

# Morphometric comparisons of *Diaphorina citri* (Hemiptera: Liviidae) populations from Iran, USA and Pakistan

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The Asian citrus psyllid (ACP), *Diaphorina citri* Kuwayama (Hemiptera: Liviidae), vector of citrus greening disease pathogen, Huanglongbing (HLB), is considered the most serious pest of citrus in the world. Prior molecular based studies have hypothesized a link between the *D. citri* in Iran and the USA (Florida). The purpose of this study was to collect morphometric data from *D. citri* populations from Iran (mtCOI haplotype-1), Florida (mtCOI haplotype-1), and Pakistan (mtCOI haplotype-6), to determine whether different mtCOI haplotypes have a relationship to a specific morphometric variation. 240 samples from 6 ACP populations (Iran - Jiroft, Chabahar; Florida - Ft. Pierce, Palm Beach Gardens, Port St. Lucie; and Pakistan - Punjab) were collected for comparison. Measurements of 20 morphological characters were selected, measured and analysed using ANOVA and MANOVA. The results indicate differences among the 6 ACP populations (Wilks' lambda= 0.0376, F= 7.29, P <0.0001). The body length (BL), circumanal ring length (CL), antenna length (AL), forewing length (WL) and Rs vein length of forewing (RL) were the most important characters separating the populations. The cluster analysis showed that the Iran and Florida populations are distinct from each other but separate from the Pakistan population. Thus, three subgroups can be morphologically discriminated within *D. citri* species in this study, 1) Iran, 2) USA (Florida) and 3) Pakistan population. Morphometric comparisons provided further resolution to the mtCOI haplotypes and distinguished the Florida and Iranian populations.

Morphometric comparisons of *Diaphorina citri* (Hemiptera: Liviidae) populations from Iran, USA and Pakistan

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24 **ABSTRACT**

25 The Asian citrus psyllid (ACP), *Diaphorina citri* Kuwayama (Hemiptera: Liviidae), vector of citrus  
 26 greening disease pathogen, Huanglongbing (HLB), is considered the most serious pest of citrus in the  
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 29 Iran (mtCOI haplotype-1), Florida (mtCOI haplotype-1), and Pakistan (mtCOI haplotype-6), to  
 30 determine whether different mtCOI haplotypes have a relationship to a specific morphometric  
 31 variation. 240 samples from 6 ACP populations (Iran - Jiroft, Chabahar; Florida - Ft. Pierce, Palm  
 32 Beach Gardens, Port St. Lucie; and Pakistan - Punjab) were collected for comparison. Measurements  
 33 of 20 morphological characters were selected, measured and analysed using ANOVA and MANOVA.  
 34 The results indicate differences among the 6 ACP populations (Wilks' lambda= 0.0376, F= 7.29, P  
 35 <0.0001). The body length (BL), circumanal ring length (CL), antenna length (AL), forewing length  
 36 (WL) and Rs vein length of forewing (RL) were the most important characters separating the  
 37 populations. The cluster analysis showed that the Iran and Florida populations are separate from the  
 38 Pakistan population. Thus, two groups can be morphologically discriminated within *D. citri* species in  
 39 this study: 1) Iran and the USA (Florida), 2) the Pakistan population.

40 Keywords: Asian citrus psyllid, ACP, Citrus, Huanglongbing, HLB

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43 **INTRODUCTION**

The Asian citrus psyllid (ACP), *Diaphorina citri* Kuwayama (Hemiptera: Liviidae) (Burckhardt & Ouvrard, 2012), is the vector of the bacteria ‘*Candidatus Liberibacter spp.*’, the causal agent associated with Huanglongbing (HLB) or citrus greening disease (Bové, 2006; Grafton-Cardwell et al., 2013; Hall et al., 2013). Huanglongbing is considered the world’s most important disease of citrus (Gottwald, 2010; Grafton-Cardwell et al., 2013, Hall et al., 2013). The Asian citrus psyllid has been reported from the Arabian Peninsula, Afghanistan through to the Indian subcontinent, Japan, Taiwan, Hong Kong, China, the Philippine Islands, the Pacific Islands of Hawaii and Guam, the continental USA, the Caribbean, Central and South America and the Indian Ocean islands of Mauritius and Réunion (Boykin et al., 2012). In southern Iran, the ACP was discovered in 1997 followed by the HLB disease in 2006 (Bové et al., 2000; Faghihi et al., 2009) and now it has established in the citrus growing regions of Hormozgan, Sistan–Baluchistan, Kerman and Fars Provinces. In the USA, it was first reported from Florida in 1998 (Bové, 2006) and now occurs from Florida to California (Boykin et al., 2012). Also it was reported from Pakistan in 1927 (Husain & Nath, 1927), and has become a serious pest in all citrus growing areas of Pakistan (Mahmood et al., 2014).

