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IS INCREASED RESISTANCE TO PARASITISM AT THE ORIGIN OF POLYGYNY IN A MEXICAN POPULATION OF THE ANT ECTATOMMA TUBERCULATUM (HYMENOPTERA: FORMICIDAE)?

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ABSTRACT

The Neotropical ant Ectatomma tuberculatum (Olivier) shows a polymorphic social organization with strictly monogynous or facultative polygynous populations along its distribution range. The causative factors of these variations in social organization are unknown but biotic and abiotic environmental factors might explain the differences. In particular, recent studies have suggested that parasites and parasitoids could have an impact on host colony phenotype. Here, we report on eucharitid wasps and mermithid nematodes that attack the only known Mexican polygynous population of E. tuberculatum in Veracruz, Mexico. The results were compared with those previously obtained from a monogynous population in Chiapas, Mexico; the aim being to investigate environmental influences on colony social organization. The rather similar prevalence of parasitism between the 2 populations suggests that the hypothesis of an increased resistance to parasites and parasitoids as one of the factors responsible for the emergence of polygyny in E. tuberculatum populations does not apply. More likely, other kinds of environmental stress or constraints facilitate the development of polygyny, such as nest site limitations, risks of dispersal and/or high rates of predation.

Key Words: ectatommine ants, prevalence of parasitism, social structure, facultative polygyny, nest site limitation, Mexico

RESUMEN

La hormiga neotropical Ectatomma tuberculatum (Olivier) presenta una organización social polimorfa, con poblaciones estrictamente monogámicas o bien, polígámicas facilitativas a lo largo de su rango de distribución. Los factores responsables de estas variaciones en su organización social se desconocen, pero factores ambientales bióticos y abióticos podrían explicar estas diferencias. En particular, estudios recientes sugieren que parásitos y parasitoides pueden afectar el fenotipo de las colonias hospederas. Aquí reportamos sobre avispas eucharítidas y nemátodos mermitidos que atacan a la única población polígina de E. tuberculatum conocida para México, en Veracruz. Se comparan estos resultados con los que se obtuvieron anteriormente de una población monógama de Chiapas (México), con el objetivo de analizar las influencias ambientales sobre la organización social de las colonias. La prevalencia de parasitismo bastante similar entre ambas poblaciones sugiere que la hipótesis de una mayor resistencia a parásitos y parasitoides como uno de los factores responsables de la aparición de la polígina en E. tuberculatum no es válida. Parece más probable que otras clases de estrés o limitantes ambientales estén implicadas, tales como limitación en sitios de nidificación, riesgos de dispersión o altas tasas de depredación.

Translation provided by the authors.
Variation in social organization (the number of individuals in a social group, their behavioral and genetic relationships, and the way in which reproduction is partitioned among them) has been regarded as the product of diverse extrinsic and intrinsic selection pressures related both to local ecological factors and to competitive and cooperative interactions among group members (Ross & Keller 1995; Smith et al. 2008). By comparison with the numerous studies focused on behavioral and genetic mechanisms (see for example: Keller 1993; Chan et al. 1999; Clark et al. 2006; Goodisman et al. 2008; Gotzek & Ross 2008; Haag-Liautard et al. 2009; Suni & Gordon 2010), little attention has been paid to biotic environmental factors such as predation, competition, parasitism, or disease. However, such environmental factors are known to be important in structuring ant communities and populations. Predation by army ants on other ant species, for example, is thought to be a major factor affecting leaf litter community structure and life-history traits of ant colonies in the neotropics (Kaspari & O'Donnell 2003; O'Donnell et al. 2007), while high densities of predators such as larval ant lions (Gotelli 1996) or lizards (Huang et al. 2008) can strongly alter ground-foraging ant assemblages. Furthermore, various studies have shown that attacks by parasitoid phorid flies (Peener 2000; Philpott et al. 2009), eucharitid wasps (Lachaud & Pérez-Lachaud 2009; Pérez-Lachaud et al. 2010), or pathogens (Keller 1995b; Gardner et al. 2007; Nonacs 1988; Keller 1995b). Differing hypotheses have been generated to account for variation in queen number including competition, nest site limitation, food availability, risks of dispersal, and inter- and intraspecific social parasitism (Keller 1995b; Gardner et al. 2007; McGlynn 2010). Polygyny and/or multiple mating (polyandry) are thought to be beneficial to the colony by increasing resistance to parasites and pathogens due to an enhancement in the genetic diversity of the workers (Schmid-Hempel 1994, 1998; Hughes et al. 2008). More specifically, genetically diverse host populations are predicted to have lower levels of infection or parasitism prevalence, but studies addressing this issue are scant (Hughes & Boomsma 2004; Gardner et al. 2007; Reber et al. 2008), and the results are not always consistent (see Castella et al. 2010).

