Studies on the Asian sawflies of *Formosempria* Takeuchi (Hymenoptera, Tenthredinidae), with notes on the suitability of *F. varipes* Takeuchi as a biological control agent for skunk vine, *Paederia foetida* L. (Rubiaceae) in Florida

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Abstract

*Formosempria* Takeuchi, 1929, is distributed in southeastern Asia from Taiwan and China to Vietnam, Myanmar, and possibly northern India. Three species are included: *F. crassicornis* Wei & Nie, 2002, *F. shanensis* Malaise, 1961, and *F. varipes* Takeuchi, 1929 (= *F. annamensis* Malaise, 1961, syn. n.; = *F. metallica* Wei, 2003, syn. n.). *Formosempria varipes* was reared from larvae feeding on *Paederia foetida* L. (Rubiaceae) in Hong Kong and was a potential biological agent for the invasive *P. foetida* in Florida. Larval feeding tests indicate more than one species of *Paederia* are suitable hosts for *F. varipes* and further study for use as a biological control agent in Florida is unwarranted. Descriptions and illustration of the species are given, and life history notes on *F. varipes* are presented.

Keywords

Symphyta, biological control, skunk vine
Introduction

Interest in the small and little-known genus *Formosempria* Takeuchi was prompted by discovery of larvae of a species in Hong Kong feeding on skunk vine, *Paederia foetida* L. (Rubiaceae), a target for a biological control program in Florida (Pemberton and Pratt 2002). Adults reared from these larvae were identified as a species of *Formosempria*, an Asian genus in the subfamily Allantinae. Taeger et al. (2010) listed five described species in this genus, distributed from Taiwan and western and southern mainland China to Vietnam, Myanmar, and possibly northeastern India. Very few specimens of *Formosempria* were previously available for study, most of which are types of the described species, but additional specimens from Taiwan and those reared from Hong Kong have become available. The identity of the Hong Kong species, as outlined here, is *Formosempria varipes* Takeuchi, 1929, which was described from Taiwan.

*Formosempria* was based on a single species, *F. varipes*, described from a single specimen (Takeuchi 1929). Malaise (1961) added two species, *F. annamensis* from Vietnam and *F. shanensis* from Myanmar, and gave a key to the three species. Wei and Nie (2002) and Wei (in Wei and Nie 2003) added *F. crassicornis* and *F. metallica* from China. Saini and Deep (1994) and Saini (2006) reported *F. shanensis* Malaise, the species described from Myanmar, from India, but this is questionable. The genus probably does not occur in India. Haris (2012) recorded *F. crassicornis* from Vietnam. Critical to the study of *Formosempria* is the identity of the type species of the genus, *F. varipes* Takeuchi. Unfortunately, the type could not be located. However, among some specimens from Taiwan we found a series identical to Takeuchi’s description of *F. varipes*. We use these specimens to characterize *Formosempria* and the included species.

Discovery of *F. varipes* feeding on *Paederia* in Hong Kong initiated studies on the species as a possible biological control agent for *P. foetida* in Florida. However, host preference studies showed that several species of *Paederia* may serve as host plants, thus making it unsuitable for release as a biological control agent of *P. foetida* in Florida.

Material and methods

Figures 1-15 were obtained using an EntoVision Imaging Suite that included a firewire JVC KY-75 3CCD digital camera mounted to a Leica M16 zoom lens via a Leica z-step microscope stand. Multiple focal planes were merged using Cartograph 5.6.0 (Microvision Instruments, France) software. Figures 16–18 were prepared by PDP.

Acronyms used are: CSCS (Collection of Central South University of Forestry and Technology, Changsha, Hunan, China); FSCA (Florida State Collection of Arthropods, Gainesville, FL, USA); IPRL (Invasive Plant Research Laboratory, Agricultural Research Service, U. S. Department of Agriculture. Ft. Lauderdale, FL, USA); NHR (Naturhistoriska Riksmuseet, Stockholm, Sweden); UOP (University of Osaka Prefecture, Sakai, Japan); SDEI (Senckenberg Deutsches Entomologisches Institut,
Müncheberg, Germany); USNM (National Museum of Natural History, Smithsonian Institution, Washington, D.C., USA).

