana*, which is a synonym to *Mackaya neesiana* (Wall.) (POWO, 2021), can both be found in Malaysia, although these two weedy species are not yet recorded so far in our oil palm plantations.

The differences in D. metesae diet shown in *Figure 2* could be due to food availability in the respective areas (Folgarait and Gilbert, 1999; Jervis et al., 1992). In an agroecosystem, two important resources (i.e., host and food) for the parasitoids are rarely located within the same site, hence, the adult parasitoids must be able to fly regularly between the host site and the plant food site in order to reproduce successfully (Lewis et al., 1998). Furthermore, a study by Coudrain et al. (2014) found that the parasitoid species richness was higher in areas surrounding the forests than in isolated sites due to higher plant variability (richness) in the former. The importance of plant species such as M. acuminata and C. nucifera, in which the DNA of both have been detected as plant foods of *D. metesae* in this study cannot be overlooked, since both species can be planted as side crops or ornamentals in the oil palm plantations (Mathews et al., 2007). The intercropping strategy has been applied in many



Figure 5. Comparison of the alpha diversity box plots between the GAP (Red) and non-GAP (Blue) plantations. The p-value threshold for significance is set to 0.05.

plantations for many purposes. One of it is to reduce or control the infestation of crop pests such as okra and legumes (Singh *et al.*, 2017), pickerelweed and rice (Xiang *et al.*, 2021) by increasing the population abundance and diversity of parasitoids (Amala and Shivalingaswamy, 2018; Devi *et al.*, 2020), and the second objective is to provide two-in-one crop production (Nchanji *et al.*, 2016; Rocha *et al.*, 2020).

In this study, the molecular data and the mathematical analysis showed that there was no significant difference in plant diversity in the D. metesae diet between the GAP and non-GAP plantations (Figure 5), even though the agricultural practices differed significantly between the two in terms of chemical application (Saswattecha et al., 2015), which would have had very significant impacts on the diversity of weeds and the associated biota (Darras et al., 2019). This might be because in our study, the weed species commonly found in the oil palm plantations such as A. gangetica, Clidemia hirta and Drymaria cordata (Ali et al., 2021; Satriawan and Fuady, 2019) could not be traced or detected in the diet of D. metesae (Figure 2). However, A. gangetica had been documented in other studies as having an important role in the abundance of parasitoids other than D. metesae (Wood and Norman, 2019a). Although this study showed that there was no significant different in terms of plant species consumed by D. metesae in GAP and non-GAP plantations, Fuat et al. (2022) found that the parasitoid abundance was significantly higher in GAP plantations compared to non-GAP plantations.

CONCLUSION

This is the first study to use metabarcoding analysis in determining the plant communities consumed by the *D. metesae* as their food sources in the wild. Overall, the natural diet of *D. metesae* was dominated by the Coral vine or Bride's Tears (*A. leptopus*), a common creeper with abundant pink flowers and rich in nectar. Therefore, it is highly recommended that this beneficial plant should be planted in the oil palm plantations to attract more *D. metesae*, the dominant parasitoid species of *M. plana* as a control measure for bagworm infestation.

ACKNOWLEDGEMENT

The authors would like to express their gratitude to the Department of Crop Protection, Faculty of Agriculture, Universiti Putra Malaysia for enabling us to conduct some experiments in their laboratory. We wish to thank Maimon Abdullah for her useful comments and final editing of the manuscript. This project was funded by the research grants, code ZF-2019-005, GP-K013317-2021 and GP-K013317-2020.

REFERENCES

Achterbeg, C V (1993). Illustrated key to the subfamilies of the Braconidae (Hymenoptera: Ichneumonoidea). *Zool. Verh., 283*: 1-189.

Aini, K; Tan, Y A; Norman, K and Yeoh, C B (2007). Pesticide application in the oil palm plantation. *Oil Palm Bulletin*, 54: 52-67.

Ali, N B M; Karim, M F A; Saharizan, N; Adnan, N S; Mazri, N H; Fikri, N; Amaludin, N and Zakaria, R (2021). Weeds diversity in oil palm plantation at Segamat, Johor. *IOP Conf. Ser. Earth Environ. Sci.*, *756*(1): 012034. DOI: 10.1088/1755-1315/756/1/012034.

Amala, U and Shivalingaswamy, T M (2018). Effect of intercrops and border crops on the diversity of parasitoids and predators in agroecosystem. *Egypt. J. Biol. Pest Control, 28*: 1-4. DOI: 10.1186/s41938-017-0015-y.

