

Effect of Honey Solution and Water Acquisition on Survival of Starved Solenopsis Mealybug, Phenacoccus solenopsis

Authors: Huang, Fang, Wang, Feifei, Lu, Yaobin, Zhang, Pengjun, Zhang, Jinming, et al.

Source: Journal of Insect Science, 14(1): 1-9

Published By: Entomological Society of America

URL: https://doi.org/10.1673/031.014.01

BioOne Complete (complete.BioOne.org) is a full-text database of 200 subscribed and open-access titles in the biological, ecological, and environmental sciences published by nonprofit societies, associations, museums, institutions, and presses.

Your use of this PDF, the BioOne Complete website, and all posted and associated content indicates your acceptance of BioOne's Terms of Use, available at <u>www.bioone.org/terms-of-use</u>.

Usage of BioOne Complete content is strictly limited to personal, educational, and non - commercial use. Commercial inquiries or rights and permissions requests should be directed to the individual publisher as copyright holder.

BioOne sees sustainable scholarly publishing as an inherently collaborative enterprise connecting authors, nonprofit publishers, academic institutions, research libraries, and research funders in the common goal of maximizing access to critical research.



Effect of honey solution and water acquisition on survival of starved solenopsis mealybug, *Phenacoccus solenopsis*

Fang Huang^a*, Feifei Wang^b, Yaobin Lu^c*, Pengjun Zhang^d, Jinming Zhang^e, Zhijun Zhang^f, Weidi Li^g, Wencai Lin^h, Yawei Beiⁱ

State Key Laboratory Breeding Base for Zhejiang Sustainable Pest and Disease Control, Institute of Plant Protection and Microbiology, Zhejiang Academy of Agricultural Sciences, Hangzhou, Zhejiang 310021, China

Abstract

The current study examined the effects of honey solution and water access on feeding behavior and survival of starving solenopsis mealybugs, *Phenacoccus solenopsis* Tinsley (Hemiptera: Pseudococcidae). The electrical penetration graph technique and an artificial membrane system were used to check whether *P. solenopsis* could imbibe free water or other liquid, such as the honey solution used here, in its natural environment. The recorded electrical penetration graph waveforms revealed that *P. solenopsis* could continuously imbibe water-honey solution for several hours, which indicated that honey solution and water acquisition could possibly occur when *P. solenopsis* had access to such liquids in its natural environment. Waveforms of water-honey solution feeding alternated between two distinct feeding phases in a regular pattern, which was assumed to reflect inherent habits of feeding attempts. The effects of honey solution and water acquisition on survival of *P. solenopsis* was also examined. Comparison between *P. solenopsis* in different treatments (starved, water feeding, honey solution feeding, and cotton plant feeding) suggested that 1) *P. solenopsis* could accept but did not favor feeding on water or the honey solution, and 2) this feeding could prolong its survival, but had no effect on body size.

Keywords: artificial membrane, cotton mealybug, electrical penetration graph technique, invasive pest, starvation **Abbreviations: CRL**, control treatment; **EPG**, electrical penetration graph; **NM**, access to honey solution treatment; **SV**, starvation treatment; **WM**, access to water treatment

Correspondence: ^a huangfang_ch@hotmail.com, ^b 347655996@qq.com, ^c luybcn@163.com, ^d peng_junzhang@hotmail.com,

e insectzhang@163.com, f 77163880@qq.com, & liwd2003@163.com, h wencailwc@sina.com, l shibeima@163.com, *Corresponding authors Editor: Todd Shelly was editor of this paper.

Received: 10 May 2012 Accepted: 21 November 2013 Published: 2 January 2014

Copyright: This is an open access paper. We use the Creative Commons Attribution 3.0 license that permits unrestricted use, provided that the paper is properly attributed.

