Elucidating the diet of the island flying fox (*Pteropus hypomelanus*) in Peninsular Malaysia through Illumina Next-Generation Sequencing

Sheema Abdul Aziz¹²³⁴, Gopala Mountains Reuben Clements¹²³⁵⁶, Lee Yin Peng⁵⁷, Ahimsa Campos-Arceiz³, Kim R. McConkey³⁸, Pierre-Michel Forget² and Han Ming Gan⁵⁷

¹ Rimba, Bandar Baru Bangi, Selangor, Malaysia
² UMR MECAD 7179 CNRS-MNHN, Muséum National d’Histoire Naturelle, Département Adaptations du Vivant, Brunoy, France
³ School of Environmental and Geographical Sciences, The University of Nottingham Malaysia Campus, Semenyih, Selangor, Malaysia
⁴ Centre for Biological Sciences, Faculty of Natural and Environmental Sciences, University of Southampton, Southampton, United Kingdom
⁵ School of Science, Monash University Malaysia, Petaling Jaya, Selangor, Malaysia
⁶ Kenyir Research Institute, Universiti Malaysia Terengganu, Kuala Terengganu, Malaysia
⁷ Genomics Facility, Tropical Medicine and Biology Platform, Monash University Malaysia, Petaling Jaya, Selangor, Malaysia
⁸ School of Natural Sciences and Engineering, National Institute of Advanced Studies, Indian Institute of Science Campus, Bangalore, India

ABSTRACT

There is an urgent need to identify and understand the ecosystem services of pollination and seed dispersal provided by threatened mammals such as flying foxes. The first step towards this is to obtain comprehensive data on their diet. However, the volant and nocturnal nature of bats presents a particularly challenging situation, and conventional microhistological approaches to studying their diet can be laborious and time-consuming, and provide incomplete information. We used Illumina Next-Generation Sequencing (NGS) as a novel, non-invasive method for analysing the diet of the island flying fox (*Pteropus hypomelanus*) on Tioman Island, Peninsular Malaysia. Through DNA metabarcoding of plants in flying fox droppings, using primers targeting the *rbcL* gene, we identified at least 29 Operationally Taxonomic Units (OTUs) comprising the diet of this giant pteropodid. OTU sequences matched at least four genera and 14 plant families from online reference databases based on a conservative Least Common Ancestor approach, and eight species from our site-specific plant reference collection. NGS was just as successful as conventional microhistological analysis in detecting plant taxa from droppings, but also uncovered six additional plant taxa. The island flying fox’s diet appeared to be dominated by figs (*Ficus* sp.), which was the most abundant plant taxon detected in the droppings every single month. Our study has shown that NGS can add value to the conventional microhistological approach in identifying food plant species from flying fox droppings. At this point in time, more accurate genus- and species-level identification of OTUs not only requires support from databases with more representative sequences of relevant plant DNA, but probably necessitates *in situ* collection of plant specimens to create a reference collection. Although this method...
cannot be used to quantify true abundance or proportion of plant species, nor plant parts consumed, it ultimately provides a very important first step towards identifying plant taxa and spatio-temporal patterns in flying fox diets.

**Subjects** Biodiversity, Conservation Biology, Ecology, Genetics, Molecular Biology

**Keywords** Amplicon, rbcL, Pteropodid, Fruit bat, Metabarcoding, Phytophagous, Frugivory, Nectarivory

### INTRODUCTION

Understanding the contribution of animals to the functioning of rainforests has become an important issue in conservation biology. Conservation studies are now recognizing the need to collect qualitative and quantitative information on trophic relationships between animals and plants, not only to identify potential ecosystem service providers (Pompanon et al., 2012; Hibert et al., 2013), but also to inform management interventions for threatened species (Valentini et al., 2009; Ando et al., 2013).

Bats (Order: Chiroptera) provide important ecosystem services such as insect pest suppression, pollination, and seed dispersal (Fujita & Tuttle, 1991; Kunz et al., 2011). Characterising their diet is a fundamental step towards understanding their ecological roles. In the Old World, fruit bats such as flying foxes (Pteropodidae: Pteropus spp., Acerodon spp.; Kingston, 2010) have become increasingly threatened by hunting for bushmeat and medicine (Mildenstein, Tanshi & Racey, 2016). Identifying their diet and roles as ecosystem service providers can help strengthen arguments for their protection. It will also help us understand the wider implications of large-scale flying fox extinctions, as these giant bats are known to interact with plants on a large landscape scale, performing ecological roles over vast transboundary areas (Epstein et al., 2009). Flying foxes are likely to be particularly important players in island ecosystems where they often serve as keystone pollinators and seed dispersers both within and between islands (Cox et al., 1991; Banack, 1998; McConkey & Drake, 2007; McConkey & Drake, 2015), and where maintaining their numbers at high densities is necessary for the survival of plant communities (McConkey & Drake, 2006). Such data are also important to understand the drivers and potential mitigation strategies for conflicts between fruit bats and humans (Aziz et al., 2016; Aziz et al., in press).

Whilst in-depth, comprehensive dietary/foraging studies have been conducted for certain flying fox species, particularly in Australia (e.g., Boulter et al., 2005; Williams et al., 2006), Oceania (e.g., McConkey & Drake, 2006; Luskin, 2010), Japan (e.g., Nakamoto, Kinjo & Izawa, 2007; Nakamoto, Kinjo & Izawa, 2009; Lee et al., 2009), South Asia (e.g., Mahmood-ul-Hassan et al., 2010; Sudhakaran & Doss, 2012), and Indian Ocean islands (e.g., Nyhagen et al., 2005; Oleksy, Racey & Jones, 2015), the diets of Southeast Asian species, which are some of the most threatened due to the additional threat of commercial hunting (Mildenstein, Tanshi & Racey, 2016), remain largely unknown. Indeed, apart from a few studies in the Philippines (Reiter & Curio, 2001; Mildenstein et al., 2005; Stier & Mildenstein, 2005), Thailand (Weber et al., 2015), and Myanmar (Win & Mya, 2015), all other dietary and foraging studies on Southeast Asian Pteropodidae have focused on the smaller
pteropodids (e.g., Hodgkison et al., 2004; Fletcher, Zubaid & Kunz, 2012; Bumrungsri et al., 2013; Stewart, Makowsky & Dudash, 2014). This is of particular concern given that out of the 67 flying fox species listed on the IUCN Red List, almost half (30 species i.e., 45%) are found in Southeast Asia (IUCN, 2016).

Due to the nocturnal and volant nature of bats, invasive analyses (by capturing individuals) or indirect methods (by collecting droppings) have traditionally been used to study their diets. Microscope analyses of pteropodid faeces have provided insights into their diet (Bumrungsri, Leelapaibul & Racey, 2007), as well as their roles in pollination (Bumrungsri et al., 2013) and seed dispersal (Sritongchuay et al., 2014). However, all these studies have relied on physical identification of food plant species—either through direct observations of foraging bats, or microhistological identification of seeds, pollen, fruit fibres and leaf fragments in faeces and ejecta (chewed-up pellets of plant parts spat out by bats after swallowing the juice or soft pulp; Nyhagen et al., 2005; Long & Racey, 2007). The successful use of these approaches relies on several important factors such as accessibility and visibility of foraging bats for the former method, and also the availability of expert botanical knowledge or resources such as reference collections. Another limitation of these conventional approaches is that they require physically identifiable remains to be expelled by the bats; any plant parts that were consumed or expelled solely in liquid form will be missed out in the analysis (Pompanon et al., 2012). Studies on the foraging ecology of wide-ranging species such as flying foxes also require the use of expensive, hi-tech equipment such as GPS collars, which is often not feasible.