Recent surveys by JM resulted in the discovery of numerous *F. varipes* larvae feeding on the foliage of *P. foetida* in the Sham Tseng area (N 22.373, E114.064), Hong Kong. Taxonomic and biological assessments were based on *F. varipes* individuals that were shipped from Hong Kong in March 2013 and April 2014 to the quarantine facility at IPRL, under permit #P526P-12-04814. In an effort to establish a laboratory colony of *F. varipes*, 25 males and 25 females from the 2013 shipment were added to each of three screen cages (33 × 33 × 61 cm) within the quarantine glasshouse (28 °C (±5), 85% (±10) relative humidity). Each cage contained a potted *P. foetida* plant as well as several cotton balls hydrated with dionized water, which were positioned on the top surface of each cage. Observations of these adults, the resulting eggs, and larvae were recorded.

A subset of adults arising from the 2014 shipment were incorporated into no-choice host range tests, which were designed to quantify the propensity of *F. varipes* to oviposit and feed on select *Paederia* species. Among the *Paederia*, only *Paederia ciliata* (Bartl. ex DC) Standley is native to North America and has a geographic range limited to Mexico (Puff 1991). The release of *F. varipes* as a biological control agent of *P. foetida* in Florida may pose significant risk to the Mexican endemic if female sawflies recognize *P. ciliata* as an ovipositional host and larvae complete development when feeding exclusively on the plant. To test these host use patterns, five replicate *P. ciliata* or *P. foetida* potted plants were placed in individual screen cages (33 × 33 × 61 cm) within a quarantine glasshouse as described above. Plants were pruned to ensure a uniform height and biomass among individuals. Five male-female pairs, each less than 24 hours old, were randomly assigned to each cage and held with the test plant until death. Each plant was reviewed for oviposition incidence 7 days following adult inoculation and larval densities per plant were noted 21 days after adult release.

Additionally, no-choice larval tests were conducted to determine if *P. ciliata* is a developmental host. Other *Paederia* species were also included to provide insights into the generic specificity of the herbivore: *Paederia bojeriana* (Rich.), *P. crud-dasiana* Prain, *P. pilifera* Hook., and the target weed *P. foetida*. Tests were conducted in a single controlled environmental chamber set at 25 °C, 75% relative humidity and 14:10 light:dark. Neonate larvae (< 24 hrs old) were collected from the laboratory colony and five individuals were transferred onto a slightly moistened filter paper within individual 90 × 15 mm petri dishes. Five replicate petri dishes with the accompanying larvae were randomly assigned one of five replicated plants for each of the five host plant treatments. Leaves were excised from assigned test plant replicates and placed within the petri dish, which was sealed with parafilm and maintained at internal conditions of 25 °C and 90% relative humidity. A mixture of soft, newly developing and mature, fully expanded leaves in excess of the herbibore’s daily consumption were added to each petri dish every 24 hours. Survivorship was monitored daily and larval development time to the prepupal stage was noted. The influence of host plant species
on oviposition, larval survival, and development rates were compared with ANOVA followed by Tukey's HSD post hoc analysis (PROC GLM; SAS 1999). Data are presented as means (± standard error).

Results
Systematics

*Formosempria* Takeuchi

*Formosempria* Takeuchi 1929: 85. Type species: *Formosempria varipes* Takeuchi, original designation; Malaise 1961: 247 (diagnosis; key to 3 species); Malaise 1963: 159 (in world key); Togashi 1990: 184 (one species, Taiwan); Abe and Smith 1991: 34 (genus listed); Chou and Naito 1991: 89 (one species, Taiwan); Saini and Deep 1994: 48 (one species, India); Wei and Nie 1998: 28 (placed in Blennocampidae, Belesininae, Atelozini); Wei 2001: 678 (one species, Zhejiang, China); Lacourt 2003: 506 (in Allantinae, Atelozini); Wei et al. 2006: 527 (three species listed from China); Saini et al. 2006: 579 (one species, India); Xiao 2006: 192 (one species listed from China); Taeger et al. 2010: 282 (world catalog, five species listed); Haris 2012: 138 (one species, Vietnam).