ISSN: 1536-2442 | Vol. 14, Number 1

Cite this paper as:

Huang F, Wang F, Lu Y, Zhang P, Zhang J, Zhang Z, Li W, Lin W, Bei Y. 2014. Effect of honey solution and water acquisition on survival of starved solenopsis mealybug, *Phenacoccus solenopsis*. *Journal of Insect Science* 14:1. Available online: http://www.insectscience.org/14.1

Introduction

The solenopsis mealybug, Phenacoccus so-Tinsley lenopsis (Hemiptera: Pseudococcidae), is a sap-sucking pest of many agricultural, horticultural, and ornamental crops, including cotton (Dutt 2007; Wang et al. 2010). Recently, it has caused serious economic losses in transgenic and conventional cotton production in India (Jhala et al. 2008) and Pakistan (Abbas et al. 2009). Its polyphagous nature, high reproductive capacity, and short life cycle (Huang et al. 2011), and wide morphological plasticity have enabled P. solenopsis to survive and even thrive under a wide variety of environmental conditions (Hodgson et al. 2008). The species has spread rapidly in Asia and beyond, threatening the world's cotton industry and other crops as an invasive pest (Wang et al. 2010). Consequently, P. solenopsis has been added to the European and Mediterranean Plant Protection Organization list (EPPO 2011) and Chinese quarantine pest list (MOA and AQSIQ 2009).

Population fitness homoeostasis, defined as the ability of an individual or population to maintain relatively constant fitness over a range of environments, determines species invasiveness in large part (Rejmánek 2000). Within parameters of this 'fitness,' resistance to food deprivation was suggested to be an important adaptation in flightless insects (Stockhoff 1991). Under severe defoliation, individuals may be forced to wander extensively to locate other hosts. Utilization of liquid (dew and/or proximal honey solution) in the environment was reported to be important for survival of some insects, i.e., Drosophila (Karan et al. 1998), mites (van Rijn and Tanigoshi 1999), mosquitoes (Gary and Foster 2004), and many coccinellids (Lundgren 2009). Little information could be found to reveal whether or how plant sapsucking insects with specialized mouthparts could use these fluids. Mittler and Dadd (1963) used the electronic penetration graph technique (EPG) and confirmed that the green peach aphid, Myzus persicae, could ingest considerable amounts of water and sugary fluids entirely by their own sucking efforts. Kehat and Wyndham (1972) reported that survival of the female Rutherglen bug, Nysius vinitor, with access to water was uninfluenced. If starving, could P. solenopsis consume water and/or other fluids when offered, and will this consumption increase its odds of survival? It has been suggested that P. solenopsis, as an invasive pest with limited mobility, was dispersed mainly through commercial transportation (Giliomee 2011; Fand 2013). Travel between suitable locations is potentially a lengthy process for *P. solenopsis*, and changes in food quality, falling from the host plant, and food depletion could occur during movement. If dew or another fluid could be used by P. solenopsis, individuals would be expected to survive longer, which would improve their chances to find a suitable environment. In other words, benefits from the utilization of these fluids could promote their survival in a new environment, which is an essential trait for an invasive insect.

The EPG technique was originally derived from the electronic feeding monitor, and it allows real-time tracking of the invisible stylet penetration (probing) activities by insects with piercing-sucking mouthparts (Tjallingii and Hogen Esch 1993; Walker 2000). An artificial membrane system that made the process of water sucking visible was developed (Huang et al. 2013) in conjunction with EPG. Plant sap sucking procedures of *P. solenopsis* were described, and water being sucked through a stretched parafilm membrane was observed in a previous study (Huang et al. 2012)..

To confirm whether water consumption is beneficial, the present study was conducted to investigate whether access to honey solution or water influenced the feeding behavior and survival of starving *P. solenopsis*. The importance of the consumption of these substances to the invasive ability of the species is discussed.

Materials and Methods

Insects and plants

Phenacoccus solenopsis specimens were originally collected from the ornamental plant Rose of Sharon, Hibiscus syriacus L. (Malvales, Malvaceae), in Hangzhou, China. The specimens were reared on cotton, Gossypium hirsutum L., cultivar Zhefengmian No.1, in a climate-controlled room at 27 ± 1° C, RH 65~75%, and 12:12 L:D. The G. hirsutum plants for use in the culture were grown in 13 cm plastic pots in a mixture of peat moss, Sphagnum sp. (Sphagnalus, Sphagnaceae), vermiculite, organic fertilizer, and perlite (10:10:10:1 ratio), and kept in a climatecontrolled room (25 \pm 3° C, R.H. 60~ 70%, 12:12 L:D). All the plants used in the experiments had two or three fully-expanded leaves.