Although molecular analysis of pteropodid diets can potentially be used to overcome the obstacles outlined above, this approach has yet to be applied. DNA analyses of faeces collected non-invasively have already been conducted to determine the herbivorous diets of animals such as primates (Bradley et al., 2007; Quéméré et al., 2013; Srivathsan et al., 2016), marmots, bears, capercaillies, grasshoppers, molluscs, slugs (Valentini et al., 2009a), pigeons (Ando et al., 2013) and tapirs (Hibert et al., 2013), but this has never before been attempted for pteropodids in the Palaeotropics. To date, molecular analyses of bat diets in the Old World have only been conducted for insectivorous species (e.g., Clare et al., 2009; Razgour et al., 2011; Zeale et al., 2011). In fact, to our knowledge, the only successful attempt to identify the diet of plant-visiting bats through molecular analysis has been done by one study in the Neotropics (Hayward, 2013).

We evaluated the utility of Illumina Next Generation Sequencing (NGS) to identify plant species present in the droppings of the island flying fox (Pteropus hypomelanus) from Tioman Island in Peninsular Malaysia. In addition, we evaluated the potential of NGS analysis in complementing or even replacing conventional microhistological analysis to elucidate flying fox diets. First, we created a site-specific reference collection of potential flying fox food plants—DNA sequencing of these plants facilitated the construction of a phylogenetic tree to support morphology-based identification of these food plant species and primer design for NGS. Next, we assessed the feasibility of extracting plant DNA from flying fox droppings, and evaluated whether DNA sequences obtained from NGS could be matched with those from online and site-specific DNA reference databases. Finally,
we compared the performance of NGS with conventional microhistological analyses to identify food plant species from flying fox droppings.

**MATERIALS AND METHODS**

**Study site**

We conducted this study on Tioman Island (2°48′38″N, 104°10′38″E; 136 km²; Fig. 1A), located 32 km off the east coast of Peninsular Malaysia in the State of Pahang. This research was approved by the Economic Planning Unit of Malaysia (Permit number: 3242). Much of the island inland is still covered by primary tropical rainforest, which has been designated as Pulau Tioman Wildlife Reserve (83 km²). It has a hilly topography, with flat areas only along the coast (Abdul, 1999). The area designated as a wildlife reserve is composed of lowland mixed dipterocarp forest and hill dipterocarp forest. Most forested areas are still inaccessible due to the rugged topography, with many steep slopes and rocky outcrops (Latiff et al., 1999). The climate is tropical, uniformly warm and humid throughout the year (Hasan Basyri et al., 2001), but the island experiences the northeast monsoon from November to March (Bullock & Medway, 1966).

There are currently seven villages on the island, situated along the coastline (Fig. 1A). The majority of the local people are Muslim, and therefore due to religious dietary restrictions...
do not hunt bats for food or medicine (Aziz, 2016). As the island’s marine area is also a designated Marine Park and a popular tourist destination, many of the local people are heavily involved in the tourism industry (Abdul, 1999).

**Study species**
The island flying fox (*Pteropus hypomelanus*), also known as the variable flying fox and the small flying fox, roosts gregariously, forming colonies of up to 5,000 individuals. It is a widespread insular species, considered to be abundant throughout a distribution range that extends from the Maldives and Indian islands in the west to Melanesia in the east. Because of this, it is considered to be Least Concern on a global scale by the IUCN Red List; however, populations are decreasing (Francis et al., 2008; Olival, 2008), and the species is now listed as endangered on the Malaysian Red list (DWNP, 2010).

On Tioman, the island flying fox can be found roosting permanently in two villages: Tekek, on the west coast, and Juara, on the east coast (Fig. 1A), and forages throughout the island (Medway, 1966; Ong, 2000). Monthly roost counts conducted during March–October 2015 using a thermalscope (Pulsar Quantum HD38S) yielded estimated ranges of 2,178–5,385 individuals for the entire island (see Fig. S1 for further details). Local people have reported that the flying foxes do forage in other villages on the island.

**Flying fox dropping collection**
Collection of flying fox droppings took place once a month during March-October 2016 (i.e., eight months). Samples of droppings consisting of faeces and ejecta were collected for three mornings in the last week of each month from three separate day roosts in Juara (east coast) and two separate day roosts in Tekek (west coast). The number of roosts and sampling days were determined based on species accumulation curves of pollen morphospecies that were detected through preliminary microhistological analysis in June 2014. Program EstimateS (version 9.1.0; http://viceroy.eeb.uconn.edu/estimates/) indicated that sampling completeness (i.e., observed/estimated number of species; Soberón, Llorente & Onate, 2000) was around 97% using this sampling regime.

In Juara, three suitable roost trees (Fig. 1B) for sampling were selected based on accessibility and also on the highest/largest amount of faecal/ejecta splatter produced under the roost, in order to maximise sample yield. As flying foxes often shifted roosts or temporarily abandoned degraded roosts, this meant that sometimes different roosts were sampled in each location every month or even every morning, although most roosts were consistently sampled each month due to their constant high occupancy and best accessibility.

Plastic sheets measuring 0.8 × 1.0 m were placed under each roost after dark, once the bats had exited the roost to forage. The roosts were then visited the next morning for collection starting at 0700 h and ending at 1200 h (bats typically returned to the roosts around 0500–0600h); the plastic sheets were carefully moved away from the roost to a clear area for processing (Fig. 1C). As it was often difficult to differentiate faeces from ejecta, both were collected and analysed equally as ‘droppings’ (Fig. 1D). Droppings collected for processing were selected based on unique colour and texture, as this was assumed
to be representative of plant diversity in the bats’ diet. Following the approach used by Stier & Mildenstein (2005) based on short gut-passage time for flying foxes (12–34 min; Tedman & Hall, 1985), we assumed that each bat voided its last meal once, and therefore each dropping represented a different individual’s food choice. We devised our own novel collection protocol where droppings were collected by swabbing them with a cotton bud, then placing each individual dropping into a 5 ml Eppendorf tube containing ~1,000 µl of 95% ethanol. These tubes were then kept cool in the field, either by storing in a conventional freezer or by using a portable cooler box with ice packs, for 1–3 days before being transported off the island and then stored in a −80 °C freezer.

In order to simultaneously test the utility of NGS and compare it with conventional approaches, we collected two duplicate sets of 10 individual droppings from one single roost in Juara village during a single morning on 6 May 2015. One sample set was then kept in a conventional fridge for microscope analysis, whilst the other set was stored in the −80 °C freezer for molecular analysis.

Site-specific plant reference collection
We first checked a published list of genera of known food plants for Pteropus across its range (Marshall, 1985), cross-checked this against a preliminary checklist of seed plants for Tioman (Latiff et al., 1999), and also obtained information on possible flying fox food plants through talking to local people in Juara. We then searched for genera of similar plants in and around the two villages with the aid of a local plant expert. The botanical identification of plants (at least to genus) were subsequently verified by a trained botanist familiar with plants from the region. When we found an individual plant from one of these genera, we recorded its GPS location and collected pollen, fruit, and/or seeds if it was flowering or fruiting. We also collected 3–5 mature leaves for DNA extraction. The leaves were stored in Ziploc bags with silica gel under cool conditions to retard decomposition rates. Leaf samples from 19 different plant species were obtained for this purpose, constituting a preliminary library (Table S1).

Genomic DNA was extracted from approximately 25 mg of one leaf from each plant species using DNAeasy Plant Mini Kit (Qiagen, Halden, Germany) according to the manufacturer’s protocols. Primers targeted the rbcl gene, a protein-coding gene associated with the chloroplast genome of all living plants. DNA amplifications were performed in a mastermix containing 1 µL of DNA, 25 µL of OneTaq Quick-Load 2X Master Mix with Standard Buffer, (New England Biolab, Ipswich, MA, USA), 1 µL of 10mM forward primer rbclAf-M13, 1 µL of 10 mM reverse primer rbclA-revM13 (Table S2), and 22 µL of nuclease-free water. The PCR protocol was started with an initial denaturation step for 30 s at 94 °C, followed by 30 cycles of 30 s at 94 °C, 30 s at 48 °C, 40 s at 68 °C, and final elongation for 2 min at 68 °C. The PCR products were purified using 0.8× volume ratio of Agencourt Ampure XP beads (Beckman Coulter, Inc). The purified samples were sent to 1st BASE laboratories (http://www.base-asia.com/) for Sanger sequencing. The sequencing results were quality trimmed using CodonCode TraceViewer (http://www.codoncode.com/traceviewer/) and aligned using MAFFT version 7.0 (Katoh & Standley, 2013).
A phylogenetic tree (Fig. S4) was constructed to support morphology-based identification of site-specific reference plants. Additional rbcL gene sequences that displayed high similarity to the sequenced plant specimens were mined from the BOLD/NCBI databases, aligned with MAFFT version 7.0 and trimmed using TrimAl version 1.9 optimized for maximum likelihood tree construction (Capella-Gutiérrez, Silla-Martínez & Gabaldón, 2009). The trimmed alignment was subsequently used to construct a maximum likelihood tree with FastTree using the –nt (nucleotide) and –gtr (generalised time-reversible model) setting.

