Title: Mutualism enhances *Wolbachia* infection rates in ant-attended *Tuberculatus* aphid species (Hemiptera: Aphididae)

Author name: Izumi Yao[1]

1: Department of Ecology and Systematics, Graduate School of Agriculture, Hokkaido University

Corresponding Author: Izumi Yao

Address: Kitaku, Kita9 Nishi9, Sapporo 060-8589, Japan

E-mail: iyao@res.agr.hokudai.ac.jp

Abstract

Some aphid species form close associations with ants: offering them honeydew and obtaining protection from ants in return. However, mutualistic interactions with ants can also have a negative influence on aphid physiological and morphological traits. *Wolbachia* are intracellular bacteria whose major genotypes are classified into 17 supergroups (A to S except G and R). Aphid species within the genus *Tuberculatus* feed on Fagaceae leaves and exhibit two contrasting ecological characteristics, ant-attendance and non-attendance. Previous work has found that ant-attended species exhibit lower dispersal and are likely to form aggregated colonies. Considering that host-parasitoid interactions may well be one of the most common horizontal transmission routes of *Wolbachia*, it is therefore expected that ant associations will be associated with higher *Wolbachia* infection rate in *Tuberculatus* aphid species. This study compared *Wolbachia* infection rates between 11 ant-attended and 12 non-attended *Tuberculatus* aphid species, which were collected throughout Japan and around Mt. Kariwangsan in South Korea. Mean infection rates of *Wolbachia* were 30.2% in ant-attended species and 3.1% in non-attended species. The *Wolbachia* haplotypes detected were classified into supergroups B, M, N, and O. A phylogenetic tree of *Tuberculatus*

aphids constructed from a mitochondrial gene of cytochrome oxidase subunit I (*COI*) and nuclear gene of *18S rRNA* was used to examine the correlation between *Wolbachia* infection rates and ant associations. The phylogenetic comparative analysis showed that *Wolbachia* infection rates were significantly higher in ant-attended species. Possible *Wolbachia* infection routes are discussed in terms of the differences in the ecological characteristics between ant-attended and non-attended aphid species. This study shows that the spread of microorganisms is affected by host species interactions, and contributes new insights into the evolution of mutualistic interactions.

43 **Introduction**

44 Aphids (Insecta: Hemiptera: Aphididae) feed on plant phloem sap using their sucking and
45 piercing mouthparts, and excrete honeydew including carbohydrates and amino acids (Yao
46 and Akimoto 2001, 2002; Leroy et al. 2011; Renyard et al. 2021). Some aphid species form
47 close associations with ants by offering them honeydew and obtain protection from ants in
48 return (Way 1963; Yao et al. 2000). However, there is growing evidence that mutualistic
49 interactions with ants can have negative effects on aphid physiological and morphological
50 traits, such as changes in sugar and amino acid composition of honeydew (Fischer and
51 Shingleton 2001; Yao and Akimoto 2001, 2002) and decreases in colony size, body size and
52 embryo numbers (Stadler and Dixon 1998; Flatt and Weisser 2000; Yao et al. 2000;
53 Katayama and Suzuki 2002). These examples show that evolution of ant-aphid interactions
54 has resulted in both benefits and costs to aphids (Stadler and Dixon 2005; Yao 2014).

55 Recently, a number of studies have raised the possibility that microorganisms are
56 involved in the establishment of aphid-ant mutualisms. For example, the bacterium
57 *Staphylococcus xylosus* in *Aphis fabae* produces a blend of semiochemicals that attracts ant

58 scouts (Fischer et al. 2015). Additionally, it is reported that the ant *Lasius niger* could

59 potentially use cuticular hydrocarbons cues to discriminate among aphid lines (*Aphis fabae*)

60 harbouring different endosymbionts (Hertaeg et al. 2021). Henry et al. (2015) demonstrated

61 that two symbiont species, *Hamiltonella defensa* and *Regiella insecticola*, which protect

62 aphids from natural enemies (Oliver et al. 2003; Scarborough et al. 2005; Vorburger et al.

63 2010), were more likely to occur in aphid species that are *not* tended by ants.

64 *Wolbachia* are intracellular bacteria that occur in arthropods and nematodes (Werren

65 et al. 2008; Kauer et al. 2021). It is suggested that more than half of arthropod species are

66 infected with *Wolbachia* (Hilgenboecker et al. 2008; Weinert et al. 2015). At present, it has

67 been reported that the major genotypes of *Wolbachia* are highly diverse and classified

68 phylogenetically into 17 supergroups (A to S except for G and R) (Glowska et al. 2015;

69 Lefoulon et al. 2020). The roles of *Wolbachia* in hosts range from parasitism to mutualism.

70 *Wolbachia* infection can alter host reproduction by inducing feminization, parthenogenesis,

71 male killing, and cytoplasmic incompatibility (Werren et al. 2008). By contrast, *Wolbachia*

72 has been observed associating mutualistically with the bedbug *Cimex lectularius*, providing

73 B vitamins to the host (Hosokawa et al. 2010).

74 *Wolbachia* has been found in some aphid species (Gómez-Valero et al. 2004; Wang et

75 al. 2009; Augustinos et al. 2011; De Clerck et al. 2014; Yao 2019; Ren et al. 2020).

76 However, the roles of *Wolbachia* in host aphids are unknown. De Clerck et al. (2015)

77 claimed that *Wolbachia* in the banana black aphid *Pentalonia nigronervosa* could provide

78 nutrition to the host by association with *Buchnera aphidicola*, the primary endosymbiont of

79 aphids, while Manzano-Marín (2020) rejected the nutrition provision hypothesis by arguing

80 that it was based on a biased interpretation of antibiotic treatment analyses and incorrect

81 genome-based metabolic inference.

82 *Tuberculatus* aphids feed on Fagaceae (oak, chestnut, and beech) leaves and do not

83 alternate host plants during the season (Quednau 1999) (Table S1). This group encompasses

84 species with two contrasting ecological characteristics, ant-attendance and non-attendance

85 (Yao 2011). In a previous phylogenetic independent contrasts analysis, it was found that

86 ant-attended species have higher wing loading (the ratio of wing area to body size) (Yao

87 2011), suggesting that ant-attended aphids have allocated more resources to their bodies

88 than to their wings, resulting in lowered dispersal. Lower dispersal is likely to result in the
89 formation of aggregated colonies (Stadler et al. 2003). It has also been demonstrated that
90 ant-attended colonies attract more parasitoid wasps compared to ant-excluded colonies
91 (Völkl 1992; Kaneko 2002, 2003; Sadeghi-Namaghi and Amiri-Jami 2018). Considering
92 that host-parasitoid interactions may well be one of the most common horizontal
93 transmission routes of *Wolbachia* (reviewed by Sanaei et al. 2021), it is expected that ant
94 associations will be associated with higher *Wolbachia* infection rates in *Tuberculatus* aphid
95 species.

96 This study (1) examined *Wolbachia* infection rates and the type of *Wolbachia*
97 supergroup in *Tuberculatus* aphid species collected throughout Japan and around Mt.
98 Kariwangsan in South Korea, (2) estimated molecular phylogenetic trees based on a
99 mitochondrion gene and a nuclear gene, and (3) evaluated the correlation with *Wolbachia*
100 infection rates and ant associations using a phylogenetic comparative method. Infection
101 routes of *Wolbachia* to aphids are discussed in terms of horizontal transmission via
102 parasitoid wasps and ants.

103

104 **Materials & Methods**

105 **DNA extraction and *Wolbachia* infection rate**

106 *Tuberculatus* aphids (Aphididae: Calaphidini), 11 ant-attended and 12 non-attended species

107 (Table 1 and Table S1), were collected from regions throughout Japan and around Mt.

108 Kariwangsan of South Korea (Fig. S1 and Table S2). A species was regarded as

109 ant-attended if aphids offered honeydew directly from their anus to attending ants. Because

110 it was difficult to physically identify in the field three of the ant-attended aphid species (*T.*

111 *fulviabdominalis*, *T. indicus*, and *T. pilosulus*) and seven of the non-attended aphid species

112 (*T. higuchii* A- and B-types, *T. kashiwae* A- and B-types, *T. yokoyamai*, *T*. sp. D, and *T*. sp.