EPG analysis

The stylet penetration activities of *P. solenop*sis on the artificial membrane were recorded by a Giga-4 DC-EPG system (EPG Systems, http://epgsystems.com). Newly-emerged *P.* solenopsis adults were carefully collected from the colony and starved for 24 hr. Their dorsal wax was partially removed with a longhaired goat's wool brush to permit attachment of an EPG insect electrode (a thin gold wire, diameter 18 μ m, length 2 cm) using a droplet of water-based silver glue. The insect electrode (+ve) was connected to the EPG ampliamplifier, which had an input resistance of 10⁹ ohm and a gain of 50× (Tjallingii 1985). In assessments of water or honey solution acquisition, the other electrode (-ve) was inserted into the liquid through the Parafilm (BEMIS, www.parafilm.com) membrane. Assessment of EPG on the G. hirsutum plant was set up as a control and was conducted using the same materials and methods as Huang et al. (2012). Test material was placed in a Faraday cage at an ambient temperature of $27 \pm 2^{\circ}$ C to shield it from external electric noise. All recorded signals were analyzed using Probe 3.4 soft-(provided W.F. ware bv Tjallingii, Wageningen, The Netherlands).

Before bioassay, *P. solenopsis* was held above the Parafilm using an attached gold wire and then lowered onto the membrane surface, at which point EPG output was recorded. Preliminary overview of stylet activity was observed continuously for a 24-hr period. Data of 12-hr recordings were used to analyze EPG variables. In each treatment, EPG signals of 80 individual *P. solenopsis* specimens were recorded, and no less than 20 sets of EPG data were analyzed.

Female adults' survival and body size

To evaluate the effect of water or honey solution acquisition on *P. solenopsis* survival rates, body size and survival time were examined for each specimen.

Three hundred newly-emerged adult female *P*. *solenopsis* were isolated individually in 9 cm Petri dishes prior to being randomly allocated to one of the following four treatments: (1) access to water (WM) (n = 80): a Petri dish containing a *P. solenopsis* specimen was covered with a 5 µm thick membrane and then upended over distilled water, which was changed daily; (2) access to honey solution (NM) (n = 80): a Petri dish containing a *P. solenopsis* to honey solution (NM) (n = 80): a Petri dish containing a *P. solenopsis* specimen was set up using the same procedure as the WM treatment, except

that they were upended over honey solution, which was changed daily; (3) starvation (SV) (n = 60): females were kept individually in empty Petri dishes covered by Parafilm; (4) control (CRL) (n = 80): each Petri dish base was lined with a piece of moist filter paper, on which G. hirsutum leaves were placed adaxially; the leaves were replaced daily. Each P. solenopsis specimen was teased with a fine brush into a non-feeding mode without injury, then carefully transferred onto the new leaves by a long-haired goat's wool brush. Each Petri dish was then covered by a 5 µm thick membrane. P. solenopsis in Petri dishes were checked twice daily (at 08:00 and 20:00) to record their survival rates. When the insects died, their body size (length and maximum width) was measured.

Statistics

Body size and EPG data were analyzed using analysis of variance (ANOVA) followed by Tukey's HSD test.

Results

EPG waveforms of *P. solenopsis* in water and honey solution conditions

In the WM treatment, the four major and diswaveforms tinct EPG recorded were characterized and labeled as A, B, C, and G (Figure 1). Waveform A occurred at the commencement of each probe; it showed the greatest amplitude, with high frequency and irregular waveforms (Figure 1A). Waveform B, which followed waveform A, was characterized by gradually decreasing amplitude over time and usually had fewer than 10 repetitions (Figure 1A). Waveform C coincided with or was superimposed on waveform B (Figure 1A). When waveform B faded away, waveform C remained with a continuation of the B wave periodicity, as in most of the signal shown in Figure 1B. Waveform G (Figure 1C) was the only waveform directly related to feeding activity and was characterized by a frequency of 3–6 Hz.