**Laboratory procedure**

Based on the alignment of rbcL gene fragments from the site-specific reference plant DNA (Fig. S2), we designed a primer pair targeting 198 bp of the rbcL gene (minus primer sequence) using Primer3 (http://bioinfo.ut.ee/primer3-0.4.0/) on default settings (Table S2). Prior to primer synthesis, partial Illumina adapter sequences were added to the 5’ end of the designed primers, rbcl-357F and rbcl-556R, to allow barcoding and sequencing on the Illumina platform.

Individual droppings were pooled according to roost (n = 5, 2 in Tekek and 3 in Juara) and month (n = 8), creating 40 separate mixtures for analysis. The tubes containing the daily samples were first vortexed for 2 min to homogenise the content and subsequently, 1,000 µL of the sample was pipetted into another tube to form the mixture. Next, 100 µL of the mixture was used for DNA extraction similarly using DNAeasy Plant Mini Kit (Qiagen, Halden, Germany) instead of a stool-specific DNA extraction kit to improve the recovery of plant-derived DNA from faecal samples.

PCR reaction was performed using IlluM_rbcLF and IlluM_rbcLR. The 20 µL PCR cocktail consists of 10 µL Q5 Hot Start High-Fidelity 2× Master Mix (New England Biolab, Ipswich, MA, USA), 1 µL each of 10 µM forward and reverse primer, 1 µL gDNA and 7 µL nuclease-free water. All reactions were performed in a Veriti® 96-Well Fast Thermal Cycler with the following protocol: initial denaturation for 30 s at 98 °C, 25 cycles of 10 s at 98 °C, 30 s at 55 °C and 10 s at 65 °C, with a final 1 min extension at 65 °C. The PCR product was purified using 0.8× vol. ratio Agencourt Ampure XP beads (Beckman Coulter, Inc). Then, 1 uL of Index 1 and Index 2 primers from Nextera XT kit were added to 3 uL of purified PCR product and combined with 5 uL of Q5 Hot Start High-Fidelity 2X M aster Mix (New England Biolabs, Ipswich, MA). The PCR protocol was as followed: initial denaturation for 30 s at 98 °C, 8 cycles of 10 s at 98 °C and 1 min at 65 °C, with a final 1 min extension at 65 °C.

**Sequence analysis and taxon assignation**

The purified amplicons containing the full length Illumina adapter and appropriate unique barcode were then quantified using KAPA Library Quantification kit (Kapa Biosystems, CapeTown, South Africa) on the EcoRealTime PCR system (Illumina, San Diego, CA, USA). Based on the qPCR data, the amplicons were normalised, pooled and subsequently sequenced on the MiSeq (2 × 250 bp paired-end run) located at the Monash University Malaysia Genomics Facility.
Illumina Nextera adapters and primer sequences of the reads were trimmed off using Trimmomatic v 0.33 and FastX trimmer, respectively (Bolger, Lohse & Usadel, 2014; http://hannonlab.cshl.edu/fastx_toolkit/). The trimmed paired-end reads were then merged using PEAR (Zhang et al., 2014) using default settings. Dereplication, singleton removal and Operationally Taxonomic Unit (OTU) clustering (default setting of 97% identity clustering; -id 0.97) were performed using the pipeline implemented in UPARSE (Edgar, 2013). The filtered OTUs were translated into protein sequence and manually inspected for translated sequence containing stop codon(s), which were removed from subsequent analyses. Taxonomic assignment of OTUs was conducted by matching sequences against those from the BOLD/NCBI non-redundant nucleotide database as of December 2016. The BLASTn output for each OTU was downloaded in “XML” format and imported into MEGAN6 (Huson et al., 2016) to calculate their Lowest Common Ancestor (LCA). LCA predicts the taxonomic rank of OTUs based on matches to a set of reference lineages such that the predicted taxonomic rank is shared by all matches within user pre-defined parameters. The implemented parameters for MEGAN6 were: Minimum Blast bit score = 200, Max expected E-value = 0.01, Min Percent Identity = 90.0, Top percent (the percentage of the best score a match needs to lie within) = 10 and Weighted LCA% = 75. To estimate the OTU relative abundance in each sample, reads were mapped to the filtered OTUs via USEARCH (97% similarity cut-off) and normalised to 10,000 reads (Caporaso et al., 2010).

**Microhistological analysis**

For the 10 dropping samples collected in May, we sent one set for NGS analysis (following the protocol above) and used another set for microscope analysis. For the latter, we first manually broke up the dropping contents in the tube to produce a relatively more representative liquid sample. We then droppped 1–3 drops of this liquid onto a microscope slide using a pipette. Fuchsin jelly was added to this in order to stain pollen grains within the dropping, a slip cover was placed on top, and the jelly was then melted over an open flame, sealing the slip cover to the slide. The slide was then cooled down in a conventional fridge in order to allow the jelly to solidify again before examination.

Once cooled, we placed the slide under a conventional light microscope (Leica DM E) and first examined it using 10/0.25 magnification in order to detect pollen grains and other plant parts. We used a self-made reference collection as well as photos from Start (1974), S. Bumrungsri (http://www.seabcru.org/seabcru-resources) and Mohamed (2014) to identify pollen and other plant parts. When necessary, we used higher magnification (40/0.65) to view the pollen grains. We used just ‘presence/absence’ to assess pollen to avoid quantification biases towards species that naturally produce greater amounts of pollen. Following Thomas (2009), we considered a species present if we found three or more pollen grains on a single slide.
RESULTS

Site-specific plant reference database
For the 19 specimens in the site-specific plant reference database, the phylogenetic tree (Fig. S4) based on a 689 bp aligned rbcL gene region recovered the monophyly of several plant genera with high SH-like local support (>0.90). However, the sister grouping among genera in general was not well-supported, possibly due to the lack of comprehensive plant taxon sampling, and the use of a single genetic marker. The rbcL-based phylogeny generally supported the morphology-based identification of reference specimens of potential flying fox food plants, with 15 specimens correctly identified at least to genus level as shown by their monophyletic clustering with plant species of the same genus (Table 1). Sample PTMN02 (identified as Streblus asper based on morphology) was an interesting exception as it formed a sister group with the Ficus clade (SH-like local support = 0.938), but not with other congeners. rbcL sequences representing the plant species collected on Tioman were scarce in the BOLD/NCBI databases, suggesting these databases have insufficient sequence representation that can affect taxonomic assignment of plant DNA in the droppings.

Feasibility of using NGS to study flying fox diet
With our newly designed rbcL primer, we were able to successfully extract, amplify, and subsequently identify plant DNA from all of the collected flying fox droppings. Initially, a total of 160 Operationally Taxonomic Units (OTUs) were recovered from the sequencing reads, of which 29 OTUs (Table 1) were retained after filtering based on cumulative relative abundance (>0.5%) and presence of stop codon(s) in reading frame. Using a conservative LCA approach, we identified at least three different plant genera and at least 14 plant families from the droppings (Table 1). In addition, 8 OTUs matched with specimens from the site-specific plant reference collection.