113 F), those species were identified through the genetic sequencing (Table S1). Sampling was

114 conducted on viviparous females, which appears from April to September. Since

115 *Tuberculatus* aphid species parthenogenetically produce nymphs in summer, several nymph

116 individuals on a leaf are a high likely to be clones. Therefore, aphids were collected from

117 more than ten leaves in a tree, to avoid collecting clonal aphids. Individuals were placed

118 into 99.5% ethanol and stored at -20℃. Before DNA extraction, the collected aphids were

119 dissected to check for the presence of parasitoid wasps. Aphids with parasitoid wasps were

120 excluded from DNA extraction. Total DNA was extracted from each dissected aphid (whole

121 body) with the Wizard genomic DNA purification kit (Promega, Tokyo, Japan). Since the

122 *16S rRNA* gene is highly conserved in a wide variety of microorganisms, it was used for

123 polymerase chain reaction (PCR) amplification to determine the presence or absence of

124 *Wolbachia*. In the small-scale experiment, using a gene map of the *16S rRNA* locus of

125 *Wolbachia* (Simões et al. 2011), seven pairs of primers were selected and tested for each of

126 23 *Tuberculatus* species, in which two to three individuals per species were tested (Table 2).

127 One pair of primers, *16SWolbF* (*16S-3f*) (Casiraghi et al. 2001) and *WspecR* (*16S-2r*)

128 (Werren and Windsor 2000), was identified as the most appropriate for assessing the 23

129 species because it was able to amplify *Wolbachia* at the maximum number of species

130 (seven species) of the 23 species (Table 3). After the small-scale experiment, a full-scale

131 experiment using the pair of primers was conducted on all collected samples (Table 1). To

132 check whether DNA extraction was successful, the barcoding region (in mitochondrion) of

133 primer pairs, *LCO1490* and *HCO2198*, was also used (Table 2). Because more than 90% of

134 individuals of *T. macrotuberculatus* in the Ishikari site (site 4 in this study) harboured

135 *Wolbachia* (Yao 2019), one individual of the species from the site was used for a positive

136 control sample for *Wolbachia* detection. PCR was performed in 10 μL volumes which

137 included 2 μL of 5×KAPATaq Extra buffer (Nippon Genetics, Tokyo, Japan), 1 μL 25 mM

138 $MgCl_2$, 0.3 μL dNTP mixture (10 mM of each), 0.5 μL of 10 μM of each primer, 1 μL

139 template DNA, and 0.05 μL KAPATaq Extra DNA polymerase (5 units/ μL). Reaction

140 cycle parameters were: 94 °C for 1 min; 40 cycles of 94 °C for 20 sec, 50 °C for 20 sec,

141 and 68 °C for 1 min, followed by a final extension of 68 °C for 1 min. When PCR products

142 had faint bands, the samples were rechecked by PCR in 20μL reaction volume. If the bands

143 were false, nothing was amplified in 20μL reaction volume. The PCR product was checked

144 using 1.5% agarose gel electrophoresis with ethidium bromide stain illuminated by UV

145 light. The *Wolbachia* infection rate of each species was defined as the percentage of

146 individuals amplified with the *Wolbachia*-specific primer out of all individuals amplified

147 with the barcoding region primer. The correlation between the *Wolbachia* infection ratio in

148 each collection site and geographical distance was tested by a Mantel test (Mantel 1967)

149 using the package vegan (Oksanen et al. 2012) in R (R Development Core Team 2021). The

150 values of latitude and longitude of collection sites were obtained from Google Maps and

151 were used for the geographic distance matrix. *Wolbachia* infection rates at the collection

152 sites were used for an environmental parameter distance matrix. Except for exhaustive

153 infection of *T.* sp. B, a Mantel test was applied to the species that was collected from more

154 than a single site.

155

156 **Phylogenetic trees for *Tuberculatus* aphids**

157 A phylogenetic tree of the 23 *Tuberculatus* aphid species was constructed from the

158 nucleotide sequences of a mitochondrion gene of a partial of cytochrome oxidase subunit I

159 (*COI*) (940bp) from DDBJ (DNA Data Bank of Japan) (Table 1). Besides the *COI* gene, a

160 partial of the nuclear gene of *18S rRNA* (approx. 670bp) was amplified and used to

161 construct phylogenetic trees. For reading the sequences of *18S rRNA* gene, PCR was

162 performed in 20µL reaction volume with a pair of primers (*Ns1* and *Ns2a*; Table 2), the

163 same reagents, and reaction cycles, as mentioned in the previous section were used, but the

164 annealing temperature was changed to 47 °C. PCR products were purified and sent to a

165 sequencing service (using Sanger sequencing) (Eurofins, Japan). The sequence data of the

166 *18S rRNA* gene (515bp) were deposited in the DDBJ and accession numbers are listed in

167 Table 1. A combined sequence of *COI* and *18S rRNA* genes (1,455bp) was used for the

168 construction of phylogenetic trees. The appropriateness of the combined sequence was

169 checked by a homogeneity test implemented in PAUP* 4.0b10 PPC (Swofford 2002) ($P >$

170 0.05). Maximum likelihood (ML) analysis was performed using PAUP* 4.0a 169. For the

171 ML tree, parameters were chosen based on the Akaike Information Criterion, as

172 implemented in Modeltest ver 3.7 (Posada and Crandall 1998). The GTR + I + G model

173 was selected for the combined sequence of *COI* and *18S rRNA* genes. ML trees were

174 searched heuristically with TBR branch swapping. For the bootstrap test on ML, 1,000

175 replicates were performed using fast stepwise addition as a starting option. Because

176 phylogenetic tree for the comparative analysis of independent contrasts must be fully

177 dichotomous with no gaps in the data, outgroup species were excluded from the analysis.

178

179 **Phylogenetic independent contrasts**

180 As a consequence of their common ancestry, closely related species share many
181 characteristics, and similarity between lineages is often influenced by relatedness rather
182 than by independent evolution. Most statistical tests assume independence of data points
183 and, therefore, data that are phylogenetically non-independent will tend to inflate the
184 degrees of freedom (Felsenstein 1985; Harvey and Pagel 1991). Comparative analysis by
185 independent contrasts (CAIC) uses independent comparisons of components within a
186 phylogeny, with each comparison being made at a different nodes in the phylogeny (Purvis
187 and Rambaut 1995). To examine the correlation between *Wolbachia* infection rates
188 (continuous data as dependent variables) and ant association (discrete data as independent
189 variables) in *Tuberculatus* species, phylogenetically independent contrasts were calculated
190 using the pic function implemented in the package ape (Paradis and Schliep 2019) in R.
191 Discrete data of ant association were coded as continuous variable using the contr.treatment
192 function in R. The extent of ant association was categorized as either 0 (non-attendance) or

193 1 (facultative and obligate ant-attendance). *Wolbachia* infection rates were arcsine-square

194 root transformed before analysis. The regression of contrasts between ant association and

195 *Wolbachia* infection rates passes through the origin (the intercept is set to zero) as

196 recommended by Garland et al. (1992).

197

198 ***Wolbachia* supergroups**

199 For *Wolbachia* that were detected in aphids (Table S2), the PCR products were sequenced

200 with the same primers (*16S-3f* and *16S-2r*) (Table 2). PCR products were purified with

201 FastGene Gel/PCR Extraction Kit (Nippon Genetics, Tokyo, Japan). The cycle sequencing

202 reaction was performed with a 5 μL volume consisting of 2 μL of Quick Start Mix

203 (Beckman Coulter, Tokyo, Japan), 0.5 μL of 10 μM forward or reverse primers, and 2.5 μL

204 of 10 ng/μL template DNA. The reaction cycle was 40 cycles of 94 °C for 20 sec, 50 °C for

205 20 sec, and 60 °C for 1 min. DNA sequencing was analyzed using the CEQ2000XL DNA

206 Analysis System (Beckman Coulter, Tokyo, Japan). The length of sequences that were

207 successfully read through the samples were from about 500bp to 900bp.

