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Microhabitats of pholcid spiders (araneae: pholcidae) in Mt. Kitanglad Agri-Ecological and Techno-Demo Center, Imbayao, Bukidnon, Philippines

Eddie P. Mondejar*, Olga M. Nuñez

Department of Biological Sciences, MSU-Iligan Institute of Technology, Philippines

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Key words: Broad-leaf, Leaf-litter, Rock crevice, *Spathiphyllum sp.*, Species-specific.

Abstract

This study was conducted in Mt. Kitanglad Agri-Ecological and Techno-Demo Center, Brgy. Imbayao, Malaybalay, Bukidnon. The study aims to determine the microhabitats of pholcid spiders. Using Cruising method combined with visual search, pholcid spiders were searched on the underside of broad-leaf plants, buttresses, between rock and rock crevices, fallen logs, forest floor litter, and other possible microhabitats. Fourteen microhabitats were identified. Leaf litter and understory plant were the most species-rich with moderate diversity of pholcid spiders. High species richness of pholcid species was found between the paired microhabitats of bamboo sheath and understory plant, with degree of complementarity or distinctness value of 100%. This indicates that pholcid spiders tend to be more species-specific in the selection of microhabitats.

*Corresponding Author: Eddie P. Mondejar ✉ edmondejar2014@gmail.com

Introduction

Pholcidae is well adapted to various microhabitats such as under objects, dead leaves, rocks, logs on the ground, between buttresses, in little holes or caves, or on the underside of live leaves. The different species exhibit cryptic coloration that indicates avoidance with visually hunting predators, but desiccation resistance and wetness may be also involved. Most if not all pholcids are sedentary predators and exhibit nocturnality (Deeleman-Reinhold, 1986; Huber, 2009; Lapinski and Tschapka, 2014; Huber, 2014).

Microhabitat preferences are assumed to be adaptive in higher forms of organisms, and microhabitat-associated microclimate differs among species, thus, some species partition either microhabitat or microclimate within each guild in order to maintain the physiological equilibrium (Martin, 1998). However, it seems this is also true with some invertebrates like spiders. Litter-dwelling spiders for instance tend to aggregate in groups and use large, complex tree buttresses with more leaf litter than sparse litter that depends on the available herbaceous vegetation and variable microclimates (Cady, 1984; Chapin, 2014). In addition, some spiders show seasonal variability in a certain microhabitat that indicates the low rates of emergence of their prey (Yuen and Dudgeon, 2015).

Our current knowledge on other families of spiders may be of help in understanding the microhabitat preference of pholcid spiders. Many studies associate selection of microhabitat based on physical structure of host plants such as the leaf architecture that provides shelter, forage, and reproductive sites (Figueira and Vasconcellos-Neto, 1993; Harwood *et al.*, 2003; Harwood and Obrycki, 2005; de Omena and Romero, 2008; de Omena and Romero, 2010; Carvalho *et al.*, 2012).

However, only few studies on pholcid microhabitats were conducted. Huber and Schütte (2009) conducted a pioneering work in the study of microhabitat of two leaf dwelling pholcid spiders, *Metagonia uvita* and *M. osa*. The result showed that the two pholcid species prefer certain microhabitat and tend to be species-specific. Hence, this study was conducted to determine the microhabitats of pholcid spiders found in Imbayao, Malaybalay, Bukidnon.

Materials and methods

The Mt. Kitanglad Agri-Ecological Techno-Demo Center (MKAETDC) is located in Brgy. Imbayao, Malaybalay City, Bukidnon at 8°2'12"N latitude and 125°02'03"E longitude (Fig. 1).