Based on sampling completeness (calculated using EstimateS) for OTU relative abundance data from five roosts (data pooled over three days) per month using Chao 1 species richness estimator (good for datasets skewed towards low abundance classes; Chao, 1984), sampling completeness was relatively high for the months March (99%), April (100%), June (100%), August (100%), September (88%), and October (96%). However, sampling completeness could be improved for May (55%) and July (79%). The month of May, which had the lowest overall sampling completeness, also had the highest number of droppings collected (Table S3).

The results from our NGS analysis of island flying fox droppings over eight months suggest that the diet at both Juara and Tekek during this time was dominated by four different plant taxa (Table 1; Fig. 2): OTU 1 (Ficus sp.), OTU 3 (likely to be Mangifera indica based on sequence match with site-specific plant reference collection) and OTUs 4 & 5 (Rubiaceae). Spatio-temporal patterns in the relative abundance of these four taxa in the diet were observed during the sampling period (Fig. 2). For example, OTU 5 appeared to be consumed in similar proportions at both Juara and Tekek across all months whereas OTU 4 was consistently consumed in low proportions in Tekek yet consumed irregularly in Juara over the same period (Fig. 3). Even between different roosts in the same site, spatio-temporal differences were observed, such as for OTU 7 (Fig. S5).
Table 1  Summary information of 29 OTUs detected in flying fox droppings over eight months (Mar–Oct 2015) in Tioman Island, Malaysia: OTU identities (genus in bold) based on matches with plant sequences from online reference database using the Least Common Ancestor approach, identity of plant family, OTU relative abundance (Sum of reads for particular OTU/sum of reads for ALL OTUs × 100%) and matches with sequences from plant reference specimens.

| OTU No. | Lowest common ancestor (LCA): order/family/subfamily/genus | Family          | Relative abundance | Plant reference specimen match at 100% identity (code; Table 1) |
|---------|-------------------------------------------------------------|-----------------|-------------------|---------------------------------------------------------------|
| OTU 1   | *Ficus*                                                      | Moraceae        | 66.3              | *Ficus* sp. (PTMN11/PTMN22)                                    |
| OTU 3   | Anacardiaceae                                               | Anacardiaceae   | 15.43             | *Mangifera indica* (PTMN20)                                    |
| OTU 4   | Ixoroidae                                                   | Rubiaceae       | 5.33              |                                                               |
| OTU 5   | Naucleeae                                                   | Rubiaceae       | 2.64              |                                                               |
| OTU 6   | Lamiales                                                    |                  | 2.01              |                                                               |
| OTU 7   | *Diospyros*                                                 | Ebenaceae       | 1.67              |                                                               |
| OTU 8   | *Ficus*                                                     | Moraceae        | 1.4               |                                                               |
| OTU 9   | Anacardiaceae                                               | Anacardiaceae   | 0.64              |                                                               |
| OTU 10  | Moraceae                                                    | Moraceae        | 0.57              |                                                               |
| OTU 11  | Myrtoideae                                                  | Myrtaceae       | 0.42              | *Syzygium* sp. (PTMN08/PTMN09/PTMN17)                          |
| OTU 12  | Arecoideae                                                  | Areaceae        | 0.42              | *Cocos nucifera* (PTMN07)                                     |
| OTU 13  | *Terminalia*                                                | Combretaceae    | 0.4               | *Terminalia catappa* (PTMN19)                                  |
| OTU 15  | Malpighiales                                                | ?               | 0.37              |                                                               |
| OTU 16  | *Ficus*                                                     | Moraceae        | 0.25              |                                                               |
| OTU 17  | Moraceae                                                    | Moraceae        | 0.23              |                                                               |
| OTU 18  | Malvaceae                                                   | Malvaceae       | 0.21              | *Durio zibethinus* (PTMN16)                                    |
| OTU 19  | Fabids                                                      | ?               | 0.12              |                                                               |
| OTU 20  | Chrysobalanaceae                                            | Chrysobalanaceae| 0.11              |                                                               |
| OTU 21  | Ixoroidae                                                   | Rubiaceae       | 0.1               |                                                               |
| OTU 22  | Anacardiaceae                                               | Anacardiaceae   | 0.09              |                                                               |
| OTU 23  | Annonaceae                                                  | Annonaceae      | 0.07              |                                                               |
| OTU 24  | Pentapetalae                                                | ?               | 0.06              | *Strombosisia* sp. (PTMN06)                                    |
| OTU 25  | Anacardiaceae                                               | Anacardiaceae   | 0.05              |                                                               |
| OTU 26  | Naucleeae                                                   | Rubiaceae       | 0.05              |                                                               |
| OTU 27  | Lamiales                                                    | Lamiaceae       | 0.05              | *Vitex pinnata* (PTMN05)                                      |

Notes.

*LCA parameters: Min score = 200, Max expected = 0.01, Min percent identity = 0.0, Top Percent = 10, Weighted LCA% = 80.*

Performance of NGS vs. microhistological analysis

Microscope analysis identified two plant taxa in flying fox droppings (Table 2). Out of 10 individual droppings, three contained durian (*Durio* sp.) pollen. All the other droppings contained fig parts exclusively; no other plant parts were detected. Durian pollen occurred at extremely low abundance; in all cases, only 3–4 grains were detected per slide. No other
Figure 2  Relative abundance of 8 OTUs detected in flying fox droppings. Relative abundance of 8 OTUs detected in flying fox droppings across 8 months (Mar–Oct 2015) at two different roosting sites on Tioman Island, Tekek (two roosts) and Juara (three roosts). OTU 1, Ficus; OTU 3, Anacardiaceae; OTU 4, Rubiaceae; OTU 5, Rubiaceae; OTU 6, Diospyros; OTU 7, Moraceae; OTU 8, Lamiales; OTU 13, Terminalia; Others, pooled OTUs with <5% relative abundance at each roost.

Table 2 Comparative effectiveness of microhistological vs. NGS analyses in identifying plants from two duplicate sets of 10 samples of flying fox droppings collected on 6 May 2015. Plant ID (probable genus/family) for the microhistological analysis was based on visual identification from our plant reference collection, while plant ID for the NGS analyses was based on NGS sequence matches with online reference plant databases and DNA extracted from our plant reference collection.

| Plant ID | Microhistological analysis | NGS analysis |
|----------|---------------------------|--------------|
| 17 (Durio) | x x x x x x x x x x | x x x x x x x x x x |
| 1 (Ficus) | x x x x x x x x x x | x x x x x x x x x x |
| 3 (Mangifera) | x x x x x x x x x x | x x x x x x x |
| 19 (Strombosia) | x x x x x x x | x x x x x x |
| 13 (Terminalia) | x x x x x x x | x x x x x x |
| 12 (Arecaceae) | x x x x x x x x x x | x x x x x x |
| 5 (Rubiaceae) | x x x x x x x x x x | x x x x x x |

pollen or plant parts were detected. NGS identified the same two plant taxa detected by microhistological analysis, and further identified an additional six plant taxa. However, Durio was not detected in the same samples as those identified via microscope.

DISCUSSION

Our study is the first to describe the diet of the island flying fox, which was previously unknown. To our knowledge, this is also the first use of NGS to identify plant taxa in the diet of a pteropodid, which has been difficult to characterise due to these animals’ volant nature, large home ranges and nocturnal foraging behaviour. Figs consistently formed the highest amount of plant taxa detected in the droppings each month, at both sampling sites. This strongly suggests that figs compose the core diet of flying foxes on the island. It is thus
highly likely that the island flying fox plays a key role in dispersing fig seeds throughout Tioman, making these bats important keystone species for the island (Cox et al., 1991; McConkey & Drake, 2015); future studies on seed dispersal and germination are required to confirm this.

**NGS is a reliable tool to study flying fox diet**

We have demonstrated that identification of plant taxa to family level is generally possible based on the partial sequence of *rbcL* using the LCA approach. In addition, some OTUs in our study were successfully assigned to the genus level. In order to be conservative, however, we avoided assigning most OTUs to species level, unless there were matches with BOLD/NCBI database sequences and site-specific reference plant sequences. As
species-level plant identification based solely on NGS is not straightforward, the use of a site-specific food plant DNA reference database is vital to help identify plant species in flying fox diets.