**Description.** Antenna (Figs 3, 10) hairy, stout and slightly thickened medially; scape longer than broad; pedicel as long as broad, antennomere 3 longer than 4; apical 3 antennomeres each with ventral sensory area. Malar space linear; clypeus (Figs 4, 8) truncate or bluntly protruding anteriorly; each mandible with large subapical tooth; inner margins of eyes slightly converging below (Figs 4, 8); genal carina absent; head from above strongly narrowing behind eyes (Figs 5, 9). Epicnemium absent. Forewing (Fig. 2) with 4 cubital cells; hind wing (Fig. 2) without cells Rs and M, anal cell with short petiole. Inner fore tibial spur appearing simple but with very slight preapical projection. Tarsal claws (Fig. 11) with long inner tooth, nearly as long as outer tooth and lateral to outer tooth, with basal lobe. Pulvilli small on hind tarsomeres 3 and 4. Head and body usually with slight metallic luster.

**Remarks.** Species of *Formosempria* are superficially similar to species of *Empria*, being similar in size and color and similar wing venation; thus, the reason for the name. Takeuchi (1929) mentioned that they are like *Empria*, but without a malar space. *Formosempria* will key to the same couplet as *Hemibeleses* in Malaise's 1963 key, but are separated from *Hemibeleses* which has the pedicel as long or longer than the scape and the 3-toothed hind claws in the male. Males of *Formosempria* were not known at that time, but the hind claw is similar to the fore- and midclaws, not 3-toothed as in *Hemibeleses*.

Based on this study, two species of *Formosempria* are recognized, *F. varipes* from Taiwan, southeastern China, and Vietnam, and *F. shanensis* from Myanmar. *Formosem-
Formosempria crassicornis Wei and Nie is listed, but the type was not examined and its status cannot be determined. The record of Formosempria from India by Saini and Deep (1994) and Saini (2006) is questionable.

**Formosempria crassicornis** Wei & Nie

*Formosempria crassicornis* Wei and Nie 2002: 837, 839, 846, 848, figs 10, 11 (female sheath, saw on p. 848); Wei et al. 2006: 527 (listed); Haris 2012: 138 (recorded from Vietnam).

**Comments.** This species was described from Hainan, China. The holotype, deposited in the Institute of Zoology, Chinese Academy of Sciences, Beijing, was not examined.

The figures by Wei and Nie (2002) of the female sheath and lancet do not differ from *F. varipes*, and *F. crassicornis* may be a synonym. Even though distinguishing characters for the species are not evident in the description and the species seems to fall within the range of *F. varipes*, it is kept valid until the type can be examined. Haris (2012) recorded a female from Vietnam (“N. Vietnam, Ninh Binh, Cue Phuong NP, near centre, c. 225 m, 20.xii.1999-19.ii.2000”), but gave no discussion. This specimen was not examined.

**Formosempria shanensis** Malaise

Figs 1–5

*Formosempria shanensis* Malaise 1961: 249, fig. 8A,B,C (head front; antenna, sheath dorsal); Malaise 1963: figs 73, 131; Saini and Deep 1994: 48 (listed; first record from India); Saini 2006: 68, 152, figs 215–222 (India record; male described); Saini et al. 2006: 579 (listed).

**Comments.** Malaise described this species from five females from “Burmese Southern Shan States (Taunggyi at 1500 m)”. The syntypes are in NHR and were examined. The lectotype, here designated, is a female labeled “S. Shan States, Burma, 1500 m, Taunggyi, 1-VIII-22.IX.34 [“22.IX” crossed out], Malaise”, “TYPUS” [red], “Formosempria shanensis n. sp., R. Malaise det 1958”, “NHRS-HEVA 000000834” with a lectotype label added. The other four females are paralectotypes with the same data but with the label “PARATYPUS” [in red]; three do not have dates crossed out and one has “1-VIII” crossed out.