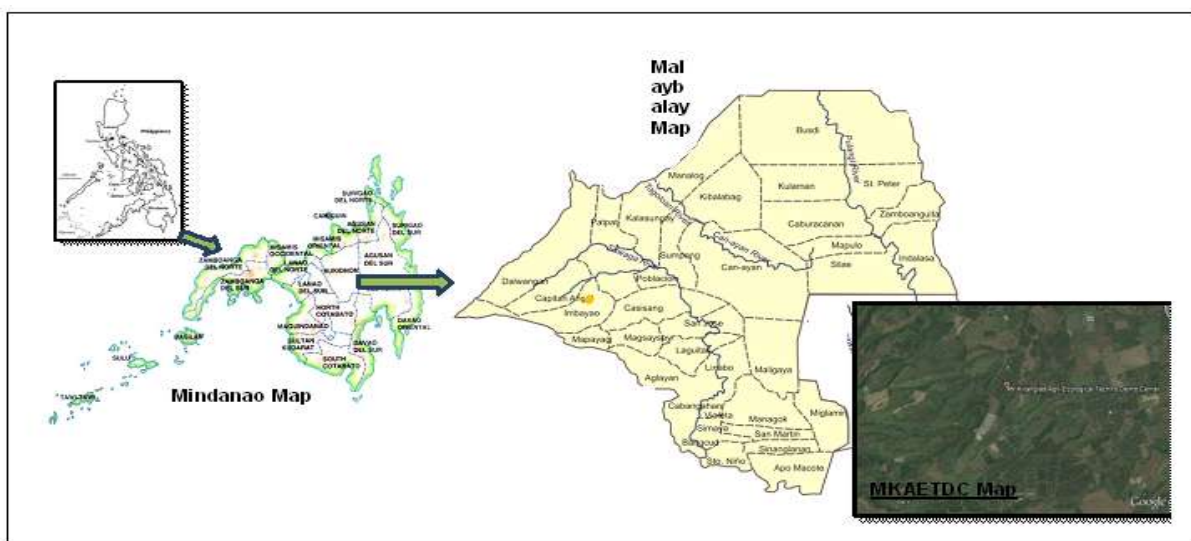


Fig. 1. Map showing the location of the sampling sites in Mt. Kitanglad Agri-Ecological and Techno-Demo Center, Imbayao, Malaybalay, Bukidnon (www.ivoteph.com, 2013; www.googleearth.com, 2015).

Sampling site was selected according to the vegetation and distance to water source. Habitat was described using the Habitat Description Form (Heaney, 2001) and vegetation cover was estimated using chart for estimation of foliage cover (Luttmerding *et al.*, 1990). Four sampling sites were established. The first site was within the natural forest and ecological walk area. The second site was within the abaca experimental site and the Natural forest. The third site was within the rain forestation area while the fourth site was between the Ceremonial ground area and the buffer zone of Mt. Kitanglad Range Natural Park, Bukidnon.

Collection, processing, and identification of samples

The collection of pholcid spiders was done on February 9-10, 2015 for a total of 24 man-hours. The specimens were collected using the cruising method combined with visual search of possible microhabitats such as under fallen and decaying logs, leaf litter, on the underside of live leaves, tree buttresses, rock surface and crevice, and other microhabitats where pholcid spiders are usually found, with a maximum time limit of two hours in each site. Hand collecting was used to capture the pholcid spiders according to microhabitat. The specimens were photographed in their microhabitat before putting them into the collecting jar. Specimens were then placed in Eppendorf tubes containing 95% ethanol. All collected unidentified specimens in the field were taken as voucher specimens for identification. Surveys were conducted in the morning between 700

hrs to 1100 hrs and in the afternoon and evening from 1500 hrs to 1900 hrs. Collection of samples was carried out as soon as all the necessary local and DENR permits were available. Specimens were identified by Dr. Bernhard Huber of the Alexander Koenig Zoological Research Museum, Germany. Each specimen that is probable new species was assigned a specific code in order to facilitate future record tracking.

Statistical analysis

Data analyses were performed using Paleontological Statistics software (PAST) package version 2.17c (Hammer *et al.*, 2001) for Multivariate analysis - Seriation; Statistical Process Control software (SPC) for excel to calculate ANOVA; and Microsoft excel 2010 software to determine the degree of complementarity or distinctness between microhabitats, given the formula $C_{jk} = U_{jk}/S_{jk}$, where U_{jk} is the number of species unique to either site, and S_{jk} is the species richness for both sites combined; $U_{jk} = S_j + S_k - 2V_{jk}$, and $S_{jk} = S_j + S_k - V_{jk}$; S = the number of species in sites j and k , and V_{jk} = the number of species shared by sites j and k (Colwell and Coddington, 1994).