Aman-Zuki, A; Ghazali, S Z; Badrulisham, A Z; Hazmi, I R; Nurul Wahida, O and Yaakop, O (2021). Proof on the divergence times of two sympatric species, *Rhynchophorus ferrugineus* and *R. vulneratus* (Coleoptera: Curculionidae) by molecular clock analysis. *J. Entomol. Res. Soc.*, 23(1): 11-26. DOI: 10.51963/jers.v23i1.1851.

Apriani, E; Kim, Y S; Fisher, L A and Baral, H (2020). Non-state certification of smallholders for sustainable palm oil in Sumatra, Indonesia. *Land Use Policy*, 99: 105112. DOI: 10.1016/j. landusepol.2020.105112.

Avanesyan, A (2014). Plant DNA detection from grasshopper guts: A step-by-step protocol, from tissue preparation to obtaining plant DNA sequences. *Appl. Plant Sci.*, *2*(2): 1300082. DOI: 10.3732/apps.1300082.

Balmer, O; Pfiffner, L; Schied, J; Willareth, M; Leimgruber, A; Luka, H and Traugott, M (2013). Noncrop flowering plants restore top-down herbivore control in agricultural fields. *Ecol. Evol.*, *3*(*8*): 2634-2646. DOI: 10.1002/ece3.658.

Basri, M W; Norman, K and Hamdan, A B (1995). Natural enemies of the bagworm, *Metisa plana* Walker (Lepidoptera: Psychidae) and their impact on host population regulation. *Crop Prot.*, *14*(*8*): 637-645. DOI: 10.1016/0261-2194(95)00053-4.

Caporaso, J G; Bittinger, K; Bushman, F D; Desantis, T Z; Andersen, G L and Knight, R (2010a). PyNAST: A flexible tool for aligning sequences to a template alignment. *Bioinformatics*, *26*(2): 266-267. DOI: 10.1093/bioinformatics/btp636.

Caporaso, J G; Kuczynski, J; Stombaugh, J; Bittinger, K; Bushman, F D; Costello, E K; Fierer, N; Peña, A G; Goodrich, J K; Gordon, J I; Huttley, G A; Kelley, S T; Knight, D; Koenig, J E; Ley, R E; Lozupone, C A; Mcdonald, D; Muegge, B D; Pirrung, M; Reeder. J; Sevinsky, J R; Turnbaugh, P J; Walters, W A; Widmann, J; Yatsunenko, T; Zaneveld, J and Knight, R (2010b). QIIME allows analysis of high-throughput community sequencing data. *Nat. Methods*, 7(5): 335-336. DOI: 10.1038/ nmeth0510-335.

Cayuela, L; Gotelli, N J and Colwell, R K (2015). Ecological and biogeographic null hypotheses for comparing rarefaction curves. *Ecol. Monogr.*, *85*(3): 437-455. DOI: 10.1890/14-1261.1.

Chakravarthy, A K and Thyagaraj, N E (2012). The palm squirrel in coconut plantations: Ecosystem services by therophily. *Mammalia*, *76*(2): 193-199. DOI: 10.1515/mammalia-2011-0073.

Cooper, W R; Horton, D R; Unruh, T R and Garczynski, S F (2016). Gut content analysis of a phloem-feeding insect, *Bactericera cockerelli* (Hemiptera: Triozidae). *Environ. Entomo.*, 45(4): 938-944. DOI: 10.1093/ee/nvw060.

Coudrain, V; Schüepp, C; Herzog, F; Albrecht, M and Entling, M (2014). Habitat amount modulates the effect of patch isolation on host-parasitoid interactions. *Front. Environ. Sci.*, 2: 27. DOI: 10.3389/fenvs.2014.00027.

Damien, M; Llopis, S; Desneux, N; Van Baaren, J and Le Lann, C (2020). How does floral nectar quality affect life history strategies in parasitic wasps? *Entomol. Gen.*, 40(2): 147-156. DOI: 10.1127/ entomologia/2020/0906.

Darras, K F A; Corre, M D; Formaglio, G; Tjoa, A; Potapov, A; Brambach, F; Sibhatu, K T; Grass, I; Rubiano, A A; Buchori, D; Drescher, J; Fardiansah, R; Hölscher, D; Irawan, B; Kneib, T; Krashevska, V; Krause, A; Kreft, H; Li, K; Maraun, M, Polle, A; Ryadin, A R; Rembold, K; Stiegler, C; Scheu, S; Tarigan, S; Valdés-Uribe, A; Yadi, S; Tscharntke, T and Veldkamp, E (2019). Reducing fertilizer and avoiding herbicides in oil palm plantations - Ecological and economic valuations. *Front. For. Glob. Change*, 2: 65. DOI: 10.3389/ ffgc.2019.00065.