The color and structure are very similar to *F. varipes*, but *F. shanensis* differs by its slightly larger size, the more truncate clypeus with more acute lateral corners (Fig. 4, more rounded laterally in *F. varipes*, Fig. 8), and longer antennae (Fig. 1), being more than half the length of the body (thorax and abdomen) and about 2.2× the head width. These differences seem significant to retain *F. shanensis* as a distinct species at present.
Saini and Deep (1994) and Saini (2006) recorded this species from five males from India: “Himachal Pradesh, Dalhousie, Kalatop, 2850 m, 29.6-30.6.1986”. Saini (2006) mentioned these are the first known males of the genus, although Wei (in Wei and Nie 2003) described the male of *F. metallica*. Saini (2006) also included description of the female. These specimens have three-toothed hind claws and the genitalia figured are quite different from that of associated males of *F. varipes*. We have not seen
specimens, but according to Saini’s (2006) description, we question that these are *F. shanensis* or even a member of *Formosempria*. Thus, the India record for *Formosempria* is doubtful.

**Formosempria varipes** Takeuchi  
Figs 6–18

*Formosempria varipes* Takeuchi, 1929: 85–86. Figure of wings; Malaise 1961: 249 (in key); Togashi 1990: 184, figs 46–50; Wei et al. 2006: 527 (listed).  
*Formosempria annamensis* Malaise, 1961: 249. **syn. n.**  
*Formosempria metallica* Wei, in Wei and Nie 2003: 136–137, 207, figures, male and female; Wei et al. 2006: 527 (listed). **syn. n.**

**Description.** Female: Length, 6.5–7 mm. Black, with slight metallic lustre, and following white: apices of coxae, trochanters, basal third of mid and hind femora, basal third to half of mid and hind tibiae, base of hind basitarsus, most of mid and fore basitarsi. Wings lightly, uniformly infuscated; veins and stigma black. Head and body with slight metallic luster, smooth, shiny, with short silvery hairs.  
Antenna (Figs 6, 10) with scape longer than broad; pedicel about quadrate; antennomere 3 longer than 4, remaining antennomeres gradually decreasing; length less
than half body length (thorax and abdomen) and about 1.7× head width. Clypeus truncate to slightly protruding anteriorly, with corners rounded (Fig. 8). Malar space linear. Hind basitarsus (Fig. 11) about equal to length of remaining tarsomeres combined. Sheath (Fig. 13) slender in dorsal view, in lateral view slightly truncate at apex, appearing slightly directed dorsally. Lancet (Fig. 12) with serrulae low, pointed at apices, each with about 6 fine anterior and posterior subbasal teeth.

Male: Length, 6.5 mm. Similar in color and structure to female. Genitalia in Figs 14, 15.

Neonates (Fig. 17): Similar to late feeding stage.

Last feeding stage (Fig. 18): Length, about 12 mm. Head black with postclypeus, preclypeus, area between antacorium and clypeus and mandible, mandible except apex and mouthparts except apices of palpi whitish; labrum brownish. Thorax and abdomen entirely pale with whitish bloom. Head with long scattered setae, longer than preclypeus; preclypeus with 2 setae on each side; labrum with 3 setae on each side; right mandible with 3 small ventral teeth and 2 acute and one broad dorsal teeth; left mandible with 2 acute ventral teeth and 4 acute dorsal teeth and with inner ridge extending inward from outer dorsal tooth. Thorax with scattered long setae. Abdominal segments each with 6 dorsal annules; 2–3 setae on each side of annules 2 and 4; subspiracular lobe and surpedal lobe each with about 4 long setae; apical segment with about 12 long setae dorsally and with long setae on subanal area.

Prepupa: Differs from feeding stages by head, thorax, and abdomen entirely pale, whitish. Setae on head and body absent. Each mandible with 3 linear teeth; left mandible with outer tooth largest, basal 2 teeth smaller and subequal in size; right mandible with central tooth largest, outer and inner teeth smaller and subequal in size.

**Type material.** Takeuchi described this species from a single female “Sozan near Taihoku, Formosa” “collected by M. Kato, on May 2, 1926”. The holotype should be in UOP with the Takeuchi collection. On a visit to Japan in 1979, I examined the holotype and took some brief notes. When recently requesting the type, it could not be found in the Takeuchi collection at UOP. A thorough search for the type was made by Noria Hirai (UOP) and Akihko Shinohara (National Museum of Nature and Science), but to no avail.

*Formosempria annamensis* was described from a single female. The holotype is at NHR, was examined and is labeled “Phuc Son, Annam, XI-XII, H Rosse, Berlin, s.w. 11”, “TYPUS” [red], *Formosempria annamensis* n. sp., R. Malaise det 1958, “NHRS-HEVA 000000839”.