Observations of EPG waveforms recorded from *P. solenopsis* on artificial membranes revealed repetition in some sort of pattern at 12 hr intervals (Figure 2). During each 12 hr period, two distinct phases could be determined: a continuous feeding phase (phase I) and an alternating feeding phase (phase II, Figure 2). In phase I (Figures 1A, B, and C), *P. solenopsis* were generally feeding continuously, but some non-probing events were observed for a few individuals. In phase II, periods of non-feeding without new probing activities (~ 10 min) regularly alternated with periods of feeding (2–7 min) (Figure 1D).

In the NM treatment, although the same waveforms as above were recorded, some differences between the treatments were nevertheless present in the parameters of phases I and II (Figure 3). Compared to the WM data, in NM the duration of phase I was significantly shorter. The average duration of probing and the number of feedings in phase II were not significantly different between the treatments, but the duration of non-probing time was significantly longer in NM than in WM.

Comparison of *P. solenopsis* feeding behavior among different treatments

Due to the simple EPG waveforms recorded on the artificial membrane, analysis of *P. solenopsis* EPG waveforms in the previous report (Huang et al. 2012) was simplified as follows: (1) potential drop waveform was pooled into waveform C, and (2) data of phloem feeding-related waveforms were used to describe the sap-sucking activities (Huang et al. 2012), which were represented by waveform G in WM and NM treatments. The results of the comparison between the two

treatments and the CRL are shown in Table 1. P. solenopsis initiated probing into plant leaf tissue in 6 min, whereas it took at least 1 hr to occur in the NM treatment and even longer in the WM treatment. Phenacoccus solenopsis feeding on artificial membranes probed more often than when on plant tissue. Phenacoccus solenopsis on G. hirsutum leaves spent the longest time probing, followed by P. solenopsis in the WM treatment. In the feeding phases, P. solenopsis in NM spent the least time feeding; total feeding time was longest in WM. Phenacoccus solenopsis in NM spent the most time in non-probing activity, followed by WM; insects in the CRL spent the least time in non-probing activity (< 1.5 hr in a 24-hr period). The percentage of feeding time spent probing was calculated based on the above data. The time spent probing G. hirsutum leaves in the CRL $(32.20 \pm 4.65\%)$ was much shorter than in the other treatments $(86.67 \pm 2.91\%$ and $84.30 \pm 3.02\%$ in WM and NM, respectively).

Body sizes of *P. solenopsis* in different treatments

The effect of different food source treatments on *P. solenopsis* body size was significant (length: $F_{3,51} = 32.0$, p < 0.001; width: $F_{3,51} =$ 7.8, p < 0.001). Measured upon death, *P. solenopsis* in the SV, WM, and NM treatments were significantly smaller than those in the CRL (Figure 4). Body dimensions were 1–1.2 mm long and 0.5–0.7 mm wide in the three former treatments, compared with > 1.4 mm long and > 0.7 mm wide in the CRL treatment.

Survival of adult *P. solenopsis* in different treatments

Different feeding treatments significantly influenced survival of newly-emerged adult female P. solenopsis. Some individuals in the starved group began to die within five days, whereas adults in other groups survived at least one week (Figure 5). Survivorship was progressively higher for P. solenopsis in WM, NM, and CRL treatments (Figure 6). The effect of food treatments on longevity was significant ($F_{3.50} = 85.1$, p < 0.001). The average longevity of P. solenopsis was longest $(29.26 \pm 7.42 \text{ days})$ in the G. hirsutum leaf treatment compared to the other three treatments. The adults that fed on water and honey solution survived for 11.67 ± 3.5 days and 14.17 ± 2.79 days, respectively (no significant difference). Starved mealybug adults only lived for 6.83 ± 3.13 days.