Worldwide genetic diversity of *D. citri*, based on mitochondrial cytochrome oxidase I (mtCOI) DNA sequences, suggests the existence of eight haplotypes (Dcit-1 to Dcit-8) (Boykin et al., 2012). Haplotype-1 occurs in the following countries: United States of America (USA: Florida and Texas), India, Saudi Arabia, Brazil and Mexico. Haplotype 2 includes populations from Brazil (Sao Paulo), China (Fuzhou, Gangzhou), Indonesia (Java, Bali), Mauritius, Reunion, Taiwan (Taipei), Thailand (Hat Yai), and Vietnam (Hanoi). Haplotype 3 includes populations from Puerto Rico (Univ.of PR) and Guadeloupe. Haplotypes 4-8 includes populations from China (Gangzhou, Zhejaing), Florida and Mexico (Akil, Yucatan). An additional study revealed that *D. citri* populations from Iran are genetically similar to the mtCOI Haplotype-1 group, while the Pakistan population has been designated as mtCOI haplotype-6 (Lashkari et al., 2014). Further evidence supporting the haplotype grouping

comes from *Wolbachia*, *wDi*, *wsp* sequences which indicated that the Iran population was similar to the Florida population, but was different from the Pakistan population (Lashkari et al., 2014).

Morphologically, the psyllids within *Diaphorina* can be differentiated by the shape of the genal processes the shape and pattern coloration of the forewings, the arrangement of spinules on the forewing membrane, and the shape of the female terminalia (Hollis, 1987). Six morphological measurements, including body length, wing length and width, genal process length and width, and antenna length have been used to study the morphometry of ACP populations on six Rutaceae from Mexico (García-Pérez et al., 2013). Additionally, Vargas-Madríz et al. (2013) used 4 morphological indices including body length, body width, wing length, and wing width to describe the morphometry of another psyllid species, *Bactericera cockerelli* (Hemiptera: Triozidae), on two varieties of host plant.

The purpose of this study was to explore whether the different mtCOI haplotypes of ACP populations (mtCOI haplotypes 1 and 6) have correlate with specific morphometric variation.

80

## 81 MATERIAL AND METHODS

### 82 Psyllid samples

Six genetic based populations of *D. citri* were collected from Iran (2) and Florida (3) as mitochondrial COI Haplotype 1, and Pakistan as mitochondrial COI Haplotype 6. The collected specimens were preserved in 96% ethanol (Table 1). The female adults were selected for this study for direct comparison to the previous molecular study (Lashkari et al, 2014), which included only females. Also, it has been shown that the structure of the male genitalia within *Diaphorina* is homogeneous throughout and species are defined on the shape of genal cones, the shape and coloration of the forewings, the arrangement of spinules on the forewing membrane, and the shape of the female terminalia (Hollis, 1987).

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### 93 **Morphometric analysis**

94 A total of 240 female adults (40 adults from each population) were randomly selected for  
 95 morphological analyses. In order to calculate the morphometric information of the specimens, each  
 96 insect was dissected to separate the different structures. The selected specimens were placed  
 97 individually in 1.5 ml tubes containing 96% ethanol. Twenty standard morphological characters (Table  
 98 2) were selected to survey the morphometric variation among the six populations (Burckhardt, 1986;  
 99 Hollis, 1987; Ossiannilsson, 1992; Olivares & Burckhardt, 1997; Burckhardt & Basset, 2000; Mifsud  
 100 & Burckhardt, 2002; García-Pérez et al., 2013). Descriptions of the characters are given in Table 2 and  
 101 Fig.1. The body structures (except wings) were mounted on slides with glycerin and photographs were  
 102 taken of each structure/specimen using a digital camera coupled with a stereomicroscope with 40X  
 103 magnification. The right forewing of each specimen was slide-mounted using Euparal as the mounting  
 104 medium. All measurements (mm) were performed with National Instruments Vision Assistant  
 105 Software, version 2012 (National Instruments Corporation 2012).