*Formosempria metallica* was described from nine female and male specimens from Hubei, Fujian, Zhejiang, Hangzhou provinces, China. The female holotype is at CSCS, from “Maheba, Xiangeng, Hubei, 1999-VII-25” and was examined. The holotype label is in Chinese, with the same date and “450 m.” and additional labels “F. metallica”, “Holotype: Formosempria metallica Wei female, M. Wei 1992”.

**Specimens examined.** CHINA: Hong Kong, Sham Tseng San Tsuen Temple, larvae, March 24, 2013, on *Paederia foetida*, adults April 2013, reared in quarantine, ARS, Ft. Lauderdale, FL by Paul Pratt (5 ♀, 5 ♂, USNM); same data, larvae March
2014, pupated and adults emerged IV–V 2014 (about 7 ♀, 100 ♂, 20 larvae, USNM, IPRL); Hubei, holotype and paratype of *F. metallica* (1 ♀, 1 ♂, CSCS); Henan (1 ♀, SDEI). TAIWAN: Keelung Co., Keelung City, V.2004, Malaise trap, L. Stange and H. Wang (1 ♀, 4 ♂, FSCA). VIETNAM: Holotype of *F. annamensis* (1 ♀, NHR).

**Distribution.** China (Hong Kong, Hubei, Fujian, Zhejiang, Hangzhou); Taiwan; Vietnam.

**Hosts.** *Paederia foetida*, also *P. ciliata, P. cruddasiana, P. bojeriana*, and *P. pilifera* based on host testing (Rubiaceae).

**Comments.** The interpretations of *F. varipes* by Malaise (1961), Togashi (1990) and others were based on the original description by Takeuchi (1929), not examination of the type. The type is now presumed lost. In a collection of sawflies from Taiwan (FSCA) DRS discovered a series of four males and one female that agree with Takeuchi’s description. Due to the absence of the type, we are using these specimens as representative of *F. varipes*. A key character used by Malaise (1961) and others to separate *F. varipes* is the length of the hind basitarsomere. Takeuchi (1929) stated “posterior basitarsus shorter than the following joints”. In the specimens examined, the hind basitarsomere is subequal to very slightly longer than the remaining tarsomeres combined. This is the only discrepancy between the Taiwan specimens we have and Takeuchi’s (1929) description. However, because of the overwhelming similarity of the Taiwan specimens with Takeuchi’s description, we do not believe “shorter”, “subequal”, or “slightly longer”, alone are sufficient to justify a separate species. This could have been misinterpreted by the authors, or it could be slightly variable in the species. The female ovipositor and male genitalia of the Taiwan and Hong Kong specimens were compared and were identical.

Togashi (1990) briefly described one female from Taiwan “Shin Bao Shi, near Liu Kui, 3.V.1986, K. Baba leg.” He stated that it agrees with Takeuchi’s original description. He illustrated the dorsal view of the head (fig. 46), front inner tibial spur (fig. 47), tarsal claw (fig. 48), sheath (fig. 49), and lancet (fig. 50). The front inner tibial spur was illustrated and stated to be simple; however, in specimens we have examined there is a very slight subapical projection. Otherwise, Togashi’s description and illustration agree with the specimens examined from Taiwan. Togashi’s specimen was not examined.

Malaise’s holotype of *F. annamensis* from Vietnam is identical to specimens from Taiwan and Hong Kong; therefore, we synonymize this species.

Wei (in Wei and Nie 2003) stated that *F. metallica* “…differs from its congeners in malar space absent, lateral furrows of postocellar area very weak and shallow, antenna as long as half the body length, and mesoscutellum black”. Figures in Wei and Nie (2003) of the male genitalia, tarsal claw, sheath and part of the lancet do not noticeably differ from specimens from Taiwan and Hong Kong. All of the differing characters mentioned by Wei are shared with *F. varipes*, and, upon examination of the holotype, DRS observed no differences between this and *F. varipes*. Thus, *F. metallica* is synonymized.
Biology and host specificity of *Formosempria varipes*  

