MATERNITY ROOST OF *Eptesicus brasiliensis* IN A LIANA IN THE SOUTHEAST PERUVIAN AMAZON

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ABSTRACT. We detected a maternity roost of *Eptesicus brasiliensis* in a large liana at Cocha Cashu Biological Station (Peru). Our objective was to describe the roost and its surrounding habitat employing visual methods to estimate colony size and time of foraging. At least 10 adults (9 ♂, 1 ♀) and one juvenile used a cavity in a liana that hung across a small trail. We used a night vision scope and infrared camera traps to note that bats left the roost for short foraging bouts (<1 hr). We also discuss the potential importance of lianas as roost sites in tropical forests.

Studies of roosting behavior and roost selection are important for conservation of forest bats (Betts, 1998). Some bats use cavities for roosting, breeding and hibernating, relying on roosting structures to provide protection from abiotic factors and potential predators. Most bats are secondary cavity users and select cavities in mainly live, dying, or dead trees, spending over half their lives within the roost environment (Kunz, 1982; Kunz and Lumsden, 2003; Diaz and Linares Garcia, 2012; Rengifo et al., 2013). The availability of roost sites is...
paramount to productivity and survival in cavity-roosting species. Anthropogenic land management activities, as well as natural weather events, commonly reduce roost availability in forest systems (Jones et al., 2001; Bennet et al., 2013; Tournant et al., 2013). There is a lower density of available snags for potential roosts in low latitude forests when compared to high latitude forests (Gibbs et al., 1993) potentially caused by lower rates of tree mortality, which would not favor snag creation, and higher rates of tree decomposition (Gibbs et al., 1993; Cornelius et al., 2008). Therefore, tree cavities may be a potentially unstable resource for bats in tropical systems. While some tree species have been identified as preferable for certain bat species (Evelyn and Stiles, 2003), lianas have been overlooked as potential roost sites for bats at critical life stages until now.

Nest site selection and placement is an important aspect of the natural history of a species and important for reproductive success (Goodall, 1962; Jessen et al., 2013). Persistent roost sites that are close to food sources and additionally promote a warm microclimate are particularly important for maternity roosts, which are used to suit the added demands of pregnancy and lactation and to bear and raise young (Kunz, 1982; Brigham and Fenton, 1986; Betts, 1998; Sedgeley, 2001). Since traditional tree cavities may be ephemeral, alternative and more stable vegetation types may also be used by bats as roosting sites, specifically for high cost activities such as those associated with reproduction and foraging (Sedgeley, 2001). Roost sites must also provide protection from potential predators that actively search for bats, such as carnivorous bats, primates, certain birds, and specifically those that develop a search image for roost structures (Lima and O'Keefe, 2013). It has been shown that primates are able to recognize leaves modified by tent-making bats (Boinski and Timm, 1985). The ease at which bats can leave and enter roosts will also determine the length of time of exposure to potential predators (Barclay et al., 1982; Fenton et al., 1994; VonHof and Barclay, 1996). The vertical location of roost sites within a forest may also be highly variable among and within species (Kalko and Handley, 2001), potentially hampering predator search image creation. Some species may also alleviate predation and potential roost decay through fission-fusion maternity roost systems, where many roost sites are used in the same area, with frequent switching among sites within a reproductive period (Brigham et al., 1997; Popa-Lisseanu et al., 2007).

The objective of this study was to describe the physical characteristics of a liana being used as an active maternity roost. We describe the physical characteristics and plant community surrounding the liana. We then report morphometric and reproductive state of the Eptesicus brasiliensis colony roosting within the liana and supply notes about their nocturnal activity patterns.