208 Multiple sequence alignments including the sequences of 16 *Wolbachia* supergroups (A, B,
209 C, D, E, F, H, I, J, K, L, M, N, O, Q, S) that were cited by Bing et al. (2014) (A to O), Ren
210 et al. (2020) (O found in aphids), Glowska et al. (2015) (Q), and Lefoulon et al. (2020) (S)
211 (Table S3) were processed with Clustal W (Thompson et al. 1994) on the DDBJ.
212 Supergroup P was not included in multiple sequence alignments because it had insufficient
213 sequence length for the lower region of the gene. After multiple sequence alignments, the
214 length of sequences was 471bp. To determine what types of *Wolbachia* supergroup are
215 present in *Tuberculatus* aphids, neighbor joining (NJ) with the BioNJ method was applied
216 to the constructed *Wolbachia* phylogenetic tree. NJ analysis was performed using PAUP*
217 4.0a 169. The distance matrix was calculated using the Jukes-Cantor substitution model.
218 For the bootstrap test on NJ, 1,000 replicates were performed using fast stepwise addition
219 as a starting option.
220

221 **Results**

222 ***Wolbachia* infection rate**

223 *Wolbachia* was detected in eight of 11 ant-attended aphid species (Table 1 and Fig. S1(a-f))

224 and five of 12 non-attended species (Table 1 and Fig. S1(g-l)), in which at least one

225 individual was detected. Mean *Wolbachia* infection rates were 30.2% in ant-attended

226 species and 3.1% in non-attended species (Table 1). A large variation of *Wolbachia*

227 infection rates was found in ant-attended species (0% in *T. indicus* (Fig. S1b), *T. pappus*

228 (Fig. S1e), and *T*. sp. E (Fig. S1f); 100% in *T*. sp. B (Fig. S1f). The Mantel test on *T.*

229 *fulviabdominalis* and *T. macrotuberculatus* showed that *Wolbachia* infection rates was

230 significantly correlated with distance between collection sites (for *T. fulviabdominalis*,

231 Mantel statistic $r = 0.842$, $P = 0.035$ (Table 1 and Fig. S1b); for *T. macrotuberculatus*,

232 Mantel statistic $r = 0.164$, $P = 0.03$ (Table 1 and Fig. S1d).

233

234 **Phylogenetic independent contrasts**

235 The ML phylogenetic tree based on the combined sequences of *COI* and *18S rRNA* genes

236 showed fully resolved tree topology (Fig. 1). CAIC showed a significant positive

237 correlation between contrasts of *Wolbachia* infection rates and ant association (CAIC, $F_{1, 21}$

238 = 13.7, P = 0.00134, Fig. 2); *Wolbachia* infection rates in *Tuberculatus* aphids were

239 significantly higher in ant-attended species compared to non-attended species.

240

241 ***Wolbachia* supergroups**

242 Because the sequencing for *T. pilosulus* and *T*. sp. D was unsuccesful, only 11

243 *Wolbachia*-positive were analyzed. The results of sequencing showed that each species

244 harboured one haplotype of *Wolbachia* except for *T. macrotuberculatus* (Fig. 3).

245 *Tuberculatus macrotuberculatus* harboured two haplotypes (Fig. 3): one haplotype was

246 found at nine sites (sites 1 to 8 and site 23), the other at site 22. A NJ tree showed that 12

247 haplotypes of *Wolbachia* were classified into four supergroups B, M, N, and O (Fig. 3). The

248 haplotypes of *Wolbachia* in *T. kuricola*, *T. stigmatus*, *T. higuchii* B-type and *T. paiki* were

249 placed into supergroup B. *Wolbachia* in *T. macrotuberculatus* collected from all infected

250 sites except for site 22, *T. quercicola* and *T*. sp. B belonged to supergroup M. *Wolbachia* in

251 *T. macrotuberculatus* collected from the site 22, *T. capitatus*, *T. fulviabdominalis* and *T.*

252 *japonicus* were placed into supergroup N. *Tuberculatus higuchii* A-type harboured

253 *Wolbachia* of supergroup O, which was supported with a high bootstrap value (100%).

254 Twelve DNA sequences of *Wolbachia*'s *16S rRNA* were deposited in the DDBJ and

255 accession numbers are listed in Fig. 3 and Table S3.

256

257 **Discussion**

258 The phylogenetic comparative analysis showed that *Wolbachia* infection rates were higher

259 in aphid species that have mutualistic associations with ants. One possible infection route

260 of *Wolbachia* to aphids could be horizontal transmission between *Wolbachia*-infected

261 parasitoid wasps and aphids. Regardless of whether aphids are attended by ants, aphid

262 colonies are frequently attacked by parasitoid wasps (Brodeur and Rosenheim 2000). Field

263 experiments on some ant-attended aphid species demonstrated that ant-attended colonies

264 attracted more parasitoid wasps compared to ant-excluded colonies (Völkl 1992; Kaneko

265 2002, 2003; Sadeghi-Namaghi and Amiri-Jami 2018). These behaviours of parasitoid wasps

266 are thought to be triggered by visual and chemical cues from aphid colonies attended by

267 ants (Mouratidis et al. 2021). Ant-attended species form dense colonies (Stadler et al. 2003)

268 and disperse less than non-attended species (Oliver et al. 2007; Yao 2010), which could by
269 itself an explanation for a higher *Wolbachia* prevalence. A study using fluorescence in situ
270 hybridization on the parasitoid wasp *Eretmocerus* sp. showed that *Wolbachia* were present
271 in the mouthparts and ovipositors of wasps feeding on *Wolbachia*-infected whitefly *Bemisia*
272 *tabaci* (Ahmed et al. 2015). Thus, the horizontal transmission of endosymbionts via the
273 parasitoids of insects represents a potential pathway. Besides parasitoid wasps, ants are also
274 known to harbour *Wolbachia* (Keller et al. 2001; Shoemaker et al. 2003; Tsutsui et al. 2003;
275 Viljakainen et al. 2008; Frost et al. 2010; Reeves et al. 2020) and thus could be a possible
276 agent to spread *Wolbachia* into aphid populations. In a study of scale insects and their
277 associated groups (ants, wasps, beetles, flies, mites, moths, and thrips), Sanaei et al. (2022)
278 showed that significantly higher *Wolbachia* infection rates in ant-attended scale insects,
279 suggesting a possible horizontal transfer route between ants and scale insects. This study
280 did not aim to identify the *Wolbachia* strains of parasitoid wasps or attending ants. Further
281 studies on *Wolbachia* strains for aphids and their parasitoid wasps or their mutualistic ants
282 are need to elucidate the possible routes by ants.

283 Although the average *Wolbachia* infection rates was higher in ant-attended species
284 (30.2%) than in non-attended species (3.1%), a wide range of variation was found in the
285 infection rates for ant-attended species (0-100%). The difference in realized infection rate
286 can be attributed to ecological or environmental factors affecting the cost-benefit balance of
287 *Wolbachia* infection to hosts (Gavotte et al. 2010; White et al. 2011; Okayama et al. 2016).
288 Higher infection levels across all populations of *T. capitatus* (on average 94.6% from 15
289 sites) and *T*. sp. B (100 % from 4 sites) could be responsible for positive selection
290 favouring benefits from *Wolbachia* infection such as nutrition provision (Hosokawa et al.
291 2010; De Clerck et al. 2015; but see Manzano-Marín 2020) or resistance to parasitoid
292 wasps (Oliver et al. 2003). Hence, it could be possible that *Wolbachia* plays obligate
293 mutualistic roles in *T. capitatus* and *T*. sp. B. On the other hand, for the species with
294 infection rate of between 10 and 52 %, it is difficult to determine whether *Wolbachia*
295 infection is a mutualistic or parasitic interaction with the species. The previous study of
296 seasonal changes in *Wolbachia* density in a population (site 4 in this study) of *T.*
297 *macrotuberculatus* showed that 315 of 316 (99.7%) of the aphids harboured *Wolbachia* and

298 *Wolbachia* density in an individual aphid exhibited no significant fluctuations during the
299 survey period, implying that seasonal deterioration of host plants did not affect *Wolbachia*
300 density, even though host aphids decreased in their body size and embryo numbers (Yao
301 2019). *Wolbachia* of the aphids in this site seems to give a beneficial effect on the
302 nutritional status of aphids during the harsh summer. However, in this study, no
303 *Wolbachia*-infected aphids were found in 13 of 23 collection sites of *T. macrotuberculatus*.
304 Furthermore, there was a significant correlation between geographical distance and
305 difference in infection rates in two species *T. fulviabdominalis* and *T. macrotuberculatus*.
306 This means that there is an isolation-by-distance effect among the collection sites. Indeed, it
307 has been demonstrated that the genetic structure of *T. macrotuberculatus* in Hokkaido
308 populations shows a higher inbreeding coefficient in each subpopulation and less dispersal
309 due to ant attendance (Yao 2010), suggesting that region-specific patterns as to whether
310 *Wolbachia* infection is costly or beneficial could occur in isolated populations. For the
311 species with less than 5 % infection rate, three of four species (*T. japonicus*, *T. paiki*, and *T.*
312 sp. D) are non-attended species and sometimes have been observed with ant-attended

313 species (*T. fulviabdominalis*, *T. macrotuberculatus*, *T. stigmatus* and *T*. sp. B) on the same

314 host plant (*Quercus dentata*). This sympatric host plant use might provide the non-attended

315 species with an opportunistic infection of *Wolbachia*, such as via plant-mediated horizontal

316 transmission (Li et al. 2017).