Ectatomma tuberculatum (Olivier) (Hymenoptera: Formicidae: Ectatomminae) is a Neotropical ground-nesting, predatory ant that occurs in both wet forest and dry forest habitats from Mexico to northern Argentina, and from sea level to about 1500 m in altitude (Weber 1946; Brown 1958; Valenzuela et al. 1995; Quiroz-Robledo & Valenzuela-González 2007). This species, regarded as a potential biocontrol agent in different agroecosystems (Cook 1904a, b; Weber 1946; Lachaud et al. 1996; Ibarra-Núñez et al. 2001), is ecologically dominant in coffee and cocoa plantations in Chiapas, Mexico (Schatz & Lachaud 2008), and in the arboreal ant mosaic of cocoa plantations in Brazil (Majer et al. 1994). Colony structure and social organization of E. tuberculatum populations seem to vary according to as yet unidentified regional causative factors. For example, the population at Rosario Izapa, in the Soconusco region of Chiapas, Mexico, is characterized by the presence of a single, large queen per colony, well-differentiated from the workers (Pérez-Lachaud et al. 2010); from 253 nests of E. tuberculatum collected from the whole Soconusco region over 16 yr, only 3 were digynous and one contained 4 non-functional, mermithised dealate females (Lachaud, unpublished data). At that site, nest population ranged from 22 to 1849 adults (mean colony size = 515, n = 104, Lachaud & Pérez-Lachaud, unpublished data), and nest density was estimated at between 600 and 1500 nests/ha (Schatz & Lachaud 2008). In contrast, facultative polygyny (colonies coexisting as monogynous or polygynous in the same population) is characteristic of populations in Brazil (Hora et al. 2005b), but also in a population found at Apazapan, Veracruz, Mexico (Hora et al. 2001, 2005a), where nest population ranged from 6 to 429 adults (mean colony size = 106, n = 29, recalculated from Hora et al. 2001).

Because of the clearly established differences between the 2 Mexican E. tuberculatum populations studied so far (Chiapas and Veracruz), some doubts emerged concerning species identification. However, the results of current genetic polymorphism analyses using specific microsatellite loci (see Poteaux et al. 2003) confirmed that both populations belong to the same species and only differ by a reduced number of nucleotide base pairs (Nettel et al., unpublished data). Different selective environmental pressures have been put forward to explain the observed differences in social organization (Hora et al. 2005b; Zinck et al. 2007, 2008), including differences in genetic diversity.
and resistance to parasitism (Pérez-Lachaud et al. 2010). Though the 2 above-mentioned populations have been the object of extensive studies, no comparative study has been made concerning the diversity of parasites attacking them.

Here we report on 2 groups of parasites, eucharitid wasps and mermithid nematodes, which attack the polygynous E. tuberculatum population of Apazapan, Veracruz, with the aim of adding information on the ecology of this species in order to better understand the mechanisms underlying facultative polygyny.

**MATERIALS AND METHODS**

**Study Site**

The study site was located at Apazapan, Veracruz, Mexico (19°19'38"N; 96°43'21"W, 300 m above sea level); nests were collected at the same site (referred to as Apz 1, see Hora et al. 2005a) where both polygynous and monogygous E. tuberculatum colonies infested by Ectatomma parasiticum Feitosa & Fresneau had been previously collected by Hora et al. (2001). This site is a remnant of tropical dry forest and has an irregular orography. Surrounding natural vegetation is characteristic of lowland deciduous scrub (Castillo 1995). The climate of the region is warm, subhumid, with heavy rains during the summer months and at the beginning of autumn, and sparse rains during winter. The annual average temperature is 25.5°C with an average annual total rainfall of 1250 mm. The climate of the region is characteristic of lowland deciduous scrub (Castillo 1995).