The E. brasiliensis roost was located at Cocha Cashu Biological Station (11°53'S, 71°26'W) along the Rio Manu, in the Madre de Dios region of southeastern Peru (Foster et al., 1986). This tropical floodplain forest is an example of a pristine successional plant community with disturbance caused by the rerouting of a meandering river and seasonal flooding, rather than anthropogenically (Terborgh, 1983; Foster et al., 1986). The area around Cocha Cashu Biological Station, within Manu National Park, has also remained free from organized hunting, thus high mammal diversity has maintained in this pristine forest (Janson and Emmons, 1990). An expedition directed at surveying mammal and bird species along the elevational gradient in Manu National Park detected 222 species of mammals, of which 92 were bats (Patterson et al., 2006).

The roost was situated in a late successional floodplain forest in a canopy liana (Bignoniaceae: Arribidaea florida; Vasquez and Rojas, 2004; C.F.F-N., pers. comm.) on a levee, 430 m from the start of trail KS (Fig. 1). The elevation at the roost site is 345 m above sea level and it is approximately 510 m from the Rio Manu and 950 m from Cocha Cashu, two permanent water sources. The forest is characterized by a short canopy (~25 m high), with successional trees, understory plants, and woody shrubs commonly associated with late regenerating forest. There were no standing hardwood trees approximately 200 years old, which are
characteristic of a lowland Amazonian forest of this area, such as *Cedrela odorata*, but there were several fallen and rotted *C. odorata* trees in the immediate vicinity of the roost location, aging the forest as greater than 200 but less than 400 years old (Vasquez and Rojas, 2004). At the roost location the levees were approximately 10 m apart and 1.5-2 m above ground level. The canopy plants located on the same levee as the liana roost included: Annonaceae: *Unonopsis floribunda*; Arecales: *Iriartea deltoidea* and *Socratea exorrhiza*; and Euphorbiaceae: *Sapium* sp. The understory plants located on the levee around the roost were: Piperaceae: *Piper* spp.; Marantaceae: *Calathea* sp.; Annonaceae: *Oxandra* sp.; Myristicaceae: *Otoba parvifolia*; Arecales: *Astryocaryum* sp.; and Theophrataceae: *Clavija tarapotana*. Canopy trees located in the depressions immediately surrounding the roost were: Rubiaceae: *Calycophyllum spruceanum*; Combretaceae: *Terminalia oblonga*; Moraceae: *Soracea* sp.; and Meliaceae: *Trichilia* sp. All of these woody tree species are characteristic of a late successional forest with seasonal flooding (Vasquez and Rojas, 2004).

On September 4, 2013, while walking on Trail KS (2 m wide), we heard audible squeaking sounds coming from a liana stretched across the trail. Upon inspection, we noticed a crack in the liana where apparently two separate vines may have grafted together and never fully closed, offering a small hollow as refuge for several bats, which were situated in the farthest corner of the hollow. The bats appeared to be of the genus *Eptesicus* and one
appeared very small and dull colored, as if a juvenile. The roost opening was 1.59 m above the ground; the crevice opening was 1.75 cm wide by 24.3 cm long. The diameter at the approximate centroid of the roost opening was 17.2 cm (Fig. 2).

On September 5, 2013, we set an Austbat© (Australia) style harp trap in front of the roost in the afternoon with factory strings re-tied every 2 cm to target small Vespertilionidae. We checked the trap 110 minutes after sundown to find ten individual *E. brasiliensis* roosting under the plastic roost baffle in the catch bag of the trap. All were adults (9 ♀, 1 ♂), identified using two field guides (Eisenberg and Redford, 1993; Reid, 2009), and no other bats were present in the trap. One adult female was lactating and we did not catch any juvenile bats, so the apparent juvenile may not have been volant. Forearm, ear, and weight measurements were taken on all bats using a metal ruler and digital pocket balance. All individuals were assessed for age by ossification of wing joints and reproductive condition by inspection of genitals and nipples (Kunz and Anthony, 1982), with stomach palpation for females (Table 1).