317 *Wolbachia* haplotypes were clustered into the four supergroups B, M, N, and O. Out of

318 the 11 *Wolbachia*-infected *Tuberculatus* species in the phylogenetic tree, *T. higuchii* A-type

319 fell into supergroup O that has been firstly detected in the white fly *Bemisia tabaci* (Bing et

320 al. 2014) and recently found in the galling aphid species, *Kaburagia rhusicola* and

321 *Schlechtendalia chinensis* (Ren et al. 2020). Detection in the novel host and a monophyletic

322 group with a high bootstrap value (100%) will support existence of supergroup O. Given

323 that supergroup O has so far been found only in China, it could have originated in East Asia

324 and spread into Japan. As *Wolbachia* supergroups have evolved independently, infections

325 by different supergroups presumably represent independent gains of the trait even for two

326 species with the same ant-attendance state, but these are ignored in the current analysis.

327 This would be overcome by the comparison of characteristics of hosts infected by different

328 supergroups and distributed in close distance areas. *Tuberculatus macrotuberculatus*

329 harboured two phylogenetically-distant supergroups of M and N as previously seen in

330 Moreira et al. (2019); the two sites of southern island, site 22 and site 23 (apart from

331 approximately 40 km, Fig. S1d), had the supergroup N and supergroup M, respectively.

332 Comparison between the two populations may help to elucidate the difference of

333 independent gains of *Wolbachia* supergroup involving aphid-ant mutualisms.

334 This study has revealed that the ecological characteristics of aphid hosts have

335 influenced the extent of *Wolbachia* spread in these species. Further studies are needed to

336 clarify what roles *Wolbachia* play in aphids, especially for ant-attended aphid species.

337

338 **Acknowledgments**

339 I thank S. Sugimoto of Plant Protection Station and the members of SEHU for collecting

340 aphids. I thank K. Yoshizawa, T. Kanbe, S. Suzuki, and S. Akimoto of Hokkaido University

341 for providing help during this study. This work was supported by JSPS Kakenhi Grant

342 Numbers 24570016 and 15K07210 for IY.

343

344 **Authors' contributions** IY conceived the study, collected samples, carried out molecular

345 work, analyzed the data and wrote the manuscript.

346 **Availability of data and material** Data are available by request to the author.

347 **Declarations**

348 **Ethics approval** N/A.

349 **Conflicts of interest/Competing interests** The author declares no conflicts of interest or

350 competing interests.

351 **Consent to participate** N/A.

352 **Consent for publication** N/A.

353

354 **References**

355 Ahmed MZ, Li SJ, Xue X, Yin XJ, Ren SX, Jiggins FM, Greeff JM, Qiu BL (2015) The

356 intracellular bacterium *Wolbachia* uses parasitoid wasps as phoretic vectors for

357 efficient horizontal transmission. PLoS Pathog 11:e1004672.

358 https://doi.org/10.1371/journal.ppat.1004672

359 Augustinos AA, Santos-Garcia D, Dionyssopoulou E, Moreira M, Papapanagiotou A,

360 Scarvelakis M, Doudoumis V, Ramos S, Aguiar AF, Borges PAV, Khadem M, Latorre

361 A, Tsiamis G, Bourtzis K (2011) Detection and characterization of *Wolbachia*

362 infections in natural populations of Aphids: Is the hidden diversity fully unraveled?

363 PLoS One 6:e28695. https://doi.org /10.1371/journal.pone.0028695

364 Baldo L, Hotopp JCD, Jolley KA, Bordenstein SR, Biber SA, Choudhury RR, Hayashi C,

365 Maiden MCJ, Tettelin H, Werren JH (2006) Multilocus sequence typing system for the

366 endosymbiont *Wolbachia pipientis*. Appl Environ Microbiol 72:7098-7110.

367 https://doi.org/10.1128/AEM.00731-06

368 Barker SC, Whiting MF, Johnson KP, Murrell A (2003) Phylogeny of lice (Insecta:

369 Phthiraptera) inferred from small subunit rRNA. Zool Scr 32:407–414.

370 https://doi.org/10.1046/j.1463-6409.2003.00120.x

371 Bing XL, Xia WQ, Gui JD, Yan GH, Wang XW, Liu SS (2014) Diversity and evolution of

372 the *Wolbachia* endosymbionts of *Bemisia* (Hemiptera: Aleyrodidae) whiteflies. Ecol

373 Evol 4:2714-2737. https://doi.org/10.1002/ece3.1126

374 Brodeur J, Rosenheim JA (2000) Intraguild interactions in aphid parasitoids. Entomol Exp

375 Appl 97:93-108. https://doi.org/10.1046/j.1570-7458.2000.00720.x

376 Casiraghi M, Anderson TJ, Bandi C, Bazzocchi C, Genchi C (2001) A phylogenetic

377 analysis of filarial nematodes: comparison with the phylogeny of *Wolbachia*

378 endosymbionts. Parasitology 122:93–103. https://doi.org/10.1017/s0031182000007149

379 De Clerck C, Fujiwara A, Joncour P, Léonard S, Félix ML, Francis F, Jijakli MH, Tsuchida

380 T, Massart S (2015) A metagenomic approach from aphid's hemolymph sheds light on

381 the potential roles of co-existing endosymbionts. Microbiome 3:63.

382 https://doi.org/10.1186/s40168-015-0130-5

383 De Clerck C, Tsuchida T, Massart S, Lepoivre P, Francis F, Jijakli MH. 2014. Combination

384 of genomic and proteomic approaches to characterize the symbiotic population of the

385 banana aphid (Hemiptera: Aphididae). Environ Entomol 43:29–36.

386 https://doi.org/10.1603/EN13107

387 Felsenstein J (1985) Phylogenies and the comparative method. Am Nat 125:1–15.

388 https://doi.org/10.1086/284325

389 Fischer M, Shingleton A (2001) Host plant and ants influence the honeydew sugar

390 composition of aphids. Funct Ecol 15:544-550.