Discussion

Compared to the structure of *G. hirsutum* leaf tissue, the artificial membrane system that *P. solenopsis* penetrated to get water was comparatively simple, so only two types of EPG waveforms were recorded, i.e., waveforms for penetration and water ingestion. These waveforms occurred alternately in two distinct feeding phases, a phase of continuous feeding and a phase of alternation between penetration and feeding. The continuous feeding phase could last for several hours, indicating that *P. solenopsis* could imbibe water. In previous work, EPG waveforms of *P. solenopsis* on *G. hirsutum* involving signals of ingestion from

	FPC parameters	Food resource		
	EPG parameters	Water	Honey solution	Cotton
	Time to 1st probe (s)	6965.99 ± 900.04 a*	4722.14 ± 643.94 b	324.92± 52.53 c
	Number of probes	9.25 ±2.21 a	10.5 ±3.05 a	6.19 ± 1.61 b
	Total probing time (s)	25101.47 ± 2381.29 b	13931.56 ± 2198.73 c	39170.12±1048.92 a
	Number of feeding phases	7.25 ± 3.23 a	$4.00 \pm 1.32 \text{ b}$	10.07± 4.01 a
	Total feeding phases (s)	21407.96 ± 2064.88 a	11554.26 ± 1898.25 b	15531.11 ± 979.63 b

* Data in the same row followed by a different letter indicate significant difference (p < 0.05) among treatments, determined by ANOVA followed by Tukey's HSD test. \ddagger Total time is 12 hr.

Journal of Insect Science | http://www.insectscience.org

xylem tissue were reported (Huang et al. 2012). Such xylem ingestion rarely occurred on G. hirsutum, but it could last for a considerable period of time once it commenced. This suggests that the P. solenopsis might ingest water, which was confirmed in the current study, in which a treatment with a supply of honey solution was set up to make a parallel comparison. The results showed that the feeding behavior characteristics of P. solenopsis on an artificial membrane with different liquids underneath were similar to some extent, showing the same pattern. Such a regular pattern did not occur when P. solenopsis were feeding on G. hirsutum, which might be an inherent habit for feeding attempts when P. solenopsis encounter only unsuitable hosts. Such a hypothesis has yet to be confirmed.

Phenacoccus solenopsis individuals consumed water and a honey solution when these were made available. Their feeding behavior on these liquids was examined and compared to that on G. hirsutum. EPG data comparison of P. solenopsis in different treatments revealed that *P. solenopsis* on *G. hirsutum* leaves began to penetrate the tissues within 10 min, whereas P. solenopsis on the artificial membrane did not start to probe until > 1 hr had passed. In addition, on G. hirsutum leaves $\sim 90\%$ of the recording time was associated with stylet activities, while only 58% and 32% of the time was spent on these activities on the artificial membranes over water and honey, respectively. Moreover, stylet evulsion during probing in cotton plant tissues occurred once per two hours, whereas such probing frequency was doubled on the artificial membranes. These results suggest that P. solenopsis accepts but does not prefer feeding on water. The results in this study were obtained using the EPG system with an artificial membrane. Water in the natural environment would be devoid of a surrounding membrane; thus,

whether *P. solenopsis* would directly imbibe free water still needs to be confirmed.

The effect of feeding on water in terms of P. solenopsis survival was also studied. The longevity in treatments was SV < WM/NM < CRL, suggesting that imbibing water can prolong P. solenopsis survival. However, feeding on water alone did not significantly increase P. solenopsis body size compared to feeding on G. hirsutum leaves. Feeding on honey solution may have prolonged P. solenopsis survival, but it did not increase their body size.

In conclusion, the results of the current study suggest that water-honey solution acquisition through an artificial membrane may be a result of inherent feeding habits and may be of survival benefit to P. solenopsis. This assumed inherent feeding behavior suggests that P. solenopsis might probe into any plant it contacts. If the content of plant phloem is ideal for *P. solenopsis* nutrition, the water in the phloem and xylem may also be of survival benefit. Phenacoccus solenopsis has been recorded on plants belonging to more than 100 genera in over 50 families (Abbas et al. 2009), but the suitability of many of these host plants has yet to be determined. If P. solenopsis will probe any plant, the suitability of different plant hosts must be examined to distinguish which hosts are permanently suitable and which serve as temporary hosts. Such information would be valuable for integrated pest management of P. solenopsis.