Exit counts were conducted on two separate nights with two methodologies to assess how effective our initial harp trapping method was in quantifying roost size. The first was on September 20, 2013, using a 4x Yukon NVMT3 night vision scope with a 50 mm lens, positioned 10 m away and pointing directly at the roost opening. Sunset at the roost location on September 20 was at 17:40. The total time of the exit count was 58 minutes and 44.07 seconds and six bats were observed leaving the roost and returning within the observation time (Table 2). We do not know if the first bat to leave was the first bat to return. However, all six bats exited the roost within a 13 minute time span and at least one bat foraged for at least 36 minutes before returning to the roost. It was observed that bats circled around the roost several times before re-entering it. The second method was 2 infrared (IR) motion video camera traps, Bushnell HD Trophy Camera Model 119537, which were placed facing the

![Fig. 2. Liana roost with trail KS in the background. There was only one apparent roost opening, indicated by the white arrow. The roost opening was 1.59 m above ground, the crevice opening was 1.75 cm wide by 24.3 cm long, and the diameter of the liana at the approximate centroid of the roost opening was 17.2 cm.](image)
Table 1
Data collected on individual bats from the harp trap survey. Ten adult *Eptesicus brasiliensis* were captured by harp trap. For comparison, mean values from this study were compared to measurements presented in two field guides. Reproductive condition codes are as follows: L = lactating, P = pregnant, PA = parous, NR = non-reproductive.

<table>
<thead>
<tr>
<th>Sex</th>
<th>Reproductive Condition</th>
<th>Ear (mm)</th>
<th>Forearm (mm)</th>
<th>Weight (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>♀</td>
<td>L</td>
<td>12</td>
<td>42</td>
<td>10.64</td>
</tr>
<tr>
<td>♀</td>
<td>PA</td>
<td>11</td>
<td>42</td>
<td>14.75</td>
</tr>
<tr>
<td>♀</td>
<td>PA</td>
<td>11</td>
<td>41</td>
<td>13.6</td>
</tr>
<tr>
<td>♀</td>
<td>P</td>
<td>11</td>
<td>41</td>
<td>14.75</td>
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<tr>
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<td>PA</td>
<td>11</td>
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<td>13.29</td>
</tr>
<tr>
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<td>PA</td>
<td>12</td>
<td>41</td>
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<tr>
<td>♀</td>
<td>P</td>
<td>11</td>
<td>39</td>
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<tr>
<td>♀</td>
<td>P</td>
<td>12</td>
<td>41</td>
<td>12.85</td>
</tr>
<tr>
<td>♀</td>
<td>P</td>
<td>11</td>
<td>41</td>
<td>14</td>
</tr>
<tr>
<td>♂</td>
<td>NR</td>
<td>10</td>
<td>40</td>
<td>9.87</td>
</tr>
</tbody>
</table>

Mean (this study) 11.2 40.9 13.03
Eisenberg & Redford 1999 15 40–41 9–10
Reid 2009 12–15 39–43 7–10

Note: The juvenile bat observed in the roost was not captured and the lactating female captured suggests that he was non-volant.

Table 2
Times of bat departure and return during the exit count using a night vision scope. Times reported here only represent one bat leaving or returning to the roost and do not refer to individual bats.

<table>
<thead>
<tr>
<th>Departure</th>
<th>Return</th>
</tr>
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<tbody>
<tr>
<td>17:51</td>
<td>18:01</td>
</tr>
<tr>
<td>17:52</td>
<td>18:13</td>
</tr>
<tr>
<td>17:55</td>
<td>18:18</td>
</tr>
<tr>
<td>17:56</td>
<td>18:21</td>
</tr>
<tr>
<td>17:58</td>
<td>18:22</td>
</tr>
<tr>
<td>18:04</td>
<td>18:40</td>
</tr>
</tbody>
</table>