106

### 107 **Statistical Methods**

108 Data were analyzed using analysis of variance (ANOVA) to compare different populations for each  
 109 character, and pairwise comparisons based on Tukey's HSD (Honest Significant Difference) test were  
 110 calculated only after a significant ANOVA was found. A multivariate analysis of variance  
 111 (MANOVA) was done for the comparison of the group means of all variables. The Wilks' lambda test  
 112 was applied as the statistical significance of the MANOVA. Moreover, the Canonical Variate Analysis  
 113 (CVA) was used to determine the relative importance of characteristics as discriminators between

groups. Mahalanobis distances (D2) were calculated between all populations' centroids using a pooled variance covariance matrix. All statistical analyses were conducted using the SAS statistical program (SAS Institute, 2003). The UPGMA (Unweighted Pair Group Method with Arithmetic Mean) hierarchical cluster analysis (Sneath & Sokal, 1973) based on squared Euclidean distances and the mantel tests were performed with NTSYS-pc program (Rohlf, 1993). Geographic distances among locations were measured using Google Earth (<https://earth.google.com>).

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## 121 RESULTS

According to univariate analysis, 15 morphological characters were found to be significantly different among the six ACP populations ( $\alpha = 0.01$ ). These included body length (BL) ( $F = 78.07$ ,  $df = 5$ ,  $P < 0.0001$ ), Vertex width (VW) ( $F = 4$ ,  $df = 5$ ,  $P = 0.0019$ ), antenna length (AL) ( $F = 11.63$ ,  $df = 5$ ,  $P < 0.0001$ ), forewing length (WL) ( $F = 28.50$ ,  $df = 5$ ,  $P < 0.0001$ ) and width (WW) ( $F = 11.19$ ,  $df = 5$ ,  $P < 0.0001$ ), Rs vein length of forewing (RL) ( $F = 26.92$ ,  $df = 5$ ,  $P < 0.0001$ ), length of the line connecting apices of vein Rs and Cu1a of forewing (RC) ( $F = 7.27$ ,  $df = 5$ ,  $P < 0.0001$ ), length of the line connecting the base and apex of vein M3+4 of forewing (b) ( $F = 6.38$ ,  $df = 5$ ,  $P < 0.0001$ ), length of the line connecting apices of veins M1+2 and M3+4 of forewing (c) ( $F = 9.69$ ,  $df = 5$ ,  $P < 0.0001$ ), length of the line connecting apices of vein Cu1a and Cu1b of forewing (d) ( $F = 15.58$ ,  $df = 5$ ,  $P < 0.0001$ ), length of widest perpendicular distance to d in cell cu1 (e) ( $F = 3.85$ ,  $df = 5$ ,  $P = 0.0025$ ), metatibial length (ML) ( $F = 5.19$ ,  $df = 5$ ,  $P = 0.0002$ ), female proctiger length (FP) ( $F = 14.69$ ,  $df = 5$ ,  $P < 0.0001$ ), circumanal ring length (CL) ( $F = 15.74$ ,  $df = 5$ ,  $P < 0.0001$ ) and female subgenital plate length (SL) ( $F = 8.94$ ,  $df = 5$ ,  $P < 0.0001$ ) (Table 3).

135 There was no statistical difference between populations with regards to the following characters: head  
136 width (HW), vertex length (VL), genal process length (GL), genal process width (GW) and length of  
137 the line connecting the base and apex of vein M1+2 of forewing (a).

138 The MANOVAs of the ACP populations revealed a significant difference among the size  
139 variables of the populations (Wilks' lambda= 0.0376, F= 7.29, P <0.0001). The shortest Mahalanobis  
140 distance (D2 = 0.720) was between the two populations from Florida (Palm Beach Gardens and Port  
141 St. Lucie), whereas the longest distance was between the populations from Pakistan and Florida (Palm  
142 Beach Gardens) (D2 = 36.756) (Table 4). The cluster analysis revealed two major clusters. The first  
143 contained samples from Iran (Jiroft, Chabahar) and Florida (Ft. Pierce, Port St. Lucie, Palm Beach  
144 Gardens) and the second one contained the Pakistan population (Fig. 2).