Acknowledgements

We thank Professor Robert Spooner-Hart from the University of Western Sydney and Gillian Watson from the California Department of Food and Agriculture for making a number of valuable suggestions and com-

ments. This research was supported by the Special Fund for Agro-Scientific Research in the Public Interest of China (No. 201103026), National Key Basic Research Program (973 Program, No. 2012CB722504), the National Natural Science Foundation of China (No. 31201523; No. 31270580), and the Ministry of Science and Technology of the EU scientific and technological cooperation in the special project (0904).

References

Abbas G, Arif MJ, Saeed S, Karar H. 2009. A new invasive species of genus *Phenacoccus* Cockerell attacking cotton in Pakistan. *International Journal of Agriculture and Biology* 11(1): 54–58.

Dutt U. 2007. *Mealy Bug Infestation in Punjab: Bt. Cotton Falls Flat.* Countercurrents. Available online: www.countercurrents.org/dutt210807.htm

EPPO. 2011. *New pest records in EPPO member countries*. EPPO Global Database.

Fand BB, Suroshe SS, Gautam RD. 2013. Fortuitous biological control of insect pests and weeds: a critical review. *The Bioscan* 8(1): 1–10.

Gary RE, Foster WA. 2004. *Anopheles* gambiae feeding and survival on honeydew and extra-floral honey solution of peridomestic plants. *Medical and Veterinary Entomology* 18: 102–107.

Giliomee JH. 2011. Recent establishment of many alien insects in South Africa—a cause for concern. *African Entomology* 19(1): 151–155.

Hodgson C, Abbas G, Arif MJ, Saeed S, Karar H. 2008. *Phenacoccus solenopsis* Tinsley (Sternorrhyncha: Coccoidea: Pseudococcidae), an invasive mealybug damaging cotton in Pakistan and India, with a discussion on seasonal morphological variation. *Zootaxa* 1913: 1–35.

Huang F, Zhang PJ, Zhang JM, Zhu YY, Lu YB, Zhang ZJ. 2011. Effects of three host plants on the development and reproduction of *Phenacoccus solenopsis* (Tinsley) (Sternorrhyncha: Coccoidea: Pseudococcidae). *Plant Protection* 37(1): 58–62.

Huang F, Tjallingii WF, Zhang PJ, Zhang JM, Lu YB, Lin JT. 2012. EPG waveform characteristics of solenopsis mealybug stylet penetration on cotton. *Entomologia Experimentalis et Applicata* 143(1): 47–54.

Huang F, Li WD, Lin WC, Lu YB, Zhang JM, Bei YW, Zhang ZJ. 2013. *An artificial spawning and feeding system for piercingsucking insects*. Chinese Patents: ZL 2009 1 0155767.0.

Jhala RC, Bharpoda TM, Patel MG. 2008. Occurrence of mealy bugs, *Phenacoccus solenopsis* Tinsley and *Phenacoccus soleni* Ferris (Hemiptera: Psedococcidae) on cotton in Gujarat. *Insect Environment* 13(4): 149– 151.

Karan D, Dahiya N, Munial AK. 1998. Desiccation and starvation tolerance of adult *Drosophila*: opposite latitudinal cline in natural populations of three different species. *Evolution* 52(3): 825–831.

Kehat M, Wyndham M, 1972. The effect of food and water on development, longevity, and fecundity in the Rutherglen bug, *Nysius*

vinitor (Hemiptera: Lygaeidae). Australian Journal of Zoology 20(2): 119–130.

Lundgren JG. 2009. Nutritional aspects of non-prey foods in the life histories of predaceous Coccinellidae. *Biological Control* 51(2): 294–305.

Mittler TE, Dadd RH. 1963. Studies on the artificial feeding of the aphid *Myzus persicae* (Sulzer)-I. Relative uptake of water and sucrose solutions. *Journal of Insect Physiology* 9(5): 623–645.

MOA and AQSIQ. 2009. Ministry of Agriculture of People's Republic of China (MOA), General Administration of Quality Supervision, Inspection and Quarantine of People's Republic of China (AQSIQ). The government announcement for adding solenopsis mealybug, *Phenacoccus solenopsis*, in the Chinese quarantine pest list, Beijing, No. 1147.

Rejmánek M. 2000. Invasive plants: approaches and predictions. *Austral Ecology* 25(5): 497–506.