**Sampling Method for Eucharitid Wasps**

Eucharitidae (Hymenoptera: Chalcidoidea) are specific, specialized ectoparasitoids of ant brood, with a highly modified life cycle (Clausen 1941; Heraty 1994). Adults emerge inside the host nest, mating occurs outside, females lay eggs into or on plant tissue, and the very mobile first-instar larvae (planidia) are presumed to attach to foraging ant workers (or to intermediate hosts) for passive phoretic transport to the ant nest (Clausen 1940; Heraty 2000). The planidia actively seek the suitable developmental host stage (Clausen 1940, 1941; Heraty 1994; personal observation) and further development takes place on the newly transformed ant pupae (Clausen 1940; Pérez-Lachaud et al. 2006b).

Nine nests of Ectatomma tuberculatum were thoroughly excavated on 21 Sep and 23 Sep 2010 and taken to the laboratory. The presence or absence of alate and dealate females in each nest was noted, as well as the number of males, workers, cocoons (pupae), larvae, and the presence of eggs. Cocoons were isolated in plastic containers topped with wire mesh and kept at room temperature (28 ± 2°C and 75 ± 5% relative humidity) for about one week to wait for parasitoid emergence. Then the identity, number, and sex of emerging adult wasps were recorded. After this period, all the cocoons were dissected under a stereomicroscope (Nikon, 75x), and both the number and stage of development of parasitoids (planidia, late larvae, pupae and pharate adults) were recorded. Larvae were also inspected, and the presence of round melanized scars (evidence of the previous attachment of a planidium), and both the number and location of planidia, if present, were recorded.

**Sampling Method for Mermithid Nematodes**

No complete life cycle of any mermithid parasite of Neotropical ants is known; but in a manner similar to the European mermithid Pheromeris villosa Kaiser that parasitizes sexuals of Lasius flavus (Fab.) and L. niger (L.) (Kaiser 1986), they are thought to have an indirect life cycle; i.e., the ant—which corresponds to the developmental host—becomes infected by eating paratenic hosts containing infective stages of the mermithid. Infection of the ant occurs in the larval stage in that the mermithid juvenile enters the body cavity of the ant larva and initiates development. Known mermithids infecting Ectatomma species (Wheeler 1930; Weber 1946; Poinar et al. 2006) are carried into the adult stage of the ant. Upon emergence from the ant, the post-parasitic juvenile enters an aquatic or moist habitat to mature, mate, and oviposit (Poinar et al. 2006).

Adult workers parasitized by mermithids were recognized from the morphological modifications of the ants brought about by the parasites and were maintained in the laboratory. Post-parasitic juveniles emerged from worker ants, and were maintained in water until the mermithids molted to the adult stage 3 wk later. Identification of the mermithid nematodes was performed by George Poinar, Jr. (Oregon State University, USA).

Voucher specimens of ants and parasitoids were deposited in the Arthropod collection of ECOSUR at Chetumal, Quintana Roo, Mexico. Mermithids were deposited in the collection of the USDA Nematode Laboratory, Beltsville, Maryland.
RESULTS

Nest Architecture

*Ectatomma tuberculatum* nests at Apazapan were characterized by a loose architecture. The entrance of the nest exhibiting the characteristic external chimney gallery, made of rough plant fibers and soil particles, was located at the base of a tree and could run vertically up to a height of 60-90 cm (2 to 3 ft) (Wheeler 1924). However at Apazapan there was no central vertical gallery with several horizontal chambers spreading from it, as is typical for nests in Chiapas and Brazil (Cook 1904a; Valenzuela et al. 1995; Hora et al. 2005b). Instead, the main gallery just under the entrance, followed the roots of the supporting tree, skirted round large and medium sized stones, and was extended backwards almost horizontally following the slope of site. By contrast with the 3 to 6—and sometimes up to 10—chambers (empty spaces whose soil and ceiling were compacted with debris and cocoon remains) found in nests in Chiapas (Lachaud, unpublished data), no chambers were observed in the nests in Apazapan. Workers and brood were found scattered in hollow cavities formed behind stones or between roots and stones, and loosely dispersed in the soil. Finally, while *E. tuberculatum* nests collected in Chiapas reached depths of up to 2 m (Valenzuela et al. 1995), nests at Apazapan hardly reached 0.60 m.