145 The canonical discriminant analysis indicated that the first two canonical variables (CVA1 and  
146 CV2) described 65.15 % and 24.85 % of the total variance, respectively. The first and second together  
147 (CVA1+CVA2) equaled 90 % (Table 5). The body length (BL), circumanal ring length (CL), antenna  
148 length (AL), forewing length (WL) and Rs vein length of forewing (RL) contributed most to this  
149 variation based on the first canonical axes (CVA1). Other characteristics also contributed, but to a  
150 lesser extent (Table 5).

151 The Mantel test showed that there was not significant correlation between geographic and  
152 morphological distances ( $r = 0.535$ ,  $p = 0.999$ ). Therefore, geographical distances did not impact the  
153 morphological differentiation found between the populations.

## 154 155 **DISCUSSION** 156



The morphometric analyses of the ACP populations from Iran, USA (Florida) and Pakistan indicated the existence of two main groups within the populations analyzed. The first group included populations from Iran (Jiroft and Chabahar) and the USA (Florida), and the second was represented by a population from Pakistan. These results support similar findings from wing structures of ACP from Iran and Pakistan (Lashkari et al., 2013). The results presented here also support previous findings indicating that *D. citri* populations in Iran and Florida are similar and separated from Pakistan populations based on a global phylogenetic analysis of mtCOI, and *Wolbachia* wsp sequences (Lashkari et al., 2014). Prior molecular based studies showed that all Iranian populations of ACP are genetically similar to the Florida populations indicating a link between the ACP in Iran and the USA (Florida) (Lashkari et al., 2014).

The morphometric data provides further resolution to the previous molecular research, which indicated that different mtCOI haplotypes of ACP populations (mtCOI haplotypes 1 and 6) correlate with specific morphometric variation. As Iran and the USA (Florida) populations (Haplotype 1) were distinguishable from Pakistan population (Haplotype 6) using mtCOI. Understanding the link between morphological and molecular characters is of vital importance for designing diagnostic tests for highly invasive species to aid global biosecurity (Boykin et al., 2012).

The Mantel test results showed that the separation of the Iran and Florida populations in this study was not due to the geographic distance. García-Pérez et al. (2013) have shown the separation of host-associated populations of ACP. They showed that the host species or variety can influence morphometric traits of different host associated populations of ACP (García-Pérez et al. 2013). They indicated that the largest ACP populations were associated with *C. sinensis* (L.) Osbeck cv. 'Marrs', *C. sinensis* (L.) cv. 'Valencia' and *Murraya paniculata* (L.) Jack, while, the smallest sizes were found in males collected from *Citrus limetta* Risso, *C. sinensis* (L.) 'Selection 8' and *C. paradisi* Macfad. In the

present study the populations from Iran that were collected from the same host species (*C. sinensis*) that were clustered together and the Florida populations that were collected from *M. paniculata* were clustered together, while the Florida population collected from *C. macrophylla* was separate. The main purpose of the present study is to explore whether the mtCOI haplotypes 1 and 6 of ACP populations have specific morphometric variation. We investigated the morphological characteristics of populations from Iran and Florida as Haplotype-1 *Diaphorina citri*, and the Pakistan population as *D. citri* Haplotype-6 that were defined in Lashkari et al. (2014).

In the current study, the body length (BL), circumanal ring length (CL), antenna length (AL), forewing length (WL) and Rs vein length of forewing (RL) contributed most to the variation found among the six populations. These results were similar to a previous study conducted by Garcia-Perez et al. (2013). They indicated that wing length, wing width and body length were the main variables contributing to discrimination of populations of *D. citri* on various host plants in Mexico. A comparison of the female ACP body size from the populations in the current study with those from different countries (Mexico, Réunion, Venezuela, and India) indicated that the populations from Iran and Florida were most similar to those from India (body length 2.4 mm; forewing length 2.17 mm), while the Pakistan populations stood alone while being shorter than the others (García-Pérez et al., 2013; Étienne et al., 2001; Fonseca et al., 2007; Mathur, 1975; Chhetry et al. 2012).