391 https://doi.org/10.1046/j.0269-8463.2001.00550.x

392 Fischer CY, Lognay GC, Detrain C, Heil M, Grigorescu A, Sabri A, Thonart P, Haubruge E,

393 Verheggen FJ (2015) Chemoecology 25:223-232.

394 https://doi.org/10.1007/s00049-015-0188-3

395 Flatt T, Weisser W (2000) The effects of mutualistic ants on aphid life history traits.

396 Ecology 81:3522-3529.

397 https://doi.org/10.1890/0012-9658(2000)081[3522:TEOMAO]2.0.CO;2

398 Folmer O, Black M, Hoeh W, Lutz R, Vrijenhoek R (1994) DNA primers for amplification

399 of mitochondrial cytochrome *c* oxidase subunit I from diverse metazoan

400 invertebrates. Mol Marine Biol Biotechnol 3:294–299. PMID: 7881515

401 Frost CL, FernÁndez-MarÍn H, Smith JE, Hughes WOH (2010) Multiple gains and losses

402 of *Wolbachia* symbionts across a tribe of fungus-growing ants. Mol Ecol 19:4077-4085.

https://doi.org/10.1111/j.1365-294X.2010.04764.x

Garland TJr, Harvey PH, Ives AR (1992) Procedures for the analysis of comparative data using phylogenetically independent contrasts. Syst Biol 41:18-32. https://doi.org/10.1093/sysbio/41.1.18

Gavotte L, Mercer DR, Stoeckle JJ, Dobson SL (2010) Costs and benefits of *Wolbachia* infection in immature *Aedes albopictus* depend upon sex and competition level. J Invertebr Pathol 105:341-346. http://dx.doi.org/10.1016/j.jip.2010.08.005

Glowska E, Dragun-Damian A, Dabert M, Gerth M (2015) New *Wolbachia* supergroups detected in quill mites (Acari: Syringophilidae). Infect Genet Evol 30:140-146. https://doi.org/10.1016/j.meegid.2014.12.019

Gómez-Valero L, Soriano-Navarro M, Pérez-Brocal V, Heddi A, Moya A, García-Verdugo JM, Latorre A (2004) Coexistence of *Wolbachia* with *Buchnera aphidicola* and a secondary symbiont in the aphid *Cinara cedri*. J Bacteriol 186:6626-6633. https://doi.org/10.1128/JB.186.19.6626–6633.2004

Harvey PH, Pagel MD (1991) The comparative method in evolutionary biology. Oxford

418 University Press, New York

419 Henry LM, Maiden MCJ, Ferrari J, Godfray HCJ (2015) Insect life history and the

420 evolution of bacterial mutualism. Ecol Lett 18:516-525.

421 https://doi.org/10.1111/ele.12425

422 Hertaeg C, Risse M, Vorburger C, De Moraes CM, Mescher MC (2021) Aphids harbouring

423 different endosymbionts exhibit differences in cuticular hydrocarbon profiles that can

424 be recognized by ant mutualists. Sci Rep 11:1-10

425 https://doi.org/10.1038/s41598-021-98098-2

426 Hilgenboecker K, Hammerstein P, Schlattmann P, Telschow A, Werren JH (2008) How

427 many species are infected with *Wolbachia*? - A statistical analysis of current data.

428 FEMS Microbiol Lett 281:215-220. https://doi.org/10.1111/j.1574-6968.2008.01110.x

429 Hosokawa T, Koga R, Kikuchi Y, Meng XY, Fukatsu T (2010) *Wolbachia* as a

430 bacteriocyte-associated nutritional mutualist. Proc Natl Acad Sci USA 107:769-774.

431 https://doi.org/10.1073/pnas.0911476107

432 Kaneko S (2002) Aphid-attending ants increase the number of emerging adults of the

433 aphid's primary parasitoid and hyperparasitoids by repelling intraguild predators.
434 Entomol Sci 5:131-146. http://dl.ndl.go.jp/info:ndljp/pid/10656505.
435 Kaneko S (2003) Different impacts of two species of aphid-attending ants with different
436 aggressiveness on the number of emerging adults of the aphid's primary parasitoid and
437 hyperparasitoids. Ecol Res 18:199-212.
438 https://doi.org/10.1046/j.1440-1703.2003.00547.x.
439 Katayama N, Suzuki N (2002) Cost and benefit of ant attendance for *Aphis craccivora*
440 (Hemiptera: Aphididae) with reference to aphid colony size. Can Entomol 134:241-249.
441 https://doi.org/10.4039/Ent134241-2
442 Kaur R, Shropshire JD, Cross KL, Leigh B, Mansueto AJ, Stewart V, Bordenstein SR,
443 Bordenstein SR (2021) Living in the endosymbiotic world of *Wolbachia*: A centennial
444 review. Cell Host Microbe 29:879-893. https://doi.org/10.1016/j.chom.2021.03.006
445 Keller L, Liautard C, Reuter M, Brown WD, Sundström L, Chapuisat M (2001) Sex ratio
446 and *Wolbachia* infection in the ant *Formica exsecta*. Heredity 87:227-233.
447 https://doi.org/10.1046/j.1365-2540.2001.00918.x

448 Lefoulon E, Clark T, Borveto F, Perriat-Sanguinet M, Moulia C, Slatko BE, Gavotte L.
449 (2020) Pseudoscorpion *Wolbachia* symbionts: Diversity and evidence for a new
450 supergroup S. BMC Microbiol 20:188. https://doi.org/10.1186/s12866-020-01863-y
451 Leroy PD, Wathelet B, Sabri A, Francis F, Verheggen FJ, Capella Q, Thonart P, Haubruge E
452 (2011) Aphid-host plant interactions: does aphid honeydew exactly reflect the host
453 plant amino acid composition? Arthropod-Plant Interact 5:193-199.
454 https://doi.org/10.1007/s11829-011-9128-5
455 Li SJ, Ahmed MZ, Lv N, Shi PQ, Wang XM, Huang JL, Qiu BL (2017) Plant-mediated
456 horizontal transmission of *Wolbachia* between whiteflies. ISME J
457 11:1019-1028. http://dx.doi.org/10.1038/ismej.2016.164
458 Mantel N (1967) The detection of disease clustering and a generalized regression
459 approach. Cancer Res 27:209–220
460 Manzano-Marín A (2020) No evidence for *Wolbachia* as a nutritional co-obligate
461 endosymbiont in the aphid *Pentalonia nigronervosa*. Microbiome 8:72.
462 https://doi.org/10.1186/s40168-020-00865-2

463 Moreira M, Aguiar AMF, Bourtzis K, Latorre A, Khadem M (2019) *Wolbachia*
464 (Alphaproteobacteria: Rickettsiales) infections in isolated aphid populations from
465 oceanic islands of the azores archipelago: Revisiting the supergroups M and N.
466 Environ Entomol 48:326-334. https://doi.org/10.1093/ee/nvy189
467 Mouratidis A, Vacas S, Herrero J, Navarro-Llopis V, Dicke M, Tena A (2021) Parasitic
468 wasps avoid ant-protected hemipteran hosts via the detection of ant cuticular
469 hydrocarbons. Proc R Soc London Ser B Biol Sci 288:20201684.
470 https://doi.org/10.1098/rspb.2020.1684
471 Okayama K, Katsuki M, Sumida Y, Okada K (2016) Costs and benefits of symbiosis
472 between a bean beetle and *Wolbachia*. Anim Behav 119:19-26.
473 https://doi.org/10.1016/j.anbehav.2016.07.004
474 Oksanen J, Blanchet FG, Kindt R, Legendre P, Minchin PR, O'Hara RB et al. (2012) vegan:
475 Community Ecology Package 2012.
476 Oliver KM, Russell JA, Moran NA, Hunter MS (2003) Facultative bacterial symbionts in
477 aphids confer resistance to parasitic wasps. Proc Natl Acad Sci USA 100:1803-1807.

http://www.pnas.org/content/100/4/1803.short.

Oliver TH, Mashanova A, Leather SR, Cook JM, Jansen VAA (2007) Ant semiochemicals limit apterous aphid dispersal. Proc R Soc London Ser B Biol Sci 274:3127-3131. https://doi.org/10.1098/rspb.2007.1251

Paradis E, Schliep K (2019) ape 5.0: an environment for modern phylogenetics and evolutionary analyses in R. Bioinformatics 35:526-528. https://doi.org/10.1093/bioinformatics/bty633

Posada D, Crandall KA (1998) Modeltest: testing the model of DNA substitution. Bioinformatics 14:817–818. https://doi.org/10.1093/ bioinformatics/14.9.817

Purvis A, Rambaut A (1995) Comparative analysis by independent contrasts (CAIC): an Apple Macintosh application for analysing comparative data. Bioinformatics 11:247-251. https://doi.org/10.1093/bioinformatics/11.3.247

Quednau FW (1999) Atlas of the drepanosiphine aphids of the world. Part I: Panaphidini Oestlund, 1922-Myzocallidina Börner, 1942 (1930) (Hemiptera: Aphididae: Calaphidinae). Contributions of the American Entomological Institute, 31(1).