roost exit at 15:30 on September 28, 2013, and retrieved at 07:30 on September 29, 2013. One was placed 1 m away directly in front of the entrance and the other was positioned 2.5 m to the west but also facing the roost exit. Cameras were set at the finest settings with one minute video recording upon detection of movement as an experimental effort to determine their use for unmanned exit counts at known roost sites. The camera placed 1 m in front of the roost detected two bats entering the roost: one at 18:18:56 and one at 18:31:13. The camera 2.5 m to the side detected one bat leaving the roost at 20:26:08 and one bat returning to the roost at 23:27:30. We suspect that the cameras missed many bats exiting and entering the roost site as these activities may happen faster than the cameras can be triggered to record (trigger speed = 0.6 sec). IR cameras that can continuously record for an entire night may be better suited for unmanned exit counts. However, the extra battery power required may make this unfeasible at remote field sites.

Considering the night vision survey and the IR camera survey, it appears that *E. brasiliensis* leaves the roost to forage shortly after sunset and returns within an hour. Though there are missing data from the IR camera survey, it appears that bats left the roost later in the night again for a foraging bout and were active until almost midnight, when one bat returned.

While some bat species use vine tangles as roost sites (Fenton et al., 2001) this is, to the best of our knowledge, the first record of roost site selection by bats in a liana cavity. Lianas (woody vines) are important components of tropical ecosystems, where they currently make up to 40% of the woody stems and more than 25% of the woody species (Schnitzer and Bongers, 2011). There is evidence that liana abundance and biomass is increasing in tropical forests which may increase the importance of lianas for bat roost sites (Schnitzer and Bongers, 2011). Since the 1970's biomass and abundance of...
Lianas in Neotropical forests have increased with some Amazonian forests showing a doubling in liana abundance in a 20-year period (Phillips et al., 2002; Schnitzer and Bongers, 2011). As liana abundance increases in Neotropical forests, tree diversity, recruitment, growth, fecundity, and survival will likely decrease (Schnitzer and Bongers, 2011), potentially making lianas a newly important roost site for bats.

Lianas may offer protection that larger tree cavities do not. Amazonian lianas may be more abundant than they were historically, therefore it is possible that predators do not have a search image for lianas as roost sites as they do for tree cavities. Neotropical lianas are smaller and less stable than paleotropical lianas and large lianas are less common in Neotropical forests than African forests (Emmons and Gentry, 1983). Smaller sized, more fragile lianas are presumably related to the evolution of prehensile tails in vertebrates that need to travel across canopy gaps (Emmons and Gentry, 1983). Neotropical lianas are used more for locomotion whereas African lianas are used more as a food source (Emmons and Gentry, 1983). Thus, it is logical that Neotropical predators may travel and search for food in the canopy rather than along the bases of lianas. Though lianas quite literally connect large trees in the canopy and provide canopy travel access for vertebrates (Charles-Dominique et al., 1981), the spatial lack of horizontal tree branches proximal to lower portions of large lianas may reduce predation potential for Neotropical bats roosting in liana cavities closer to the ground. Cavities on tree limbs or tree trunks may be more accessible to arboreal predators via horizontal branches.

The observed liana roost was located in the center of a 2 m wide trail. Therefore, it is possible that the maternity roost may be capturing a greater percentage of sunlight than other roost sites located in the dense forest understory. The retention of heat and absorption of solar radiation is particularly important for keeping pups warm when mothers leave the roost to forage (Sedgeley, 2001). The proximity of the roost site to the two permanent water sources may also be important if E. brasiliensis is foraging over water or drinking from those sources.

Lianas should not be overlooked as quality roosting sites in forests. Studies on roost selection in tropical forests should include lianas as potential roosting structures and researchers should add lianas to their search image repertoire. Similarly, forest management practices should consider the importance of lianas to bats in addition to other functions within the tropical forest system. Particularly, large Neotropical lianas should be investigated as a limiting resource on reproductive success in forest bats. As far as we know, this is the first documented report of a maternity roost of E. brasiliensis in lianas.

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