There are two pieces of evidence that suggest the invasion of ACP into Iran and the USA (Florida) originated from southwestern Asia, particularly India: 1- Southwestern Asia, i.e. India, has been suggested as the origin of ACP based on plant host origins and historical information (Hall, 2008). 2- The mitochondrial haplotype network for *D. citri* suggests a basal and thus ancestral position for Dcit-1 haplotype (Boykin et al., 2012). Boykin et al. (2012) showed that the Indian, USA, Saudi Arabian, Brazilian and Mexican populations of ACP belong to the mtCOI Haplotype-1 group.

204 However, additional studies based on the morphological and other molecular markers such as  
205 microsatellite on the phylogenetic relationships among worldwide ACP populations are needed to  
206 confirm our hypothesis. Boykin et al. (2007) developed twelve polymorphic microsatellite markers for  
207 ACP and should be explored on a global scale.

208 The differentiation of populations may originate from one of the following events: insect  
209 migration, a new host or a new habitat or both of them, landscape changes (bottleneck effect), and  
210 genetic changes by stochastic events, such as gene flow, genetic drift and mutation or natural selection  
211 (Kim & McPheron, 1993; Berlocher & Feder, 2002). These variations may be changes in morphology,  
212 physiology, behavior, and life history traits, and subsequently would lead to the manifestation of the  
213 different taxonomic status of local populations such as biotype and ecotype (Kim & McPheron, 1993).

214 We conclude that *D. citri* populations related to the mtCOI haplotypes-1 (Iran and Florida) and  
215 6 (Pakistan) have distinct morphometric characters based on multivariate analysis of morphological  
216 data. Future ACP studies are needed to confirm the relationship found here between the mtCOI  
217 haplotypes and morphology.

218

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310 (Hemiptera: Triozidae), grown on two varieties of tomato under greenhouse conditions. *Florida*  
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**Table 1** (on next page)

Table 1

Collection sites, mtCOI Haplotype, hosts and number of examined specimens for populations of *Diaphorina citri*.



2

3 **Table 1** Collection sites, mtCOI Haplotype, hosts and number of examined specimens for populations  
4 of *Diaphorina citri*.

Country	Province/State	County	mtCOI Haplotype	Host	<i>n</i>
IRAN	Sistan & Baluchestan	Chabahar	1	<i>Citrus sinensis</i> (L.) Osbeck	40
	Kerman	Jiroft	1	<i>Citrus sinensis</i> (L.) Osbeck	40
		Palm Beach Gardens, Palm Beach County	1	<i>Murraya paniculata</i> (L.) Jack.	40
USA	Florida	Port St. Lucie, St. Lucie County	1	<i>Murraya paniculata</i> (L.) Jack.	40
		USDA ARS colony, Ft. Pierce, St. Lucie County	1	<i>Citrus macrophylla</i> Wester	40
Pakistan	Punjab	Punjab	6	<i>Citrus sinensis</i> (L.) Osbeck	40

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6

**Table 2** (on next page)

Table 2

Morphological traits used for morphometric analysis of populations of *Diaphorina citri*.

**Table 2** Morphological traits used for morphometric analysis of populations of *Diaphorina citri*.

Character No.	Acronym	Character
1	BL	Body length (from the apex of the genal process to the distal part of the proctiger)
2	HW	Head width
3	VW	Vertex width
4	VL	Vertex length
5	GL	Genal process length
6	GW	Genal process width
7	AL	Antenna length
8	WL	Forewing length
9	WW	Forewing width
10	RL	Rs vein length of forewing
11	RC	Length of the line connecting apices of vein Rs and Cu1a of forewing
12	a	Length of the line connecting the base and apex of vein M1+2 of forewing
13	b	Length of the line connecting the base and apex of vein M3+4 of forewing
14	c	Length of the line connecting apices of veins M1+2 and M3+4 of forewing
15	d	Length of the line connecting apices of vein Cu1a and Cu1b of forewing
16	e	Length of widest perpendicular distance to d in cell cu1
17	ML	Metatibial length
18	FP	Female proctiger length
19	CL	Circumanal ring length
20	SL	Female subgenital plate length

**Table 3**(on next page)

Table 3

Size (MM $\pm$ SE) of the 20 morphological traits in the populations of Asian citrus psyllid, *Diaphorina citri* from Iran, Florida and Pakistan.