Devi, S; Ram, P and Rolania, K (2020). Effect of intercropping on the parasitoids, *Encarsia* spp. and *Trichogramma* spp. in cotton fields, India. *Egypt. J. Biol. Pest Control, 30*(1): 71. DOI: 10.1186/s41938-020-00275-4.

Edgar, R C (2010). Search and clustering orders of magnitude faster than BLAST. *Bioinformatics*, *26(19)*: 2460-2461. DOI: 10.1093/bioinformatics/ btq461.

Edgar, R C (2013). UPARSE: Highly accurate OTU sequences from microbial amplicon reads. *Nat. Methods*, *10*(*10*): 996-998. DOI: 10.1038/nmeth.2604.

Fahimee, J; Badrulisham, A S; Zulidzham, M S; Reward, N F; Muzammil, N; Jajuli, R; Md-Zain, B M and Yaakop, S (2021). Metabarcoding in diet assessment of *Heterotrigona itama* based on *trnL* marker towards domestication program. *Insects*, *12*(*3*): 1-16. DOI: 10.3390/insects12030205.

Folgarait, P J and Gilbert, L E (1999). Phorid parasitoids affect foraging activity of *Solenopsis richteri* under different availability of food in Argentina. *Ecol. Entomol.*, 24(2): 163-173. DOI: 10.1046/j.1365-2311.1999.00180.x.

Fuat, S; Adam, N A; Hazmi, I R and Yaakop, S (2022). Interactions between *Metisa plana*, its hyperparasitoids and primary parasitoids from good agriculture practices (GAP) and non-gap oil palm plantations. *Community Ecol.*, 23: 429-438. DOI: 10.1007/s42974-022-00092-9.

Halim, M; Aman-Zuki, A; Syarifah Zulaikha, S A; Muhaimin, A M D; Atikah, A R; Masri, M M M; Md-Zain, B M and Yaakop, S (2018). Exploring the abundance and DNA barcode information of eight parasitoid wasps species (Hymenoptera), the natural enemies of the important pest of oil palm, bagworm, *Metisa plana* (Lepidoptera: Psychidae) toward the biocontrol approach and it's application. *J. Asia. Pac. Entomol.*, *21*(*4*): 1359-1365. DOI: 10.1016/j.aspen.2018.10.012.

Halim, M; Muhaimin, A M D; Syarifah Zulaikha, S A; Nor Atikah, A R; Masri, M M M and Yaakop, S (2017). Evaluation of infestation in parasitoids on *Metisa plana* Walker (Lepidoptera: Psychidae) in three oil palm plantations in Peninsular Malaysia. *Serangga*, 22(2): 135-149.

Halim, M; Syarifah Zulaikha, S A; Muhaimin, A M D; Din, M and Yaakop, S (2019). The diversity and abundance of potential hymenopteran parasitoids assemblage associated with *Metisa plana* (Lepidoptera: Psychidae) in three infested oil palm plantations in Peninsular Malaysia. *AIP Conf Proc.*, 2111: 060024. DOI: 10.1063/1.5111286.

Hariyadi and Syahlan, A (2021). The role of planting media and mulch in the growth of *Leptopus antigonon* as a beneficial plant on oil

palm plantation. *IOP Conference Series: Earth and Environmental Science*, 694(1): 012029. DOI: 10.1088/1755-1315/694/1/012029.

Heimpel, G E and Jervis, M A (2005). Does floral nectar improve biological control by parasitoids? *Plant-Provided Food for Carnivorous Insects: A Protective Mutualism and its Applications* (Wäckers; van Rijn and Bruin eds.). Cambridge University Press, Cambridge, England. p. 267-304. DOI: 10.1017/CBO9780511542220.010.

Jervis, M A; Kidd, N A C and Walton, M (1992). A review of methods for determining dietary range in adult parasitoids. *Entomophaga*, *37*(*4*): 565-574. DOI: 10.1007/BF02372326.

Landis, D A; Wratten, S D and Gurr, G M (2000). Habitat management to conserve natural enemies of arthropod pests in agriculture. *Annu. Rev. Entomol.*, 45: 175-201. DOI: 10.1146/annurev.ento.45.1.175.