The objective of importing *F. varipes* into the IPRL quarantine facility was to determine the sawfly’s host range, from which inferences can be drawn concerning the herbivore’s suitability as a biological control agent of *P. foetiida* in Florida. Females from the 2013 survey were observed dragging their ovipositors across the leaf surface shortly after release into the cages, creating 1-2 cm longitudinal incisions through the leaf and typically located midway between the midvein and the leaf margin. Eggs (Fig. 16) were inserted between the adaxial and abaxial leaf surfaces, usually on the leaf margin but occasionally along the leaf midribs. Oviposition was consistently adjacent to the incision created by the female. The reason for this leaf cutting behavior is unknown but may be related to assessment of host plant suitability or the disruption of host plant defenses. In an effort to force mating, the heads of several males were excised and multiple attempts were made to mate females but all efforts were unsuccessful.

 Neonates (Fig. 17) exited their leaf-enclosed eggs approximately one week following oviposition. Groups of first instar larvae were observed feeding gregariously, consuming the foliage between the leaf veins. First instar larvae were transferred to small, ventilated plastic cages and fed exclusively on *P. foetiida* until pupation, demonstrating that the target weed is a developmental host. Later instars consumed entire leaves. Larvae (Fig. 18) possess a black head capsule but only later instars also have a fine wax covering the thorax and abdominal integument. The last instar larvae (prepupae) were observed wandering within the cages and lacked the wax covering, possessing instead a pale colored head capsule and integument. Prepupae were transferred to plastic boxes filled with a loose sandy soil, where the larvae readily burrowed below the surface. Pupation occurred within an oblong casing (cell) constructed, in part, from the surrounding soil. Adults emerged from their pupal cells approximately two weeks following pupation and tunneled to the surface of the soil. The subsequent (F₁) generation, however, were exclusively males and the laboratory colony was lost. These observations confirm that females of *F. varipes* are arrhenotokous.

 A second collection of *F. varipes* larvae in Hong Kong was made in April 2014 and efforts to colonize the species were repeated. Adults arising from this 2014 shipment were also used in no-choice host range tests, which were designed to quantify the propensity of *F. varipes* to oviposit and feed on select *Paederia* species. Oviposition was observed among all replicates of *P. ciliata* and *P. foetiida* tested. There was no difference in the number of larvae recovered from the *Paederia* species used in the ovipositional test ($F_{1,9} = 3.0; P \leq 0.1214$). From these data it is clear that *F. varipes* females readily oviposit on the Mexican endemic *P. ciliata* so no-choice larval feeding tests were conducted to determine if *P. ciliata* is also a developmental host. Survivorship varied among hosts ($F_{4,24} = 10.38; P < 0.001$), with no individuals completing development when held with *P. pilifera*. Larval survivorship did not vary when feeding on the remaining species including *P. foetiida* (90.0% (±0.05)), *P. cruddasiana* (88.0% (±0.08)), *P. ciliata* (76.0% (±0.10)), or *P. bojeriana* (52% (±0.22)). Host plant species also
influenced development times ($F_{3,17} = 6.69; P < 0.005$). Development from neonate to prepupa was shortest when feeding on *P. foetida* (9.5 days ($\pm 0.3$)) and *P. bojeriana* (9.8 ($\pm 0.5$)) but slowest when held with *P. ciliata* (13.1 ($\pm 0.9$)), with *P. cruddasiana* (11.6 ($\pm 0.6$)) intermediate. Like those of 2013, all efforts to establish a laboratory colony were unsuccessful with the 2014 material.

Although *P. ciliata* may not be an optimal host based on development rates, these data demonstrate that *F. varipes* will readily oviposit and complete development on the Mexican native under no-choice conditions. The risk *F. varipes* would pose to *P. ciliata* populations is a function of 1) host use, as reported herein, but also 2) the permeability of geographic barriers that inhibit landscape level dispersal of the herbivore from its intended range (Florida) to the native range of *P. ciliata* (central Mexico). While the dispersal pathway(s) remain unknown, there are several examples of biological control agents spreading far beyond their intended geographic ranges through natural or anthropogenic means (Pratt and Center 2012). Considering the ever increasing levels of tourism and trade between Florida and Mexico, the authors consider the probability of *F. varipes* dispersing to Mexico and threatening *P. ciliata* populations too great a risk to justify its release. Therefore, additional resources dedicated to the development of this species as a biological control agent are unwarranted.

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**References**