**Table 3** Size (MM±SE) of the 20 morphological traits in the populations of Asian citrus psyllid, *Diaphorina citri* from Iran, Florida and Pakistan.

Variable <sup>a</sup>	Population					
	Iran-Chabahar	Iran-Jiroft	Florida- USDA ARS colony	Florida- Palm Beach County	Florida- St. Lucie County	Pakistan
BL	2.477±0.029 a <sup>b</sup>	2.459±0.029 a	2.49±0.018 a	2.557±0.009 a	2.535±0.013 a	1.980±0.035 b
HW	0.575±0.002 a	0.569±0.002 a	0.558±0.005 a	0.573±0.003 a	0.567±0.004 a	0.573±0.006 a
VW	0.375±0.003 a	0.372±0.003 ab	0.359±0.003 b	0.376±0.003 a	0.370±0.003 ab	0.364±0.004 ab
VL	0.136±0.002 a	0.132±0.002 a	0.136±0.003 a	0.141±0.002 a	0.139±0.003 a	0.137±0.004 a
GL	0.122±0.003 a	0.121±0.003 a	0.124±0.003 a	0.132±0.002 a	0.128±0.003 a	0.122±0.004 a
GW	0.101±0.002 a	0.095±0.002 a	0.097±0.002 a	0.103±0.002 a	0.102±0.002 a	0.094±0.003 a
AL	0.447±0.006 a	0.441±0.006 a	0.436±0.005 a	0.447±0.003 a	0.442±0.004 a	0.403±0.003 b
WL	1.990±0.014 c	2.091±0.012 b	2.151±0.011 ab	2.172±0.009 a	2.164±0.010 a	2.090±0.017 b
WW	0.869±0.007 b	0.910±0.006 a	0.909±0.004 a	0.923±0.003 a	0.914±0.004 a	0.912±0.006 a
RL	1.135±0.010 c	1.222±0.007 ab	1.246±0.008 ab	1.261±0.006 a	1.250±0.007 a	1.206±0.012 b
RC	0.728±0.008 b	0.760±0.007 a	0.762±0.005 a	0.775±0.005 a	0.770±0.005 a	0.775±0.007 a
a	0.555±0.005 a	0.592±0.005 a	0.579±0.004 a	0.768±0.179 a	0.587±0.003 a	0.571±0.006 a
b	0.490±0.005 b	0.520±0.004 a	0.499±0.003 b	0.508±0.001 ab	0.504±0.003 ab	0.503±0.005 ab
c	0.319±0.004 b	0.336±0.005 ab	0.350±0.004a	0.355±0.002 a	0.353±0.003 a	0.337±0.006 ab
d	0.409±0.004 c	0.423±0.005 cb	0.440±0.003 ab	0.446±0.002 a	0.445±0.003a	0.426±0.003 cb
e	0.259±0.002 b	0.260±0.003 ab	0.259±0.002 ab	0.266±0.002 ab	0.264±0.002 ab	0.271±0.003 a
ML	0.563±0.005 a	0.557±0.005 ab	0.540±0.003 b	0.556±0.003 ab	0.548±0.004 ab	0.539±0.004 b
FP	0.511±0.007 a	0.509±0.007 a	0.479±0.004 b	0.495±0.004 ab	0.489±0.004 ab	0.450±0.007 c
CL	0.133±0.001 ab	0.130±0.001 b	0.133±0.003 ab	0.141±0.002 a	0.138±0.003 ab	0.116±0.002 c
SL	0.423±0.005 a	0.421±0.005 a	0.409±0.004 a	0.418±0.003 a	0.414±0.003 a	0.386±0.005 b