Stockhoff BA. 1991. Starvation resistance of gypsy moth, *Lymantria dispar* (L.) (Lepidoptera: Lymantriidae): tradeoffs among growth, body size, and survival. *Oecologia* 88(3): 422–429.

Tjallingii WF. 1985. Electrical nature of recorded signals during stylet penetration by aphids. *Entomologia Experimentalis et Applicata* 38: 177–186.

Tjallingii WF, Hogen Esch Th. 1993. Fine structure of aphid stylet routes in plant tissues in correlation with EPG signals. *Physiological Entomology* 18: 317–328.

van Rijn PC, Tanigoshi LK. 1999. The contribution of extrafloral honey solution to survival and reproduction of the predatory mite *Inphiseius degenerans* on *Ricinus communis*. *Experimental and Applied Acarology* 23: 281–296.

Walker GP. 2000. A beginner's guide to electronic monitoring of homopteran probing behavior. In: Walker GP, Backus EA, Editors. *Principles and Applications of Electronic Monitoring and Other Techniques in the Study of Homopteran Feeding Behavior*. pp. 14–40. Thomas Say Publications in Entomology, Entomological Society of America.

Wang YP, Watson GW, Zhang RZ. 2010. The potential distribution of an invasive mealybug *Phenacoccus solenopsis* and its threat to cotton in Asia. *Agricultural and Forest Entomology* 12(4): 403–416.

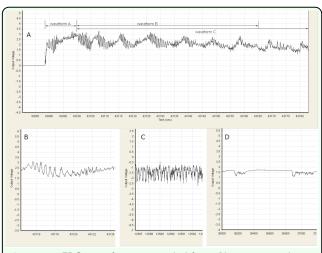


Figure 1. EPG waveforms recorded from *Phenacoccus solenopsis* in the WM treatment (artificial membrane on distilled water). A: waveforms of the phase I feeding pathway, including waveforms A, B and C; B: waveform C; C: waveform of water acquisition (waveform G); D: alternate feeding in phase II. High quality figures are available online.

Phase II Phase I 12 h

Figure 2. EPG signals of *Phenacoccus solenopsis* feeding on water on artificial membrane, showing incidence of phases I and II in repeated 12-hr period. High quality figures are available online.

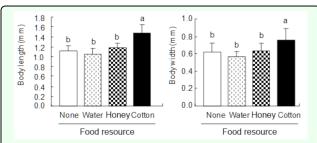
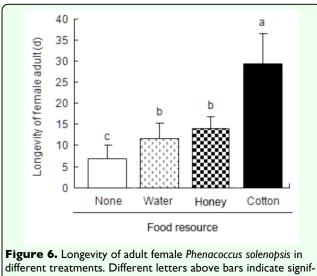


Figure 4. Body sizes of dying *Phenacoccus solenopsis* in different food source treatments (n = 80 per treatment). Different letters above the bars indicate significant difference at p < 0.05, as determined by ANOVA followed by Tukey's HSD test. High quality figures are available online.



different treatments. Different letters above bars indicate significant difference at p < 0.05 as determined by ANOVA followed by Tukey's HSD test. High quality figures are available online.

Journal of Insect Science | http://www.insectscience.org

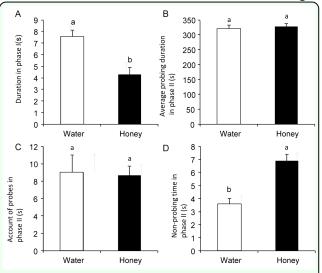


Figure 3. Characteristics of EPG waveforms of *Phenacoccus* solenopsis fed on water-honey solution through artificial membranes over 12 hr. Different letters above the bars indicate significant difference at p < 0.05, as determined by one-way ANOVA. High quality figures are available online.

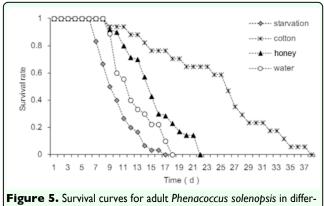


Figure 5. Survival curves for adult *Phenacoccus solenopsis* in different treatments from the time of emergence. High quality figures are available online.

Huang et al.