Results and discussion

Fourteen plant microhabitats were identified for the pholcid spiders in Imbayao, Malaybalay, Bukidnon (Table 1).

Table 1. Microhabitats of Pholcid spiders in Imbayao, Bukidnon.

Microhabitats	No. of individuals	Species Richness, S_{obs}	Shannon, H'	Evenness, E
Bamboo sheath	5	3	1.055	0.9572
<i>Schimatoglottis sp.</i>	3	1	0	1
Understory plant	11	4	1.295	0.9123
Leaf litter	10	4	1.168	0.8041
Ground fern	4	1	0	1
Ground/Soil	1	1	0	1
Decaying <i>Calamus</i>	1	1	0	1
<i>Dracaena sp.</i>	33	2	0.2286	0.6284
<i>Spathiphyllum sp.</i>	9	2	0.5297	0.8492
<i>Molinaria sp.</i>	2	2	0.6931	1
<i>Cordyline sp.</i>	3	1	0	1
<i>Calamus sp.</i>	1	1	0	1
<i>Livistona sp.</i>	1	1	0	1
<i>Medinilla sp.</i>	1	1	0	1

The most utilized microhabitats of pholcid spider species were leaf litter and understory plant, with four species each. In leaf litter, *Belisana sp.PSt365*, *Belisana sp.Phi129?*, *Holocneminus sp.PSt1291*, and

Spermophora sp.Phi109 were found. In understory plants, *Belisana sp.PSt365*, *Belisana sp.Phi129?*, *Calapnita sp.Phi90?*, and *Panjange sp.Phi125?* were documented.

Table 2. Degree of Complementarity (C) between microhabitats of Pholcid spiders. S_{jk} = combined species richness; V_{jk} = number of species shared between two microhabitats.

Microhabitats	Observed				V_{jk}	S_{jk}	Distinctness (%)	
	Species Richness							
	S_j		S_k					
Bamboo sheath	-	<i>Schismatoglottis sp.</i>	3	vs	1	0	4	100
Bamboo sheath	-	Understory plant	3	vs	4	0	7	100
Bamboo sheath	-	Leaf litter	3	vs	4	1	6	83
Bamboo sheath	-	Ground fern	3	vs	1	0	4	100
Bamboo sheath	-	<i>Dracaena sp.</i>	3	vs	2	0	5	100
Bamboo sheath	-	<i>Spathiphyllum sp.</i>	3	vs	2	0	5	100
Bamboo sheath	-	<i>Molinieria sp.</i>	3	vs	2	0	5	100
Bamboo sheath	-	<i>Cordyline sp.</i>	3	vs	1	0	4	100
Bamboo sheath	-	<i>Calamus sp.</i>	3	vs	1	0	4	100
Bamboo sheath	-	<i>Livistona sp.</i>	3	vs	1	0	4	100
Bamboo sheath	-	<i>Medinilla sp.</i>	3	vs	1	0	4	100
<i>Schismatoglottis sp.</i>	-	Leaf litter	1	vs	4	0	5	100
<i>Schismatoglottis sp.</i>	-	Ground/Soil	1	vs	1	0	2	100
<i>Schismatoglottis sp.</i>	-	Decaying <i>Calamus</i>	1	vs	1	0	2	100
<i>Schismatoglottis sp.</i>	-	<i>Dracaena sp.</i>	1	vs	2	0	3	100
<i>Schismatoglottis sp.</i>	-	<i>Molinieria sp.</i>	1	vs	2	0	3	100
<i>Schismatoglottis sp.</i>	-	<i>Cordyline sp.</i>	1	vs	1	0	2	100
<i>Schismatoglottis sp.</i>	-	<i>Calamus sp.</i>	1	vs	1	0	2	100
Understory plant	-	Ground/Soil	4	vs	1	0	5	100
Understory plant	-	Decaying <i>Calamus</i>	4	vs	1	0	5	100
Leaf litter	-	Ground fern	4	vs	1	0	5	100
Leaf litter	-	<i>Dracaena sp.</i>	4	vs	2	1	5	80
Leaf litter	-	<i>Spathiphyllum sp.</i>	4	vs	2	1	5	80
Leaf litter	-	<i>Molinieria sp.</i>	4	vs	2	1	5	80
Leaf litter	-	<i>Calamus sp.</i>	4	vs	1	0	5	100
Leaf litter	-	<i>Livistona sp.</i>	4	vs	1	0	5	100
Leaf litter	-	<i>Medinilla sp.</i>	4	vs	1	0	5	100
Ground fern	-	Ground/Soil	1	vs	1	0	2	100
Ground fern	-	Decaying <i>Calamus</i>	1	vs	1	0	2	100
Ground fern	-	<i>Dracaena sp.</i>	1	vs	2	0	3	100
Ground fern	-	<i>Molinieria sp.</i>	1	vs	2	0	3	100
Ground fern	-	<i>Cordyline sp.</i>	1	vs	1	0	2	100
Ground fern	-	<i>Calamus sp.</i>	1	vs	1	0	2	100
Ground/Soil	-	Decaying <i>Calamus</i>	1	vs	1	0	2	100