Other genes have been successfully used to identify plant species in animal diets. For example, Valentini et al. (2009a) found the trnL intronic region to be effective for Asian mammals, birds, and invertebrates, identifying 50% of the plant taxa found in the diets of these animals to species level. The same approach has been used for European bison (Kowalczyk et al., 2011), alpine chamois (Raye’ et al., 2011), and red-headed wood pigeons (Ando et al., 2013). The trnL intron has been previously reported to evolve more than three times faster than the protein-coding rbcL gene, thus potentially harbouring more variation and phylogenetic signal per base pair (Gielly & Taberlet, 1994). However, we chose the rbcL gene instead of the P6 loop of the trnL intron to study flying fox diet, as the rbcL gene is currently one of the two genes (the other being matK) that is increasingly being used for plant species identification. This is useful for studying plants originating from a less-studied region of high biodiversity such as Southeast Asia. Unlike the trnL database, the rbcL database is consistently growing in the BOLD database, and as a result is also likely to be represented by more sequences from properly identified and vouchered specimens. In addition, given that rbcL is a protein-coding gene, it enables the screening of erroneous OTU resulting from sequencing and/or amplification errors based on the presence of stop codon(s) in the translated reading frame. Instead of using recently developed and robustly tested primers (e.g., Little, 2014), we designed a new set of rbcL minibarcode primers due to the current lack of rbcL sequence representation for plant species from Malaysia or specifically, Tioman. One caveat of this approach would be the possibility of preferential primers binding to known diet items instead of unexpected ones. However, our results seem to indicate that our newly designed primers were, to a certain extent, capable of recovering OTUs belonging to a wide variety of plant families, in part due to the diverse representation of plant reference specimens that contributed to the expanded taxon coverage of our primers. Future studies aiming to achieve greater power of identification of plants in flying fox diets could consider using more than one target region (e.g., Hibert et al., 2013; Clare, 2014) coupled with an additional in silico PCR optimisation step using ecoprimers to improve the reliability and universality of the newly designed primers (Riaz et al., 2011). Also, in order to completely eliminate bias associated with PCR, metagenomic shotgun sequencing could be performed, albeit at a relatively higher cost depending on the required sequencing coverage. (e.g., Srivathsan et al., 2015; Srivathsan et al., 2016).

NGS can complement microhistological analysis

Our results showed that NGS can provide greater insights into the diet of flying foxes than conventional microhistological approaches by detecting a wider range of plant taxa, thus highlighting the utility and discriminatory potential of the newly designed rbcL primers to study flying fox diets. More importantly, the use of NGS allowed us to identify plant species even when no physical plant parts were found in the flying fox droppings. The plant genera and families detected from NGS have also been recorded by botanists as being present on
Tioman, including the top four genera/families detected most abundantly in the droppings (Latiff et al., 1999; Mohd. Norfaizal et al., 2014).

In our study, attempts to use microscope analysis to identify plant parts in droppings proved to be challenging, as no pre-existing reference collection was available. Building our own microhistological reference collection for Tioman was time-consuming and labour-intensive—and the resulting collection often did not match up with the plant parts found in the flying fox droppings. However, obtaining DNA from plant specimens is still necessary to narrow down the identity of OTUs to species level. Indeed, 8 out of 29 OTUs had 100% matches to the sequences of plant specimens collected from the study site, highlighting the importance of building a comprehensive local sequence library beforehand, preferably specific to one’s particular study site.

It is important to note that NGS did not detect *Durio* in the same individual droppings as those identified via microscope. This is likely due to the low abundance of this plant taxon in the droppings affecting detection probability, especially since the NGS analysis used a more general primer that was not specific to *Durio*. This pollen detection probability is another caveat to be aware of; Scanlon et al. (2014) have cautioned that faecal subsampling methods can potentially lead to inaccurate detection of pollen in dietary studies, regardless of which method is used.

**Caveats**

Our sample collection method in the field, selecting only for droppings with unique colour and texture, may have introduced a bias that could result in underestimating the relative abundance of OTUs in the droppings. In particular, sampling completeness for the months May, June and July were relatively low, showing that more roosts and/or days needed to be sampled in order to obtain a complete representation of diet for these months. However, roost count data (Fig. S1) show that this high diversity in diet was not influenced by population abundance. Instead, perhaps diet choice and/or food resource diversity were relatively higher during this period compared to the rest of the year. Future studies should aim to collect every single dropping found underneath a roost to improve representativeness.

Given the potentially short flying fox gut passage times (Tedman & Hall, 1985), droppings collected from day roosts in the morning may also bias the analysis results towards food items that were consumed only at the end of the foraging period (Schmelitschek, French & Parry-Jones, 2009). Food plants that were consumed during the start or middle of the evening may not have been detected by our methods. Although Banack & Grant (2002) observed flying foxes returning to food resources that were foraged upon earlier, before then returning to day roosts, this is still a potential caveat to bear in mind. For example, primates are known to exhibit temporal patterning in diet choice, structuring their diet throughout their foraging period with different food items; it is believed that this is due to how different foods are processed, and give energy, at different rates, and therefore helps to ensure that the animals maintain high energy levels (Robinson, 1984; Ganzhorn & Wright, 1994; Chapman & Chapman, 1991). Given the size of Tioman,
and the logistical challenges of observing flying foxes foraging, the best way to overcome this possible information gap is to conduct GPS tracking studies.

We also acknowledge that NGS approaches to diet identification are semi-quantitative because chloroplast abundance is variable in different plant species and different parts of the leaf. Ultimately, the ability of NGS to accurately identify food plants will always depend on sequence specificity of the primers. While the NGS approach has proven to be useful in elucidating the island flying fox’s varied diet on Tioman, for animals with such a diverse phytophagous diet, primer specificity will always be a limiting factor and there is a chance that unknown plant species will not be detected due to primer mispriming. Also, identical chloroplast DNA sequences can be present in different but related species, making it impossible to distinguish closely related plant species from each other in the diet. This could be one possible factor why several OTUs could only be identified to order/family/subfamily levels, suggesting that they require further phylogenetic investigation and/or may benefit from identification based on more rapidly-evolving plastid-coding genes such as matK. It is worth noting that the sequenced rbcL gene of some plant specimens collected in this study did not exhibit 100% identity matches to species in the BOLD/NCBI databases, which may be attributed to genetic diversity at the intra-species level or gaps in the database i.e., certain plant species consumed by the flying foxes may not yet have their corresponding sequences deposited in the database. Nevertheless, there is an urgent need for the BOLD/NCBI databases to have more representation of plant sequences from Peninsular Malaysia and Southeast Asia in general.

Another limitation of the NGS approach for generalist diets is that it does not identify which part of the plant was consumed. For specialised frugivores, nectarivores, or herbivores (e.g., folivorous), this may not be an issue. Flying foxes, however, are generalists which consume fruits, flowers, nectar, and even leaves (Marshall, 1985). It is this dietary plasticity which allows them to perform more than one ecological role in tropical landscapes. Therefore, identifying which plant parts are actually consumed is a crucial step towards identifying the ecosystem services that these bats provide. NGS can provide a first step towards identifying flying fox diet but should not be viewed as a replacement for microhistological analysis. Nevertheless, this approach has shed new light on flying fox diet by discovering plant taxa that were entirely missed out by the conventional approach. Ideally, studies using NGS should be combined with micro-histological analysis in order to fill in the gaps and broaden our understanding of pteropodid diet and foraging ecology. NGS can also be used in combination with comprehensive and long-term data on plant phenology, to observe which food resources are available at which time. Following on from this preliminary study, the identification of specific food plants via NGS can now help guide more in-depth plant sample collection and phenological observations.