Colony Composition

From the 9 nests collected, 6 had only workers and brood, and only 3 contained dealate females: one with 1 dealate female, 1 with 2 dealate females and 1 with at least 7 dealate females and several alate females at the moment of collection. However, the following day, when the latter nest was cleaned and individuals were counted, most of the winged alate females had lost their wings leading to a total of 18 dealate females and 3 alate females (Table 1). No small size females, belonging to the social parasite species *Ectatomma parasicicum*, which resembled *E. tuberculatum* females in all morphological characteristics but size, were found. Worker numbers ranged from 7 to 150 at the moment of collection (72.3 ± 13.5, mean ± SEM). Five out of the 9 nests collected had cocoons and only 3 had a few larvae.

Eucharitid Parasitism

Only 2 of the *E. tuberculatum* nests were parasitized by eucharitid wasps, which accounted for a global nest parasitism of 22.2%. From a total of 57 cocoons collected from 5 nests, 15 were parasitized, i.e., the global prevalence of parasitism of cocoons was 26.3% (Table 1). Three larvae, 6 pupae and 4 pupae emerged from parasitized cocoons.

| Collecting date | Dealate females | Alate females | Parasitized pupae (%) | Parasitized larvae (%) | Eucharitid adults | Mermithized workers | Eggs presence | Pupae | Larvae | Workers | Mermithized cocoons | Eucharitid cocoons |
|----------------|----------------|---------------|-----------------------|------------------------|------------------|----------------------|---------------|-------|--------|---------|---------------------|--------------------|
| 21/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 21/09/2010     | 18             | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 21/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 21/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 23/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 23/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 23/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 23/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 23/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
| 23/09/2010     | 0              | 0             | 0                     | 0                      | 0                | 0                    | ++            | 0     | 0      | 0       | 0                   | 0                  |
and 7 adult eucharitids were retrieved: 6 adults belonged to Isomerala coronata (Westwood) (2 females, 4 males) and one was a Dilocantha lachaudii Heraty male. One case of superparasitism was identified. In fact, most of the cocoons (42/57) belonged to the same nest from which 14 were parasitized by eucharitids accounting for a prevalence of parasitism of cocoons of 33.3% for that nest. Furthermore, from a total of 9 larvae collected from 3 nests only 1 larva was parasitized (i.e., a global prevalence of parasitism of larvae of 11.1%), and bore a planidium attached to the left maxilla.

Mermithid Parasitism

Six workers, all belonging to the largest polygynous nest collected, had enlarged gasters and all were found to be mermithized. One of the workers died before mermithid emergence but 5 post-parasitic juvenile mermithids emerged from the other workers, causing their death. Of these, 2 adults were retrieved and were killed in hot water at 50°C and then placed in fixative (5% formalin). Both specimens were juvenile females identified as belonging to the genus Mexinermis Poinar, Lachaud, Castillo & Infante and, most probably, to the same species, M. ectatommi Poinar, Lachaud, Castillo & Infante as that parasitizing E. ruidum Roger in Chiapas (see Poinar et al. 2006). However, in the absence of males and mature females, species identification could not be certified.

DISCUSSION

Apart from some scarce myrmecophiles found very occasionally within their nest, such as a gastropod, a small diplopod (Merocheta), a lepismatid or nicoletiid thysanuran, a worm infesting the refuse dumps, and a histerid beetle (Cook 1905; Pérez-Lachaud & Lachaud, unpublished data), colonies of E. tuberculatum are mainly parasitized by eucharitid parasitoid wasps and mermithid nematodes. Parasitism of E. tuberculatum by the eucharitid wasp I. coronata was known only from an earlier report on a Guatemalan population (Cook 1905). More recent studies in Chiapas, Mexico (Lachaud et al. 1998; Pérez-Lachaud et al. 2007a, 2010) confirmed parasitism by I. coronata and additionally reported 2 other eucharitid parasitoids attacking this ant host, belonging to 2 different genera, D. lachaudii and an undescribed, uncommon species of Kapala. Regarding nematode parasitism, originally reported for E. tuberculatum populations from Panama and Trinidad and, perhaps, from Costa Rica (Emery 1890; Wheeler 1930), it has also been observed in at least 16 out of 253 nests collected from the Soconusco population (Lachaud, unpublished data). On the other hand, the facultatively polygynous population of Apazapan is the only one where the social ant parasite E. parasiticum is known to occur (Hora et al. 2001, 2005a; Feitosa et al. 2008), but neither eucharitid nor mermithid parasitism had ever been reported for it before.