Ground/Soil	-	<i>Dracaena sp.</i>	1	vs	2	0	3	100
Ground/Soil	-	<i>Spathiphyllum sp.</i>	1	vs	2	0	3	100
Ground/Soil	-	<i>Molineria sp.</i>	1	vs	2	0	3	100
Ground/Soil	-	<i>Cordyline sp.</i>	1	vs	1	0	2	100
Ground/Soil	-	<i>Calamus sp.</i>	1	vs	1	0	2	100
Ground/Soil	-	<i>Livistona sp.</i>	1	vs	1	0	2	100
Ground/Soil	-	<i>Medinilla sp.</i>	1	vs	1	0	2	100
Decaying Calamus	-	<i>Dracaena sp.</i>	1	vs	2	0	3	100
Decaying Calamus	-	<i>Spathiphyllum sp.</i>	1	vs	2	0	3	100
Decaying Calamus	-	<i>Molineria sp.</i>	1	vs	2	0	3	100
Table 2 continued...								
Decaying Calamus	-	<i>Cordyline sp.</i>	1	vs	1	0	2	100
Decaying Calamus	-	<i>Calamus sp.</i>	1	vs	1	0	2	100
Decaying Calamus	-	<i>Livistona sp.</i>	1	vs	1	0	2	100
Decaying Calamus	-	<i>Medinilla sp.</i>	1	vs	1	0	2	100
<i>Dracaena sp.</i>	-	<i>Livistona sp.</i>	2	vs	1	0	3	100
<i>Dracaena sp.</i>	-	<i>Medinilla sp.</i>	2	vs	1	0	3	100
<i>Spathiphyllum sp.</i>	-	<i>Calamus sp.</i>	2	vs	1	0	3	100
<i>Molineria sp.</i>	-	<i>Livistona sp.</i>	2	vs	1	0	3	100
<i>Molineria sp.</i>	-	<i>Medinilla sp.</i>	2	vs	1	0	3	100
<i>Cordyline sp.</i>	-	<i>Calamus sp.</i>	1	vs	1	0	2	100
<i>Cordyline sp.</i>	-	<i>Livistona sp.</i>	1	vs	1	0	2	100
<i>Cordyline sp.</i>	-	<i>Medinilla sp.</i>	1	vs	1	0	2	100
<i>Calamus sp.</i>	-	<i>Livistona sp.</i>	1	vs	1	0	2	100
<i>Calamus sp.</i>	-	<i>Medinilla sp.</i>	1	vs	1	0	2	100

Field observation showed that some of the species of *Belisana* were found in leaf litter, but this could possibly be an accidental encounter during field sampling because this genus is known to inhabit in live leaves and also reported to occur in caves (Huber, 2005; Tong and Li, 2009).