**CONCLUSION**

Our study is the first to use NGS to identify potential plant species in flying fox diet, paving the way for a new approach to studying flying fox diets. Since our NGS analysis of flying fox diet was semi-quantitative, it is not yet possible to make any definite conclusions regarding
food preference vs. food availability; ultimately it is unclear to what extent sampling bias and detection probability may have influenced the type and relative abundance of plant taxa detected in our study. Yet some of the interesting patterns we observed are worth investigating in greater detail, particularly in combination with microhistological analysis. The results will also help to guide us in conducting more accurate and expanded phenology monitoring, and further collection of botanical samples. Further and more rigorous sampling, especially at the level of the individual animal, is required to understand the dietary patterns of this particular flying fox population, expand on the information provided here and build on our understanding of how flying foxes provide ecosystem services on Tioman Island and elsewhere.

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Competing Interests
Sheema Abdul Aziz and Gopalasamy Reuben Clements are co-founders of Rimba, Selangor, Malaysia.

Author Contributions
• Sheema Abdul Aziz and Han Ming Gan conceived and designed the experiments, performed the experiments, analyzed the data, contributed reagents/materials/analysis tools, wrote the paper, prepared figures and/or tables, reviewed drafts of the paper.
• Gopalasamy Reuben Clements conceived and designed the experiments, performed the experiments, wrote the paper, prepared figures and/or tables, reviewed drafts of the paper.
• Lee Yin Peng performed the experiments, analyzed the data, wrote the paper, prepared figures and/or tables.
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Field Study Permissions
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DNA Deposition
The following information was supplied regarding the deposition of DNA sequences:
Sanger sequencing results for the rbcL fragment of collected plant specimens were assigned accession numbers in GenBank/NCBI (see Table S1). NGS data were also deposited in the SRA database as mentioned in the Data Availability Section.

Accession numbers:
SRX1988609, SRX1988608, SRX1988607, SRX1988606, SRX1988605, SRX1988604, SRX1988603, SRX1988602, SRX1988601, SRX1988600, SRX1988599, SRX1988598, SRX1988597, SRX1988596, SRX1988595, SRX1988594, SRX1988593, SRX1988592, SRX1988591, SRX1988590, SRX1988589, SRX1988588, SRX1988587, SRX1988586, SRX1988585, SRX1988584, SRX1988583, SRX1988582, SRX1988581, SRX1988580, SRX1988579, SRX1988578, SRX1988577, SRX1988576, SRX1988575, SRX1988574, SRX1988573, SRX1988572, SRX1988571, SRX1988570, SRX1988569, SRX1988568, SRX1988567, SRX1988566, SRX1988565, SRX1988564, SRX1988563, SRX1988562, SRX1988561, SRX1988560
https://www.ncbi.nlm.nih.gov/sra/SRP080299.
Data Availability
The following information was supplied regarding data availability:

The research in this article only generated DNA sequences, which have already been listed in detail in the relevant section above. No other raw data or code were generated.

Supplemental Information
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REFERENCES

Abdul J. 1999. An introduction to Pulau Tioman. Raffles Bulletin of Zoology 6:3–4.
Ando H, Setsuko S, Horikoshi K, Suzuki H, Umehara S, Inoue-Murayama M, Isagi Y. 2013. Diet analysis by next-generation sequencing indicates the frequent consumption of introduced plants by the critically endangered red-headed wood pigeon Columba janthina nitens in oceanic island habitats. Ecology and Evolution 3:4057–4069 DOI 10.1002/ece3.773.
Aziz SA. 2016. Interactions between flying foxes, plants and people: implications for conservation. PhD thesis, Muséum National d’Histoire Naturelle, France.
Aziz SA, Clements GR, Giam X, Forget P-M, Campos-Arceiz A. 2017. Coexistence and conflict between the Island Flying Fox (Pteropus hypomelanus) and local people on a tropical island in Peninsular Malaysia. Human Ecology In Press.
Aziz SA, Olival KJ, Bumrungsri S, Richards GC, Racey PA. 2016. The conflict between pteropodid bats and fruit growers: species, legislation and mitigation. In: Bats in the anthropocene: conservation of bats in a changing world. Springer International Publishing, 377–426.
Banack SA. 1998. Diet selection and resource use by flying foxes (genus Pteropus). Ecology 79:1949–1967.
Banack SA, Grant GS. 2002. Spatial and temporal movement patterns of the flying fox, Pteropus tonganus, in American Samoa. The Journal of Wildlife Management 66:1154–1163 DOI 10.2307/3802948.
Bolger AM, Lohse M, Usadel B. 2014. Trimmomatic: a flexible trimmer for Illumina sequence data. Bioinformatics 30(15):2114–2120 DOI 10.1093/bioinformatics/btu170.
Boulter SL, Kitching RL, Howlett BG, Goodall K. 2005. Any which way will do—the pollination biology of a northern Australian rainforest canopy tree (Syzygium sayeri; Myrtaceae). Botanical Journal of the Linnean Society 149:69–84 DOI 10.1111/j.1095-8339.2005.00430.x.
Bradley BJ, Stiller M, Doran-Sheehy DM, Harris T, Chapman CA, Vigilant L, Poinar H. 2007. Plant DNA sequences from feces: potential means for assessing diets of wild primates. American Journal of Primatology 69:699–705 DOI 10.1002/ajp.20384.
Bullock JA, Medway L. 1966. The fauna of Pulau Tioman. I. General information. Bulletin of the Natural Museum 34:1–8.
Bumrungsri S, Lang D, Harrower C, Sriporaya E, Kitpipit K, Racey PA. 2013. The dawn bat, *Eonycteris spelaea* Dobson (Chiroptera: Pteropodidae) feeds mainly on pollen of economically important food plants in Thailand. *Acta Chiropterologica* 15:95–104 DOI 10.3161/150811013X667894.

Bumrungsri S, Leelapaibul W, Racey PA. 2007. Resource partitioning in sympatric *Cynopterus* bats in lowland tropical rainforest, Thailand. *Biotropica* 39:241–248 DOI 10.1111/j.1744-7429.2006.00245.x.

Capella-Gutiérrez S, Silla-Martínez JM, Gabaldón T. 2009. trimAl: a tool for automated alignment trimming in large-scale phylogenetic analyses. *Bioinformatics* 25:1972–1973 DOI 10.1093/bioinformatics/btp348.

Caporaso JG, Kuczynski J, Stombaugh J, Bittinger K, Bushman FD, Costello EK, Fierer N, Penña AG, Goodrich JK, Gordon JI, Huttley GA. 2010. QIIME allows analysis of high-throughput community sequencing data. *Nature Methods* 7:335–336 DOI 10.1038/nmeth.f.303.

Chao A. 1984. Non-parametric estimation of the number of classes in a population. *Scandinavian Journal of Statistics* 11:265–270.

Chapman CA, Chapman LJ. 1991. The foraging itinerary of spider monkeys: when to eat leaves? *Primatologica* 56:162–166.

Clare EL. 2014. Molecular detection of trophic interactions: emerging trends, distinct advantages, significant considerations and conservation applications. *Evolutionary Applications* 7:1144–1157 DOI 10.1111/eva.12225.

Clare EL, Fraser EE, Braid HE, Fenton MB, Hebert PD. 2009. Species on the menu of a generalist predator, the eastern red bat (*Lasiurus borealis*): using a molecular approach to detect arthropod prey. *Molecular Ecology* 18:2532–2542 DOI 10.1111/j.1365-294X.2009.04184.x.

Cox PA, Elmqvist T, Pierson ED, Rainey WE. 1991. Flying foxes as strong interactors in South Pacific island ecosystems: a conservation hypothesis. *Conservation Biology* 5:448–454 DOI 10.1111/j.1523-1739.1991.tb00351.x.

DWNP (Department of Wildlife and National Parks Peninsular Malaysia). 2010. *Red List of mammals for Peninsular Malaysia*. Cheras, Kuala Lumpur: DWNP.

Edgar RC. 2013. UPARSE: highly accurate OUT sequences from microbial amplicon reads. *Nature Methods* 10:996–998 DOI 10.1038/nmeth.2604.