493 R Core Team (2021). R: A language and environment for statistical computing. R
494 Foundation for Statistical Computing, Vienna, Austria. https://www.R-project.org/
495 Reeves DD, Price SL, Ramalho MO, Moreau CS (2020) The Diversity and distribution of
496 *Wolbachia*, *Rhizobiales*, and *Ophiocordyceps* within the widespread Neotropical turtle
497 ant, *Cephalotes atratus* (Hymenoptera: Formicidae). Neotrop Entomol 49:52-60.
498 https://doi.org/10.1007/s13744-019-00735-z
499 Ren W, Wei H, Yang Y, Shao S, Wu H, Chen X, Yang Z (2020) Molecular detection and
500 phylogenetic analyses of *Wolbachia* in natural populations of nine galling Aphid
501 species. Sci Rep 10:12025. https://doi.org/10.1038/s41598-020-68925-z
502 Renyard A, Gries R, Lee J, Chalissery JM, Damin S, Britton R, Gries G (2021) All sugars
503 ain't sweet: selection of particular mono-, di- and trisaccharides by western carpenter
504 ants and European fire ants. R Soc Open Sci 8:210804.
505 https://doi.org/10.1098/rsos.210804
506 Ros VID, Fleming VM, Feil EJ, Breeuwer AJ (2009) How diverse is the genus *Wolbachia*?
507 Multiple-gene sequencing reveals a putatively new *Wolbachia* supergroup recovered

508 from spider mites (Acari: Tetranychidae). Appl Environ Microbiol 75:1036-1043.
509 https://doi.org/10.1128/AEM.01109-08
510 Sadeghi-Namaghi H, Amiri-Jami A (2018) Success of aphid parasitoids and their hosts
511 varies with ant attendance: A field study. Entomol Sci 21:406-411.
512 https://doi.org/10.1111/ens.12319
513 Sanaei E, Charlat S, Engelstädter J (2021) *Wolbachia* host shifts: routes, mechanisms,
514 constraints and evolutionary consequences. Biol Rev 96:433-453.
515 https://doi.org/10.1111/brv.12663
516 Sanaei E, Lin YP, Cook LG, Engelstädter J (2022) *Wolbachia* in scale insects: a distinct
517 pattern of infection frequencies and potential transfer routes via ant associates. Environ
518 Microbiol 24:1326-1339. https://doi.org/10.1111/1462-2920.15833
519 Scarborough C, Ferrari J, Godfray H (2005) Aphid protected from pathogen by
520 endosymbiont. Science 310:1781.
521 https://www.science.org/doi/10.1126/science.1120180
522 Shoemaker DD, Ahrens M, Sheill L, Mescher M, Keller L, Ross KG (2003) Distribution

523 and prevalence of *Wolbachia* infections in native populations of the fire ant *Solenopsis*
524 *invicta* (Hymenoptera: Formicidae). Environ Entomol 32:1329–1336.
525 https://doi.org/10.1603/0046-225X-32.6.1329
526 Simões PM, Mialdea G, Reiss D, Sagot MF, Charlat S (2011) *Wolbachia* detection: an
527 assessment of standard PCR Protocols. Mol Ecol Res 11:565-572.
528 https://doi.org/10.1111/j.1755-0998.2010.02955.x
529 Stadler B, Dixon AFG (1998) Costs of ant attendance for aphids. J Anim Ecol 67:454-459.
530 https://doi.org/10.1046/j.1365-2656.1998.00209.x
531 Stadler B, Dixon AFG (2005) Ecology and evolution of aphid-ant interactions. Annu Rev
532 Ecol Evol Syst 36:345-372. https://doi.org/10.1146/annurev.ecolsys.36.091704.175531
533 Stadler B, Kindlmann P, Šmilauer P, Fiedler K (2003) A comparative analysis of
534 morphological and ecological characters of European aphids and lycaenids in relation
535 to ant attendance. Oecologia 135:422-430. https://doi.org/10.1007/s00442-003-1193-8
536 Swofford DL (2002) PAUP*. Phylogenetic analysis using parsimony (and other methods).
537 Version 4. Sinauer Associates, Sunderland, Massachusetts.

538 Thompson JD, Higgins DG, Gibson TJ (1994) CLUSTAL W: improving the sensitivity of
539 progressive multiple sequence alignment through sequence weighting, position-specific
540 gap penalties and weight matrix choice. Nucleic Acids Res 22:4673-4680.
541 https://doi.org/10.1093/nar/22.22.4673
542 Tsutsui ND, Kauppinen SN, Oyafuso AF, Grosberg RK (2003) The distribution and
543 evolutionary history of *Wolbachia* infection in native and introduced populations of the
544 invasive argentine ant (*Linepithema humile*). Mol Ecol
545 12:3057-3068. https://doi.org/10.1046/j.1365-294X.2003.01979.x
546 Viljakainen L, Reuter M, Pamilo P (2008) *Wolbachia* transmission dynamics in *Formica*
547 wood ants. BMC Evol Biol 8:55. https://doi.org/10.1186/1471-2148-8-55
548 Vorburger C, Gehrer P, Rodriguez P (2010) A strain of the bacterial symbiont *Regiella*
549 *insecticola* protects aphids against parasitoids. Biol Lett 6:109-111.
550 https://doi.org/10.1098/rsbl.2009.0642
551 Völkl W (1992) Aphids or their parasitoids: who actually benefits from ant-attendance? J
552 Anim Ecol 61:273-281. https://doi.org/10.2307/5320

553 Wang Z, Shen ZR, Song Y, Liu HY, Li ZX (2009) Distribution and diversity of *Wolbachia*
554 in different populations of the wheat aphid *Sitobion miscanthi* (Hemiptera: Aphididae)
555 in China. Eur J Entomol 106:49-55. https://doi.org/10.14411/eje.2009.007
556 Way MJ (1963) Mutualism between ants and honeydew-producing Homoptera. Annu Rev
557 Entomol 8:307-344. https://doi.org/10.1146/annurev.en.08.010163.001515
558 Weinert L, Araujo-Jnr EV, Ahmed MZ, Welch JJ (2015) The incidence of bacterial
559 endosymbionts in terrestrial arthropods. Proc R Soc London Ser B Biol Sci
560 282:20150249. http://dx.doi.org/10.1098/rspb.2015.0249
561 Werren JH, Baldo L, Clark ME (2008) *Wolbachia*: master manipulators of invertebrate
562 biology. Nat Rev Microbiol 6:741-751. https://doi.org/10.1038/nrmicro1969.
563 Werren JH, Windsor DM (2000) *Wolbachia* infection frequencies in insects: evidence of a
564 global equilibrium? Proc R Soc London Ser B Biol Sci 267:1277-1285.
565 https://doi.org/10.1098/rspb.2000.1139
566 White JA, Kelly SE, Cockburn SN, Perlman SJ, Hunter MS (2011) Endosymbiont costs and
567 benefits in a parasitoid infected with both *Wolbachia* and *Cardinium*. Heredity

568 106:585-591. https://doi.org/10.1038/hdy.2010.89

569 Yao I (2010) Contrasting patterns of genetic structure and dispersal ability in ant-attended

570 and non-attended *Tuberculatus* aphids. Biol Lett 6:282-286.

571 https://doi.org/10.1098/rsbl.2009.0781.

572 Yao I (2011) Phylogenetic comparative methods reveal higher wing loading in ant-attended

573 *Tuberculatus* aphids (Hemiptera: Aphididae). Can Entomol 143:35-43.

574 https://doi.org/10.4039/n10-050

575 Yao I (2014) Costs and constraints in aphid-ant mutualism. Ecol Res 29:383-391.

576 https://doi.org/10.1007/s11284-014-1151-4

577 Yao I (2019) Seasonal changes in the density of the symbionts *Buchnera* and *Wolbachia* of

578 the aphid *Tuberculatus macrotuberculatus* on *Quercus dentata*. Entomol Exp Appl

579 167:261-268. https://doi.org/10.1111/eea.12743

580 Yao I and Akimoto SI (2001) Ant attendance changes the sugar composition of the

581 honeydew of the drepanosiphid aphid *Tuberculatus quercicola*. Oecologia 128:36-43.