Lewis, W J; Stapel, J O; Cortesero, A M and Takasu, K (1998). Understanding how parasitoids balance food and host needs: Importance to biological control. *Biol. Control*, *11*: 175-183. DOI: 10.1006/ bcon.1997.0588.

Liu, A Z; Li, D Z; Wang, H and Kress, W J (2002). Ornithophilous and Chiropterophilous pollination in *Musa itinerans* (Musaceae), a pioneer species in tropical rain forests of Yunnan, Southwestern China. *Biotropica*, *34*(2): 254-260. DOI: 10.1111/ j.1744-7429.2002.tb00536.x.

Liu, H and Ueno, T (2012). The importance of food and host on the fecundity and longevity of a host-feeding parasitoid wasp. *J. Fac. Agr., Kyushu Univ., 57(1)*: 121-125. DOI: 10.5109/22058.

Mallott, E K; Garber, P A and Malhi, R S (2018). *trnL* outperforms *rbcL* as a DNA metabarcoding marker when compared with the observed plant component of the diet of wild white-faced capuchins (*Cebus capucinus*, Primates). *PLoS ONE*, *13(6)*: e0199556. DOI: 10.1371/journal. pone. 0199556.

Mathews, J; Yong, K K and Nurulnahar, B E (2007). Preliminary investigation on biodiversity and its ecosystem in oil palm plantation. *Proc. PIPOC 2007 Int. Palm Oil Cong. Agri., Biotechnol. & Sustain., Vol. 2*: p. 1112-1158.

Möller, G; Keasar, T; Shapira, I; Möller, D; Ferrante, M and Segoli, M (2021). Effect of weed management on the parasitoid community in Mediterranean Vineyards. *Biology* 10: 7. DOI: 10.3390/biology 10010007.

Morley, D (2015). RSPO, the global standard for sustainable palm oil. *Agro. Food Ind. Hi Tech*, 26(6): 29-30.

Nchanji, Y K; Nkongho, R N; Mala, W A and Levang, P (2016). Efficacy of oil palm intercropping by smallholders: Case study in South-West Cameroon. *Agroforest. Syst.*, *90*: 509-519. DOI: 10.1007/s10457-015-9873-z.

Norman, K and Basri, M W (2010). Interactions of the bagworm *Pteroma pendula* (Lepidoptera: Psycidae) and its natural enemies in an oil palm plantation in Perak. *J. Oil Palm Res.*, 22: 758-764.

Norman, K; Basri, M W and Masijan, Z (2017). Handbook of common parasitoids and predators associated with bagworms and nettle caterpillars in oil palm plantations. MPOB, Bangi. 29 pp.

Norman, K; Idris, A S and Masri, M M M (2019). Prospects in sustainable control of oil palm pests and diseases through the enhancement of ecosystem services - the way forward. *J. Oil Palm Res.*, *31*(*3*): 381-393. DOI: 10.21894/jopr.2019.0030.

Olson, D M; Takasu, K and Lewis, W J (2005). Food needs of adult parasitoids: Behavioral adaptations and consequences. *Plant-Provided Food for Carnivorous Insects: A Protective Mutualism and its Applications* (Wäckers; Van Rijn and Bruin eds.). Cambridge University Press, Cambridge. p. 137-147. DOI: 10.1017/CBO9780511542220.006.

Patt, J M; George, C H and James, H L (1997). Foraging success of parasitoid wasps on flowers: Interplay of insect morphology, floral architecture and searching behavior. *Entomol. Exp. Appl.*, *83*: 21-30.

POWO (Plants of the world online) (2021). *Mackaya neesiana* (Wall.) Das. http://www.plantsoftheworldonline.org/taxon/urn:lsid:ipni.org:names:52247-1, accessed on 21 July 2021.

R Core Team (2018). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria.

Rocha, R N C; Rodrigues, M D R L; Lopes, R; Cysne, A Q and Macedo, J L V D (2020). Production and cash flow of oil palm intercropping systems an Amazonian degraded area. *Nativa* (*Sinop*), 8(1): 52-57. DOI: 10.31413/nativa.v8i1.8073.

Salmah, M; Idris, A B and Basri, M W (2006). Effect of host larval stages on development and oviposition behaviour of *Apanteles metesae* Nixon (Hymenoptera: Braconidae), a parasitoid of the oil palm bagworm, *Metisa plana* Walker (Lepidoptera: Psychidae). *Malays. Appl. Biol.*, *35*(2): 21-28.

Saswattecha, K; Kroeze, C; Jawjit, W and Hein, L (2015). Assessing the environmental impact of palm oil produced in Thailand. *J. Clean. Prod.,* 600: 150-169. DOI: 10.1016/j.jclepro.2015. 03.037.