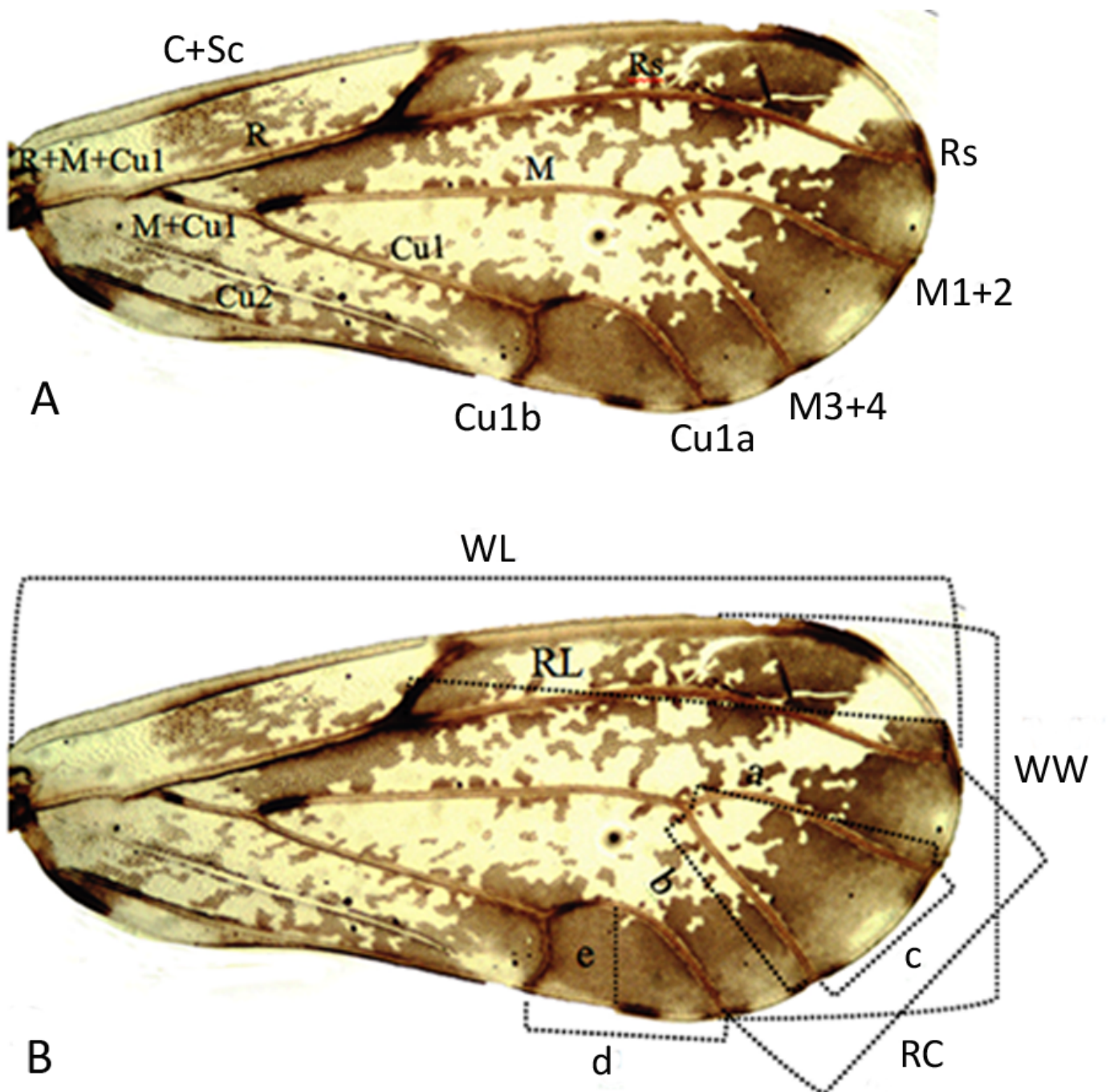
<sup>a</sup> See Table 2 for abbreviations.

<sup>b</sup> Means with the same letter within each variable are statistically equal (Tukey,  $P \leq 0.01$ ).

1

Figure 1

Forewing vein terminology based on Hodkinson and White (1979), and B- lines indicating measurements based on Mifsud and Burckhardt (2002) in the right forewing of *Diaphorina citri*. See Table 2 for abbreviations.



**Table 4**(on next page)

Table 4

Mahalanobis distances among populations of *Diaphorina citri* from Iran, Florida and Pakistan (the below diagonal).

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3

4 **Table 4** Mahalanobis distances among populations of *Diaphorina citri* from Iran, Florida and  
5 Pakistan (the below diagonal).