Epstein JH, Olival KJ, Pulliam JRC, Smith C, Westrum J, Hughes T, Dobson AP, Zubaid A, Sohayati AR, Misliiah MB, Field HE, Daszak P. 2009. *Pteropus vampyrus*, a hunted migratory species with a multinational home-range and a need for regional management. *Journal of Applied Ecology* 46:991–1002 DOI 10.1111/j.1365-2664.2009.01699.x.

Fletcher C, Zubaid A, Kunz TH. 2012. Fruit diet of frugivorous bats (*Cynopterus brachyotis* and *Cynopterus horsfieldii*) in tropical hill forests of Peninsular Malaysia. *Mammalia* 76:389–397.

Francis C, Rosell-Ambal G, Bonaccorso FA, Heaney L, Molur S, Srinivasulu C. 2008. *Pteropus hypomelanus*. The IUCN Red List of Threatened Species 2008:
Fujita MS, Tuttle MD. 1991. Flying foxes (Chiroptera: Pteropodidae): threatened animals of key ecological and economic importance. *Conservation Biology* **5**:455–463 DOI 10.1111/j.1523-1739.1991.tb00352.x.

Ganzhorn JU, Wright PC. 1994. Temporal patterns in primate leaf eating: the possible role of leaf chemistry. *Folia Primatologica* **63**:203–208.

Gielly L, Taberlet P. 1994. The use of chloroplast DNA to resolve plant phylogenies: noncoding versus rbcL sequences. *Molecular Biology and Evolution* **11**:769–777.

Hasan Basyri D, Norhan AR, Amir Hashim MK, Kamarul Azlan MN. 2001. Study on water resources in Tioman Island. *Jurnal Teknologi* **34(F)**:51–64.

Hayward CE. 2013. DNA barcoding expands dietary identification and reveals dietary similarity in Jamaican frugivorous bats. MSc thesis, University of Western Ontario, Canada.

Hibert F, Taberlet P, Chave J, Scotti-Saintagne C, Sabatier D, Richard-Hansen C. 2013. Unveiling the diet of elusive rainforest herbivores in next generation sequencing era? The tapir as a case study. *PLOS ONE* **8(4)**:e60799 DOI 10.1371/journal.pone.0060799.

Hodgkison R, Balding ST, Zubaid A, Kunz TH. 2004. Temporal variation in the relative abundance of fruit bats (Megachiroptera: Pteropodidae) in relation to the availability of food in a lowland Malaysian rain forest. *Biotropica* **36**:522–533.

Huson DH, Beier S, Flade I, Górska A, El-Hadidi M, Mitra S, Ruscheweyh HJ, Tappu R. 2016. MEGAN community edition—interactive exploration and analysis of large-scale microbiome sequencing data. *PLOS Computational Biology* **12(6)**:e1004957 DOI 10.1371/journal.pcbi.1004957.

IUCN. 2016. IUCN Red List of Threatened Species. Version 2016-1. Available at http://www.iucnredlist.org (accessed on 4 July 2016).

Katoh K, Standley DM. 2013. MAFFT multiple sequence alignment software version 7: improvements in performance and usability. *Molecular Biology and Evolution* **30**:772–780 DOI 10.1093/molbev/mst010.

Kingston T. 2010. Research priorities for bat conservation in Southeast Asia: a consensus approach. *Biodiversity Conservation* **19**:471–484 DOI 10.1007/s10531-008-9458-5.

Kowalczyk R, Taberlet P, Coissac E, Valentini A, Miquel C, Kamiński T, Wójcik JM. 2011. Influence of management practices on large herbivore diet—Case of European bison in Białowieża Primeval Forest (Poland). *Forest Ecology and Management* **261**:821–828 DOI 10.1016/j.foreco.2010.11.026.

Kunz TH, De Torrez EB, Bauer D, Lobova T, Fleming TH. 2011. Ecosystem services provided by bats. *Annals of the New York Academy of Sciences* **1223**:1–38 DOI 10.1111/j.1749-6632.2011.06004.x.

Latiff A, Faridah Hanum I, Zainudin Ibrahim A, Goh MWK, Loo AHB, Tan HTW. 1999. On the vegetation and flora of Pulau Tioman, Peninsular Malaysia. *Raffles Bulletin of Zoology* **6**:11–72.
Lee Y-F, Takaso T, Chiang T-Y, Kuo YM, Nakanishi N, Tzeng HY, Yasuda K. 2009. Variation in the nocturnal foraging distribution of and resource use by endangered Ryukyu flying foxes (Pteropus dasymallus) on Iriomotejima Island, Japan. Contributions to Zoology 78:51–64.

Little DP. 2014. A DNA mini-barcode for land plants. Molecular Ecology Resources 14:437–446 DOI 10.1111/1755-0998.12194.

Long E, Racey PA. 2007. An exotic plantation crop as a keystone resource for an endemic megachiropteran, Pteropus rufus in Madagascar. Journal of Tropical Ecology 23:1–11 DOI 10.1017/S0266467406003774.

Luskin MS. 2010. Flying foxes prefer to forage in farmland in a tropical dry forest landscape mosaic in Fiji. Biotropica 42:246–250 DOI 10.1111/j.1744-7429.2009.00577.x.

Mahmood-ul-Hassan M, Gulraiz TL, Rana SA, Javid A. 2010. The diet of Indian flying-foxes (Pteropus giganteus) in urban habitats of Pakistan. Acta Chiropterologica 12:341–347 DOI 10.3161/150811010X537927.

Marshall AG. 1985. Old World phytophagous bats and their food plants: a survey. Zoological Journal of the Linnean Society 83:351–369.

McConkey KR, Drake DR. 2006. Flying foxes cease to function as seed dispersers long before they become rare. Ecology 87:271–276 DOI 10.1890/05-0386.

McConkey KR, Drake DR. 2007. Indirect evidence that flying foxes track food resources among islands in a Pacific archipelago. Biotropica 39:436–440 DOI 10.1111/j.1744-7429.2007.00269.x.

McConkey KR, Drake DR. 2015. Low redundancy in seed dispersal within an island frugivore community. AoB Plants:plv088 DOI 10.1093/aobpla/plv088.

Medway L. 1966. The mammals. Observations on the fauna of Pulau Tioman and Pulau Tulai, Bulletin of the National Museum, Singapore 34:32–34.

Mildenstein TL, Stier SC, Nuevo-Diego CE, Mills LS. 2005. Habitat selection of endangered and endemic large flying-foxes in Subic Bay, Philippines. Biological Conservation 126:93–102.

Mildenstein T, Tanshi I, Racey PA. 2016. Exploitation of bats for bushmeat and medicine. In: Bats in the Anthropocene: conservation of bats in a changing world. Berlin Heidelberg: Springer International Publishing, 325–375.

Mohamed NZ. 2014. The role of nectar-feeding bats (Pteropodidae) in the pollination ecology of the genus Sonneratia at Setiu mangrove areas, Malaysia. PhD thesis, University of Bristol, UK.

Mohd. Norfaizal G, Masrom H, Omar Y, Aisha-Farhana S. 2014. A preliminary flora survey in Gunung Kajang, Pulau Tioman, Pahang Darul Makmur, Peninsular Malaysia. Malaysian Applied Biology 43:17–23.

Nakamoto A, Kinjo K, Izawa M. 2007. Food habits of Orii’s flying-fox, Pteropus dasymallus inopinatus, in relation to food availability in an urban area of Okinawa-jima Island, the Ryukyu Archipelago, Japan. Acta Chiropterologica 9:237–249 DOI 10.3161/1733-5329(2007)9[237:FHOOFP]2.0.CO;2.
Nakamoto A, Kinjo K, Izawa M. 2009. The role of Orii’s flying-fox (Pteropus dasymallus inopinatus) as a pollinator and seed disperser on Okinawa-jima Island, the Ryukyu Archipelago, Japan. Ecological Research 24:405–414 DOI 10.1007/s11284-008-0516-y.