Satriawan, H and Fuady, Z (2019). Short communication: Analysis of weed vegetation in immature and mature oil palm plantations. *Biodiversitas*, 20(11): 3292-3298. DOI: 10.13057/biodiv/d201123.

Shi, Z; Chen, Y L; Chen, Y S; Lin, Y R; Liu, S W; Ge, X J; Gao, T G; Zhu, S X; Liu, Y; Yang, Q E; Humphries, C J; Raab-Straube, E von; Gilbert, M G; Nordenstam, B; Kilian, N; Brouillet, L; Illarionova, I D; Hind, D J N; Jeffrey, C; Bayer, R J; Kirschner, J; Greuter, W; Anderberg, A A; Semple, J C; Štěpánek, J; Freire, S E; Martins, L; Koyama, H; Kawahara, T; Vincent, L; Sukhorukov, A P; Mavrodiev, E V and Gottschlich, G (2011). Asteraceae (Compositae). *Flora of China* (Wu, Z Y; Raven, P H and Hong, D Y eds.). Science Press (Beijing) & Missouri Botanical Garden Press (St. Louis) Vol. 20-21. pp. 894.

Singh, A; Weisser, W W; Hanna, R; Houmgny, R and Zytynska, S E (2017). Reduce pests, enhance production: benefits of intercropping at high densities for okra farmers in Cameroon. *Pest Manag. Sci.*, 73(10): 2017-2027. DOI: 10.1002/ps.4636.

Siti Ramlah, A A; Wahid, M B; Najib, M and Masri, M M M (2005). Integrated pest management: Terakil-1, *Cassia cobanensis* and beneficial insects for controlling of bagworms infestation in Malaysia. *Proc. of the PIPOC 2005 International Palm Oil Congress - Technologies Breakthroughs and Commercialization the Way Forward*. MPOB, Bangi. p. 660-690.

Tan, K T; Lee, K T; Mohamed, A R and Bhatia, S (2009). Palm oil: Addressing issues and towards sustainable development. *Renew. Sust. Energ. Rev.*, 13(2): 420-427. DOI: 10.1016/j.rser.2007.10.001.

Tiong, A (1982). Oil palm pests in Sarawak and the use of natural enemies to control them. *Proc. of the International Conference on Plant Protection in the Tropics*. Malaysian Plant Protection Society, Kuala Lumpur. p. 363-372.

Tompkins, J M L; Wratten, S D and Wäckers, F L (2010). Nectar to improve parasitoid fitness in biological control: Does the sucrose:hexose ratio matter? *Basic Appl. Ecol.*, *11*(*3*): 264-271. DOI: 10.1016/j.baae.2009.12.010.

Vandebroek, I; Picking, D; Aiken, S; Lewis, P A; Oberli, A; Mitchell, S and Boom, B (2018). A review of Coralilla (*Antigonon leptopus*): An invasive and popular urban bush medicine in Jamaica. *Econ. Bot.*, 72(2): 229-245. DOI: 10.1007/ s12231-018-9415-5.

Wood, B J; Liau, S S and Knecht, J C X (1974). Trunk injection of systemic insecticides against the bagworm, *Metisa plana* (Lepidoptera: Pyralidae) on oil palm. *Oléagineux*, 29(11): 499-505.

Wood, B J and Norman, K (2019a). A review of developments in integrated pest management (IPM) of bagworm (Lepidoptera: Psychidae) infestation in oil palms in Malaysia. *J. Oil Palm Res.*, *31*(*4*): 529-539. DOI: 10.21894/jopr.2019.0047.

Wood, B J and Norman, K (2019b). Bagworm (Lepidoptera: Psychidae) infestation in the centennial of the Malaysian oil palm industry - A review of causes and control. *J. Oil Palm Res. 31*(*3*): 364-380. DOI: 10.21894/jopr.2019.0032.

Xiang, H; Lan, N; Wang, F; Zhao, B; Wei, H and Zhang, J (2021). Reduced pests, improved grain quality and greater total income: Benefits of intercropping rice with *Pontederia cordata*. J. Sci. Food Agric., 101(14): 5907-5917. DOI: 10.1002/jsfa.11243.

Yusdayati, R; Che Salmah, M D R; Ahmad, A H and Noor, H H (2014). Diversity and distribution of natural enemies (predators and parasitoids) of bagworms (Lepidoptera: Psychidae) on selected host plants in an oil palm plantation. *Planter, 90* (1055): 91-101.