Population	Iran-Chabahar	Iran-Jiroft	Florida- USDA ARS colony	Florida- Palm Beach County	Florida- St. Lucie County	Pakistan
Iran-Chabahar	0					
Iran-Jiroft	7.486	0				
Florida- USDA ARS colony	18.277	9.264	0			
Florida- Palm Beach County	16.603	8.999	2.644	0		
Florida- St. Lucie County	15.511	8.526	1.127	0.720	0	
Pakistan	24.478	23.119	32.071	36.756	33.397	0

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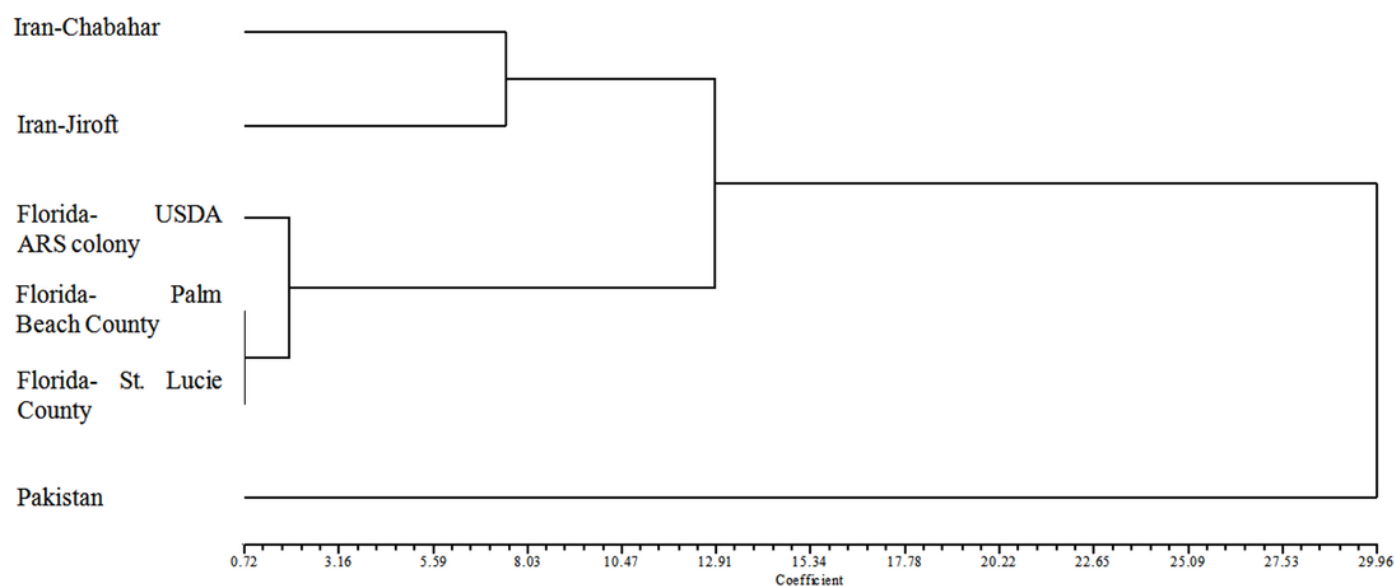
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# 2

Figure 2

Dendrogram plotted by UPGMA method based on squared Euclidean distance of *Diaphorina citri* populations.



# **Table 5**(on next page)

Table 5

Standardized coefficients for canonical variables on the first (CVA1) and second (CVA2) canonical axes.

2

3 **Table 5** Standardized coefficients for canonical variables on the first (CVA1) and second (CVA2)  
 4 canonical axes.

5

Variable <sup>a</sup>	CVA1	CVA2
BL	0.853	0.390
HW	-0.127	0.147
VW	0.068	0.261
VL	0.048	-0.079
GL	0.151	-0.107
GW	0.200	0.091
AL	0.453	0.356
WL	0.427	-0.660
WW	0.151	0.498
RL	0.417	-0.616
RC	0.039	-0.462
a	0.093	-0.046
b	0.057	-0.094
c	0.317	-0.446
d	0.390	-0.518
e	-0.156	-0.263
ML	0.083	0.398
FP	0.307	0.572
CL	0.576	0.160
SL	0.340	0.408
Eigenvalues	4.4741	1.7065
Proportion	0.6515	0.2485

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<sup>a</sup> See Table 2 for abbreviations.

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