The diversity of pholcid species was moderate in understory plants ($H' = 1.295$), leaf litter ($H' = 1.168$), and in *bamboo sheath* ($H' = 1.055$). The abundance of pholcid spiders varies with different microhabitats. *Belisana sp. Phi129?* showed dominance value of 36% in *Dracaena sp.* and lowest in understory plant and in the plant, *Molineria sp.* (1.18%). The observed dominance of one pholcid species might be due to the available resources that favor its specific resource requirements (Redborg and Redborg, 2000; Mallis and Hurd, 2005; Butt and Tahir, 2010).

Table 2 shows paired microhabitats which exhibit a high level of complementarity value ($C = 100\%$), accounted for 59% of the total number of paired microhabitats.

This result demonstrates resource partitioning among species of pholcid spiders which may be in response to interspecific competition, differences in microclimate, behavior, and ecological niche (Schoener, 1974; Hanski and Koskela, 1979; Morrison and With, 1987; Walter, 1991; Amezaga and Rodriguez, 1998; Redborg and Redborg, 2000; Albrecht and Gotelli, 2001; Blackledge *et al.*, 2003; Sushma and Singh, 2006; Butt and Tahir, 2010).

Eight percent of other paired microhabitats exhibited sharing of the observed pholcid spiders, with complementarity value of 0%.

Pholcid Species	Microhabitats													
	A	B	C	D	E	F	G	H	I	J	K	L	M	N
<i>Aetana sp.</i>	■													
<i>Belisana sp. PSt365</i>			■	■										
<i>Holocneminus sp. PSt1291</i>	■			■		■								
<i>Spermophora sp. Phi109</i>	■			■			■							
<i>Belisana sp. Phi129?</i>			■	■				■	■	■	■			
<i>Panjange sp. Phi125?</i>		■	■		■				■				■	■
<i>Calapnita sp. Phi90?</i>			■					■		■		■		

Legend for Microhabitats:

- A Bamboo sheath
- B *Schismatoglottis sp.*
- C Understory plant
- D Leaf litter
- E Ground fern
- F Ground/Soil
- G Decaying *Calamus*
- H *Dracaena sp.*
- I *Spathiphyllum sp.*
- J *Molineria sp.*
- K *Cordyline sp.*
- L *Calamus sp.*
- M *Livistona sp.*
- N *Medinilla sp.*

Fig. 2. Seriation Analysis of Microhabitats of Pholcid spiders.

This implies strong overlap in both food and habitat utilization, not because of insufficient food but rather due to the abundance of food, vegetation cover, and relative species abundance (Bengtson, 1984; Kozłowski *et al.*, 2008).

Fig. 2 shows that 16% of the collected pholcid species in Mt. Kitanglad Agri-Ecological and Techno-Demo Center (MKAETDC), Imbayao, Bukidnon were mostly found in leaf litter and in understory plant.

The distribution of litter-dwelling pholcid spiders may be relevant to the foraging mode of prey-predator interactions in leaf litter and reproduction (Casas *et al.*, 2008; Hebets *et al.*, 2008). In addition, a study has also demonstrated that different microhabitat structures (e.g. inflorescence) influence the distribution of spiders (de Souza and Modena, 2004; Rocha-Filho and Rinaldi, 2011).

Conclusion

Among the 14 identified microhabitats, leaf litter and understory plant were found to be the most utilized microhabitats of the pholcid spiders. The degree of complementarity between microhabitats of pholcid spiders was highly significant among paired microhabitats in Mt. Kitanglad Agri-Ecological and Techno-Demo Center, Brgy. Imbayao, Malaybalay, Bukidnon. Species diversity was moderate which

indicates that microhabitats may play a role in species composition of pholcid spiders.

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References

Albrecht M, Gotelli NJ. 2001. Spatial and Temporal niche partitioning in grassland ants. *Oecologia* **126**, 134-141.

Amezaga I, Rodriguez MA. 1998. Resource partitioning of four sympatric bark beetles depending on swarming dates and tree species. *Forest Ecology and Management* **109**, 127-135.