582 http://link.springer.com/10.1007/s004420100633

583 Yao I and Akimoto SI (2002) Flexibility in the composition and concentration of amino
584 acids in honeydew of the drepanosiphid aphid *Tuberculatus quercicola*. Ecological
585 Entomology 27:745-752. http://doi.wiley.com/10.1046/j.1365-2311.2002.00455.x
586 Yao I, Shibao H, Akimoto SI (2000) Costs and benefits of ant attendance to the
587 drepanosiphid aphid *Tuberculatus quercicola*. Oikos 89:3-10.
588 https://doi.org/10.1034/j.1600-0706.2000.890101.x
589 Zhou W, Rousset F, O'Neill S (1998) Phylogeny and PCR-based classification of
590 *Wolbachia* strains using *wsp* gene sequences. Proc R Soc London Ser B Biol Sci
591 265:509-515. https://doi.org/10.1098/rspb.1998.0324
592

593 Figure legends

594 Fig. 1. Maximum likelihood phylogenetic tree based on the combined sequences of *COI*
595 and *18S rRNA* genes.
596 (Note) Operational taxonomic units (OTUs) shown in bold font indicate ant-attended
597 species. The numbers on the branches of phylogenetic tree show bootstrap values (>50%).

598 Full terms of abbreviations are provided in Table 1.

599

600 Fig. 2. Correlation between contrasts of ant association and *Wolbachia* infection rates.

601 (Note) Contrastant and contrastwol represent independent contrasts that were calculated

602 based on the status of ant association and *Wolbachia* infection rates at each node (n = 22) in

603 the ML phylogenetic tree.

604

605 Fig. 3. Bootstrap 50 % majority-rule consensus tree inferred by neighbor-joining

606 (NJ) analysis for 16 *Wolbachia* supergroups

607 (Note) The labels of operational taxonomic units (OTUs) mean *Wolbachia* sp. (indicated by

608 w) and its host species. Thick vertical lines with alphabets indicate the clades of *Wolbachia*

609 supergroups. See also Table S3. Bootstrap values of more than 50% were shown on

610 branches. Full terms of abbreviations are provided in Table 1.

611

Table 1. *Tuberculatus* aphid species used in the study and *Wolbachia* infection rate

Ant-attended	Collection sites	*N*	*wol*$^{+}$	Infection rate (%)	Mantel statistic r	*P*	Abbreviation	*COI*	*18S rRNA*
T. capitatus	15	56	53	94.6	0.032	0.196	capi	AB592769	LC654240
T. fulviabdominalis	8	55	12	21.8	0.842	**0.035**	fulvi	AB592755	LC654241
T. indicus	11	53	0	0.0	-	-	ind	AB592759	LC654242
T. kuricola	10	54	15	27.8	-0.093	0.719	kuri	AB592750	LC654243
T. macrotuberculatus	23	54	28	51.9	0.164	**0.030**	mt	AB592752	LC654244
T. pappus	1	10	0	0.0	-	-	pap	AB861442	LC654245
T. pilosulus	16	79	1	1.3	0.596	0.059	pilosulus	AB592758	LC654246
T. quercicola	11	54	8	14.8	-0.105	0.551	que	AB592754	LC654247
T. stigmatus	15	56	11	19.6	-0.039	0.415	sti	AB592760	LC654248
T. sp. B	4	31	31	100.0	-	-	spB	AB592753	LC654249
T. sp. E	1	9	0	0.0	-	-	spE	AB861448	LC654250
Average	10.5	46.5	14.5	30.2					
Non-attended									
T. higuchii A-type	14	71	8	11.3	-0.100	0.560	higa	AB592762	LC654251
T. higuchii B-type	7	42	8	19.0	-0.089	0.657	higb	AB592764	LC654252
T. japonicus	7	59	1	1.7	0.814	0.143	japo	AB592756	LC654253
T. kashiwae A-type	5	39	0	0.0	-	-	kasa	AB592765	LC654254
*T. kashiwa*e B-type	4	47	0	0.0	-	-	kasb	AB592766	LC654255
T. paiki	18	51	1	2.0	0.086	0.221	paiki	AB592768	LC654256
T. pilosus	11	52	0	0.0	-	-	pilosus	AB592751	LC654257
T. querciformosanus	9	52	0	0.0	-	-	qfor	AB592761	LC654258
T. yokoyamai	3	18	0	0.0	-	-	yoko	AB592767	LC654259
T. sp.C	1	41	0	0.0	-	-	spC	AB592757	LC654260
T. sp.D	1	29	1	3.4	-	-	spD	AB592763	LC654261
T. sp.F	1	4	0	0.0	-	-	spF	AB861457	LC654262

Average	6.8	42.1	1.6	3.1

(Note): Collection sites represent the number of collection sites for aphids (see Table S2 for details). *N* and *wol*$^{+}$ mean the numbers of aphid individuals amplified with barcoding region primers and those with *Wolbachia* specific primers. Infection rate (%) was defined by the per cent of *wol*$^{+}$ divided by *N*. Except for exhaustive infection of *T*. sp. B, a Mantel test was applied to the species that were collected from more than a single site. Statistics of Mantel test, r, and *P*-values are given. The bold font shows a significant difference below 0.05 of *P*-values. Abbreviated names were used in Table 3, all figures and supplementary files. Accession numbers of *COI* and *18S rRNA* genes from DDBJ were used to create phylogenetic trees of the aphids.

Table 2. Primer set used in the small-scale experiment of *Wolbachia* detection and the amplification of *18S rRNA* gene in host aphids

Primer name	Primer sequence (5' to 3')	Product size (bp)	References
WspecF (16S-2f)	CATACCTATTCGAAGGGATAG	438	Werren and Windsor (2000)
WspecR (16S-2r)	AGCTTCGAGTGAAACCAATTC		
16SWolbF (16S-3f)	GAAGATAATGACGGTACTCAC	1014	Casiraghi et al. (2001)
16SwolbR3 (16S-3r)[*1]	GTCACTGATCCCACTTTAAATAAC		
553F_W (16S-6f)	ATACGGAGAGGGCTAGCGTTA	781	Simões et al. (2011)
1334R_W (16S-6r)	CTTCATRYACTCGAGTTGCWGAGT		
16SWup	GCCTAACACATGCAAGTCGAA	1400	Gomez-Valero et al. (2004)
16SWlo	AGCTTCGAGTGAAACCAATTCCC		
groEL-F (Wol)	CAACRGTRGSRRYAACTGCDGG	550	Ros et al. (2009)
groEL-R (Wol)	GATADCCRCGRTCAAAYTGC		
wsp81F	TGGTCCAATAAGTGATGAAGAAA	610	Zhou et al. (1998)
wsp691R	AAAAATTAAACGCTACTCCA		
FbpA_F1	GCTGCTCCRCTTGGYWTGAT	509	Baldo et al. (2006)
FbpA_R1	CCRCCAGARAAAAYYACTATTC		
16SWolbF (16S-3f)	GAAGATAATGACGGTACTCAC	972	This study
WspecR (16S-2r)	AGCTTCGAGTGAAACCAATTC		
LCO1490	GGTCAACAAATCATAAAGATATTGG	708	Folmer et al. (1994)
HCO2198	TAAACTTCAGGGTGACCAAAAAATCA		
Ns1	GTAGTCATATGCTTGTCT C	670 (approximately)	Barker et al. (2003)
Ns2a	CGCGGCTGCTGGCACCAGACTTGC		

(Note) *1. Reverse primer was not used in this study.

Table 3. Result of the small-scale experiment using seven pairs of primers

Primer combination	capi	fulvi	ind	kuri	mt	pap	pilosulus	que	sti	spB	spE
*16S-2f*16S-2r*	+	–	–	–	+	–	–	+	+	+	–
*16S-6f*16S-6r*	+	+–	–	–	+	–	+–	+	+–	+	–
*16SWup*16SWlo*	+	–	–	–	+	–	–	+	–	+	–
*groEL-F*groEL-R*	+	–	–	–	+	–	–	+	–	+	–
*FbpA_F1*FbpA_R1*	+	–	–	–	+	–	–	+	–	+	–
*wsp81F*wsp691R*	+	–	–	–	–	–	–	–	–	–	–
*16S-3f*16S-2r*	+	–	–	–	+	–	–	+	+	+	–

Primer combination	higa	higb	japo	kasa	kasb	paiki	pilosus	qfor	yoko	spC	spD	spF
*16S-2f*16S-2r*	–	+	–	–	–	–	–	–	–	–	–	+–
*16S-6f*16S-6r*	+–	+	–	–	–	–	–	–	+–	–	+–	–
*16SWup*16SWlo*	–	–	+	–	–	–	–	–	–	–	–	–
*groEL-F*groEL-R*	–	–	+	–	–	–	–	–	–	–	–	–
*FbpA_F1*FbpA_R1*	–	+	–	–	–	–	–	–	–	–	–	–
*wsp81F*wsp691R*	–	–	+	–	–	–	–	–	–	–	–	–
*16S-3f*16S-2r*	–	+	+	–	–	–	–	–	–	–	–	–

(Note) Symbols + and – indicate that a clear band appeared and no band appeared, respectively. Symbols +– mean that a faint band appeared in 10 μL of PCR reaction volume, but disappeared when rechecked with PCR in 20 μL volume. Full terms of abbreviations are provided in Table 1.