Although nest sample size at Apazapan was relatively small, it is worth noting that the same multi-species system of parasitoids and parasites attacked E. tuberculatum colonies both in Veracruz and in Chiapas, whatever the social structure (monogyny or facultative polygyny) of their populations. Both populations hosted a mermithid parasite, presumably the same species in both cases; though the specimens from Chiapas have not yet been identified. Moreover, the polygynous population from Veracruz, hosted the same predominant eucharitid parasitoid species, D. lachaudii and I. coronata, as the monogynous population from Chiapas (Pérez-Lachaud et al. 2010). In Apazapan most of the individual parasitoids belonged to I. coronata, while D. lachaudii prevailed as the main parasitoid of E. tuberculatum in the Rosario Izapa population. However, populations of both eucharitid species in Chiapas appeared to vary inversely in frequency, with predominance of one or the other species during the year (Pérez-Lachaud et al. 2010), and such patterns could also occur at Apazapan; but a larger sample, distributed along the year, is needed to verify this trend.

Nest collection at Apazapan was performed over too short a period to allow extrapolation of the real annual prevalence of parasitism at the scale of the population. However, the value of 26.3% obtained for the global prevalence of eucharitid parasitism for cocoons from the 9 nests collected at the end of Sep was fully compatible with that recorded in Chiapas during the same period of the year and with a similar sample size for the monthly collected nests: 27.1% for Oct 2004 (n = 8) and 14.4% for Sep 2005 (n = 10) (see Table 1 in Pérez-Lachaud et al. 2010). These results suggest that the prevalence of parasitism was rather similar between both populations and, therefore, that the hypothesis of increased resistance to parasites and parasitoids as one of the factors responsible for polygyny emergence in E. tuberculatum populations is not supported. This conclusion seems to be given further weight by 2 apparently adaptive biological traits related to parasitism, which make unnecessary the emergence of polygyny, i.e., i) the apparent potential of E. tuberculatum, at least for the Rosario Izapa population (Pérez-Lachaud et al. 2010), to modulate the impact of eucharitids on the colonies through differences in the seasonal timing of eucharitid main attack and sexual brood production by ants, and ii) the polydomy that seems to characterize this species (García-Pérez et al. 1991; Hora et al. 2005b) and that would reduce the probability of
any given colony having its whole brood parasi-
ized (Pérez-Lachaud et al. 2010).

Variation in morphological, behavioral, physi-
ological and life history traits is thought to enable
an individual (or a colony in the case of social in-
sects) to maximize its reproductive success under
a wide range of conditions, and thus cope better
with variable ecological constraints (Foitzik et al.
2010). The causes of the facultative polygyny
found at Apazapan or in Brazil thus seem to have
originated from environmental stress or con-
straints other than parasitism. Nest site limita-
tion, risks of dispersal, and/or high rates of preda-
tion that could be linked (see Bourke 1999, Kas-
pari & O’Donnell 2003; Linksvayer & Janssen
2009) to the lower colony sizes recorded for poly-
gynous populations (Brazil: Hora et al. 2005b;
Apazapan: Hora et al. 2001 and these results)
could be promising candidate explanations. An
important difference between Chiapas and Ver-
acruz sites is the type and structure of the soil,
which seemingly determined the structure of the
nests at both sites. At Apazapan, our observations
indicate that the number of trees available as
nest supports is more than enough, but digging
difficulties due to the very rocky soil of the site
could be a much more limiting factor for
foundresses. In such conditions, the founding of a
new colony by dispersed females would be ex-
 tremely risky and, as suggested by Keller
(1995b), could favor the selection of young mated
queens seeking adoption into established colonies
(secondary polygyny), or readoption in their
mother colony, as apparently occurs in the Brazil-
ian population (Zinck et al. 2007). Further studies
detailing the soil characteristics in the different
sites and their consequences on both nest struc-
ture and colony foundation possibilities, along
with within colony genetic diversity studies
would help reach a better understanding of the
environmental pressures controlling the emerg-
ence of polygyny in *E. tuberculatum* and the so-
cioecology of this species.

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