Bengtson DA. 1984. Resource partitioning by *Menidia menidia* and *Menidia beryllina* (Osteichthyes:Atherinidae). *Marine Ecology – Progress Series* **18**, 21-30.

- Blackledge TA, Binford GJ, Gillespie RG.** 2003. Resource use within a community of Hawaiian spiders (Araneae: Tetragnathidae). *Annales Zoologici Fennici* **40**, 293-303.
- Butt A, Tahir HM.** 2010. Resource Partitioning among five Agrobiont Spiders of a Rice Ecosystem. *Zoological Studies* **49**, 470-480.
- Cady AB.** 1984. Microhabitat selection and locomotor activity of *Schizocosa ocreata* (Walckenaer) (Araneae, Lycosidae). *Journal of Arachnology* **11**, 297-307.
- Carvalho LS, Gomes JO, Neckel-Oliveira S, Lo-Man-Hung NF.** 2012. Microhabitat use and intraspecific associations in the whip spider *Heterophrynus longicornis* (Arachnida: Amblypygi) in forest fragments formed by the Tucuruí Dam lake, Pará, Brazil. *Journal of Natural History* **46**, 1263-1272.
- Casas J, Steinmann T, Dangles O.** 2008. The Aer-dynamic Signature of Running Spiders. *PLoS ONE* **3**, e2116.
- Chapin KJ.** 2014. Microhabitat and spatial complexity predict group size of the whip spider *Heterophrynus batesii* in Amazonian Ecuador. *Journal of Tropical Ecology* **30**, 173-177.
- Colwell RK, Coddington JA.** 1994. Estimating Terrestrial Biodiversity through extrapolation. *Philosophical Transactions of the Royal Society of London, B* **345**, 101-118.
- De Omena PM, Romero GQ.** 2008. Fine-scale microhabitat selection in a bromeliad-dwelling jumping spider (Salticidae). *Biological Journal of the Linnean Society* **94**, 653-662.
- de Omena PM, Romero GQ.** 2010. Using visual cues of microhabitat traits to find home: the case study of a bromeliad-living jumping spider (Salticidae). *Behavioral Ecology* **21**, 690-695.
- de Souza ALT, Modena EDS.** 2004. Distribution of Spiders on different types of inflorescences in the Brazilian pantanal. *Journal of Arachnology* **32**, 345-348.
- Deeleman-Reinhold CL.** 1986. Leaf-dwelling Pholcidae in Indo-Australian rain forests. *International Congress of Arachnology (Panama, 1983)* **9**, 45-48.
- Figueira JEC, Vasconcellos-Neto J.** 1993. Reproductive success of *Latrodectus geometricus* (Theridiidae) on *Paepalanthus bromelioides* (Eriocaulaceae): Rosette size, Microclimate, and Prey Capture. *Ecotropicos* **5**, 1-10
- Hammer Ø, Harper D, Ryan PD.** 2001. Past: Paleontological Statistics Software Package for Education and Data Analysis. *Palaeontologia Electronica* **4**, 9.
- Hanski I, Koskela H.** 1979. Resource partitioning in six guilds of dung-inhabiting beetles (Coleoptera). *Annals of Entomologica Fennica* **45**, 1-12
- Harwood JD, Obrycki JJ.** 2005. Web-Construction Behavior of Linyphiid Spiders (Araneae, Linyphiidae): Competition and Co-Existence within a Generalist Predator Guild. *Journal of Insect Behavior* **18**, 593-607
- Harwood JD, Sunderland KD, Symondson WOC.** 2003. Web-location by linyphiid spiders: prey-specific aggregation and foraging strategies. *Journal of Animal Ecology* **72**, 745-756.
- Hebets EA, Elias MO, Mason AC, Miller GL, Stratton GE.** 2008. Substrate-dependent signaling success in the wolf spider, *Schizocosa retrorsa*. *Animal Behaviour* **75**, 605-615.
- Heaney LR.** 2001. *Island Biogeography, Paradigm*

Lost? *Frontiers of Biogeography* **2**, 94-97

Huber BA. 2005. Revision of the genus *Spermophora* Hentz in Southeast Asia on the Pacific Islands, with descriptions of the three new genera (Araneae: Pholcidae). *Zool. Med. Leiden* **79**, 61-114.