Nyhagen DF, Turnbull SD, Olesen JM, Jones CG. 2005. An investigation into the role of the Mauritian flying fox, Pteropus niger, in forest regeneration. Biological Conservation 122:491–497 DOI 10.1016/j.biocon.2004.08.012.

Oleksy R, Racey PA, Jones G. 2015. High-resolution GPS tracking reveals habitat selection and the potential for long-distance seed dispersal by Madagascan flying foxes Pteropus rufus. Global Ecology and Conservation 3:678–692 DOI 10.1016/j.gecco.2015.02.012.

Olival KJ. 2008. Population genetic structure and phylogeography of southeast asian flying foxes: implications for conservation and disease ecology. PhD dissertation, University of Columbia, USA.

Ong SY. 2000. Kajian Ekologi Pteropus hypomelanus di Kampung Ayer Batang, Pulau Tioman, Pahang. BSc thesis, National University of Malaysia, Malay.

Pompanon F, Deagle BE, Symondson WOC, Brown DS, Jarman SN, Taberlet P. 2012. Who is eating what: diet assessment using next generation sequencing. Molecular Ecology 21:1931–1950 DOI 10.1111/j.1365-294X.2011.05403.x.

Quéméré E, Hibert F, Miquel C, Lhuillier E, Rasolondraibe E, Champeau J, Rabarivola C, Nusbaumer L, Chatelain C, Gautier L, Ranirison P. 2013. A DNA metabarcoding study of a primate dietary diversity and plasticity across its entire fragmented range. PLOS ONE 8(3):e58971 DOI 10.1371/journal.pone.0058971.

Raye’ G, Miquel C, Coissac E, Redjadj C, Loison A, Taberlet P. 2011. New insights on diet variability revealed by DNA barcoding and high-throughput pyrosequencing: chamois diet in autumn as a case study. Ecological Research 26:265–276 DOI 10.1007/s11284-010-0780-5.

Razgour O, Clare EL, Zeale MR, Hanmer J, Schnell IB, Rasmussen M, Gilbert TP, Jones G. 2011. High-throughput sequencing offers insight into mechanisms of resource partitioning in cryptic bat species. Ecology and Evolution 1:556–570 DOI 10.1002/ece3.49.

Reiter J, Curio E. 2001. Home range, roost switching, and foraging area in a Philippine fruit bat (Ptenochirus jagori). Ecotropica 7:109–113.

Riaz T, Shehzad W, Viari A, Pompanon F, Taberlet P, Coissac E. 2011. ecoPrimers: inference of new DNA barcode markers from whole genome sequence analysis. Nucleic Acids Research 39:e145 DOI 10.1093/nar/gkr732.

Robinson JG. 1984. Diurnal variation in foraging and diet in the wedge-capped capuchin Cebus olivaceus. Folia Primatologica 43:216–228.

Scanlon AT, Petit S, Tuiwawa M, Naikatini A. 2014. High similarity between a bat-serviced plant assemblage and that used by humans. Biological Conservation 174:111–119.
Schmelitschek E, French KO, Parry-Jones K. 2009. Fruit availability and utilisation by grey-headed flying-foxes (Pteropodidae: Pteropus poliocephalus) in a human-modified environment on the south coast of New South Wales, Australia. *Wildlife Research* 36:592–600.

Soberón JM, Llorente JB, Onate L. 2000. The use of specimen-label databases for conservation purposes: an example using Mexican Papilionid and Pierid butterflies. *Biodiversity and Conservation* 9:1441–1466 DOI 10.1023/A:1008987010383.

Sritongchuay T, Gale GA, Stewart A, Keroka A, Bumrungsri S. 2014. Seed rain in abandoned clearings in a lowland evergreen rain forest in southern Thailand. *Tropical Conservation Science* 7:572–585 DOI 10.1177/194008291400700314.

Srivathsan A, Ang A, Vogler AP, Meier R. 2016. Fecal metagenomics for the simultaneous assessment of diet, parasites, and population genetics of an understudied primate. *Frontiers in Zoology* 13:17 DOI 10.1186/s12983-016-0150-4.

Srivathsan A, Sha J, Vogler AP, Meier R. 2015. Comparing the effectiveness of metagenomics and metabarcoding for diet analysis of a leaf-feeding monkey (*Pygathrix nemaeus*). *Molecular Ecology Resources* 15:250–261 DOI 10.1111/1755-0998.12302.

Start AN. 1974. The feeding biology in relation to food sources of nectarivorous bats (Chiroptera: Macroglossinae) in Malaysia. PhD thesis, University of Aberdeen, UK.

Stewart AB, Makowsky R, Dudash MR. 2014. Differences in foraging times between two feeding guilds within Old World fruit bats (Pteropodidae) in southern Thailand. *Journal of Tropical Ecology* 30:249–257 DOI 10.1017/S0266467414000042.

Stier SC, Mildenstein TL. 2005. Dietary habits of the world’s largest bats: the Philippine flying foxes, *Acerodon jubatus* and *Pteropus vampyrus lanensis*. *Journal of Mammalogy* 86:719–728 DOI 10.1644/1545-1542(2005)086[0719:DHOTWL]2.0.CO;2.

Sudhakaran MR, Doss PS. 2012. Food and foraging preferences of three pteropodid bats in southern India. *Journal of Threatened Taxa* 4:2295–2303 DOI 10.11609/JoTT.o2227.2295-303.

Tedman RA, Hall LS. 1985. The morphology of the gastrointestinal tract and food transit time in the fruit bats *Pteropus alecto* and *P. poliocephalus* (Megachiroptera). *Australian Journal of Zoology* 33:625–640 DOI 10.1071/ZO9850625.

Thomas DW. 2009. Analysis of diets of plant-visiting bats. In: Kunz TH, Parsons S, eds. *Ecological and behavioural methods for the study of bats*. Second edition. Baltimore: The Johns Hopkins University Press.

Valentini A, Miquel C, Nawaz MA, Bellemain EVA, Coissac E, Pompanon F, Gielly L, Cruaud C, Nasatti G, Wincker P, Swenson JE. 2009. New perspectives in diet analysis based on DNA barcoding and parallel pyrosequencing: the trnL approach. *Molecular Ecology Resources* 9:51–60 DOI 10.1111/j.1755-0998.2008.02352.x.

Weber N, Duengkae P, Fahr J, Dechmann D KN, Phensakul P, Khumbucha W, Siriroonrat B, Wacharapluesadee S, Maneenon P, Wikelski M, Newman S. 2015. High-resolution GPS tracking of Lyle’s flying fox between temples and orchards in central Thailand. *Journal of Wildlife Management* 79:957–968 DOI 10.1002/jwmg.904.

Williams N. SG, McDonnell MJ, Phelan GK, Keim LD, Van der Ree R. 2006. Range expansion due to urbanization: increased food resources attract Grey-headed

Aziz et al. (2017), *PeerJ*, DOI 10.7717/peerj.3176
Flying-foxes (*Pteropus poliocephalus*) to Melbourne. *Austral Ecology* **31**:190–198
DOI 10.1111/j.1442-9993.2006.01590.x.

**Win SS, Mya KM. 2015.** The diet of the Indian Flying Fox *Pteropus giganteus* (Brünnich, 1782) (Chiroptera: Pteropodidae) in Myanmar—conflicts with local people?. *Journal of Threatened Taxa* **7**:7568–7572 DOI 10.11609/JoTT.o4178.7568-72.

**Zeale MR, Butlin RK, Barker GL, Lees DC, Jones G. 2011.** Taxon-specific PCR for DNA barcoding arthropod prey in bat faeces. *Molecular Ecology Resources* **11**:236–244 DOI 10.1111/j.1755-0998.2010.02920.x.

**Zhang J, Kobert K, Flouri T, Stamatakis A. 2014.** PEAR: a fast and accurate Illumina Paired-End reAd MergeR. *Bioinformatics* **30**:614–620 DOI 10.1093/bioinformatics/btt593.