Huber BA. 2009. Life on leaves: Leaf-dwelling Pholcids of Guinea, with emphasis on *Crossopriza cylindrogaster*, a spider with inverted resting position, pseudo eyes, lampshade web, and tetrahedral egg-sac (Araneae: Pholcidae). *Journal of Natural History* **43**, 2491-2523.

Huber BA, Schütte A. 2009. Preliminary notes on leaf-dwelling Metagonia spiders (Araneae: Pholcidae) in the Esquinas Rainforest near La Gamba, Costa Rica: leaf preference, mimesis, and web structure. *Contributions to Natural History (Bern)* **12**, 681-697.

Huber BA. 2014. Pholcidae. In: *Biodiversidad de Artrópodos Argentinos*, Vol. 3 (eds. S. Roig-Juñent, L.E. Claps, J.J. Morrone), 131-140. *Sociedad Entomológica Argentina*.

Kozłowski AJ, Gese EM, Arjo WM. 2008. Niche Overlap and Resource Partitioning Between Sympatric Kit Foxes and Coyotes in the Great Basin Desert of Western Utah. *The American Midland Naturalist* **160**, 191-208.

Lapinski W, Tschapka M. 2014. Desiccation resistance reflects patterns of microhabitat choice in a Central American assemblage of wandering spiders. *Journal of Experimental Biology* **217**, 2789-2795.

Luttmerding HA, Demarchi DA, Lea EC, Vold T, Meidinger DV. 1990. Describing ecosystems in the field. Second edition. *British Columbian Ministry. Environment, Lands and Parks and British Columbian Ministry of For MOE Manual 11*. Victoria, B.C. 213 P.

Mallis RE, Hurd LE. 2005. Diversity among

ground-dwelling spider assemblages: Habitat Generalists and Specialists. *Journal of arachnology* **33**, 101-109

Martin TE. 1998. Are Microhabitat Preferences of Co-existing Species under selection and Adaptive? *Ecology* **79**, 656-670.

Morrison ML, With KA. 1987. Inter-seasonal and Intersexual Resource Partitioning in Hairy and White-headed Woodpeckers. *The Auckland* **104**, 225-233

Redborg KE, Redborg AH. 2000. Resource Partitioning of Spider Hosts (Arachnida, Araneae) by two Mantispid Species (Neuroptera, Mantispidae) in an Illinois Woodland. *Journal of Arachnology* **28**, 70-78.

Rocha-Filho LC, Rinaldi IMP. 2011. Crab spiders (Araneae: Thomisidae) in flowering plants in a Brazilian "Cerrado" ecosystem. *Brazilian Journal of Biology* **71**, 359-364.

Schoener TW. 1974. Resource Partitioning in Ecological Communities. *Science, New Series* **185**, 27-39.

Sushma HS, Singh M. 2006. Resource partitioning and interspecific interactions among sympatric rain forest arboreal mammals of the Western Ghats, India. *Behavioral Ecology* **17(3)**, 479-490.

Tong YF, Li SQ. 2009. Six new cave-dwelling pholcid spiders (Araneae: Pholcidae) from Hainan Island, with two newly recorded genera from China. *Zootaxa* **1988**, 17-32.

Walter GH. 1991. What is resource partitioning. *Journal of Theoretical Biology* **150**, 137-143.

www.googleearth.com. 2015. Philippines. Retrieved from www.googleearth.com.

www.ivoteph.com. 2013. Malaybalay City Barangay Elections 2013, Results. Retrieved from <http://www.ivoteph.com/barangayelections/malaybalay-city-barangay-elections-2013-results/>

Yuen E, Dudgeon D. 2015. Spatio-temporal variability in the distribution of ground-dwelling riparian spiders and their potential role in water-to-land energy transfer along Hong Kong forest streams. *PeerJ* **3**, e1134.