Fig. 1

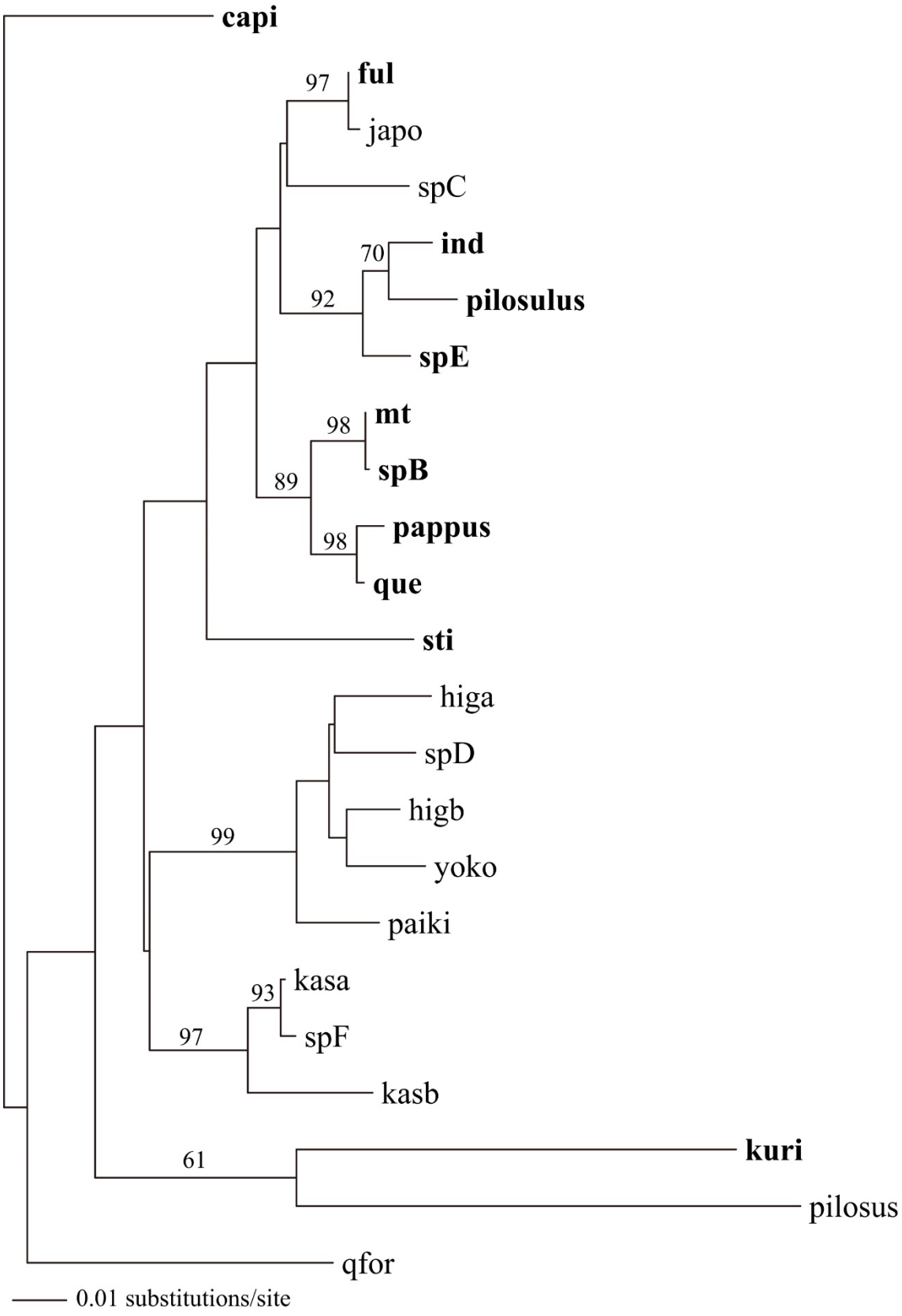

capi
97
ful
japo
spC
70
ind
92
pilosulus
spE
98
mt
89
spB
98
pappus
que
sti
higa
spD
higb
99
yoko
paiki
93
kasa
97
spF
kasb
61
kuri
pilosus
qfor
0.01 substitutions/site

Fig. 2

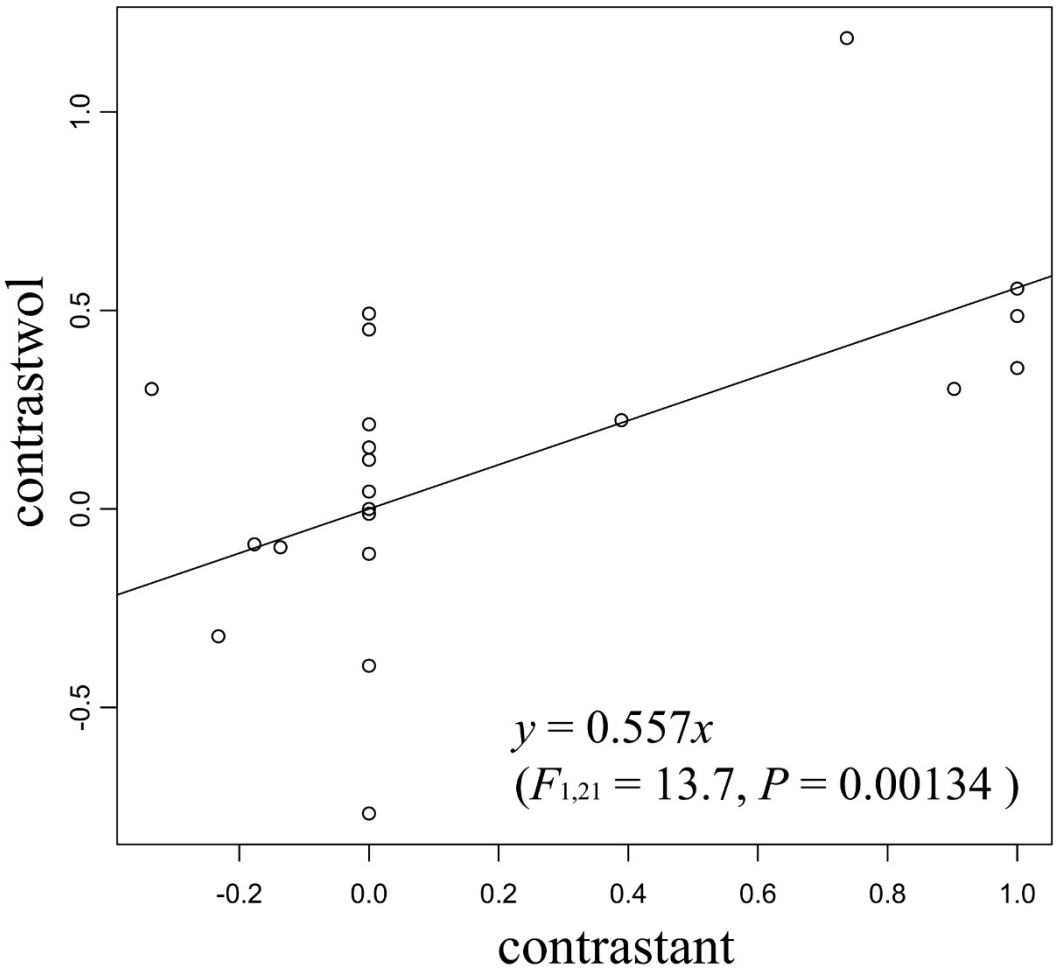
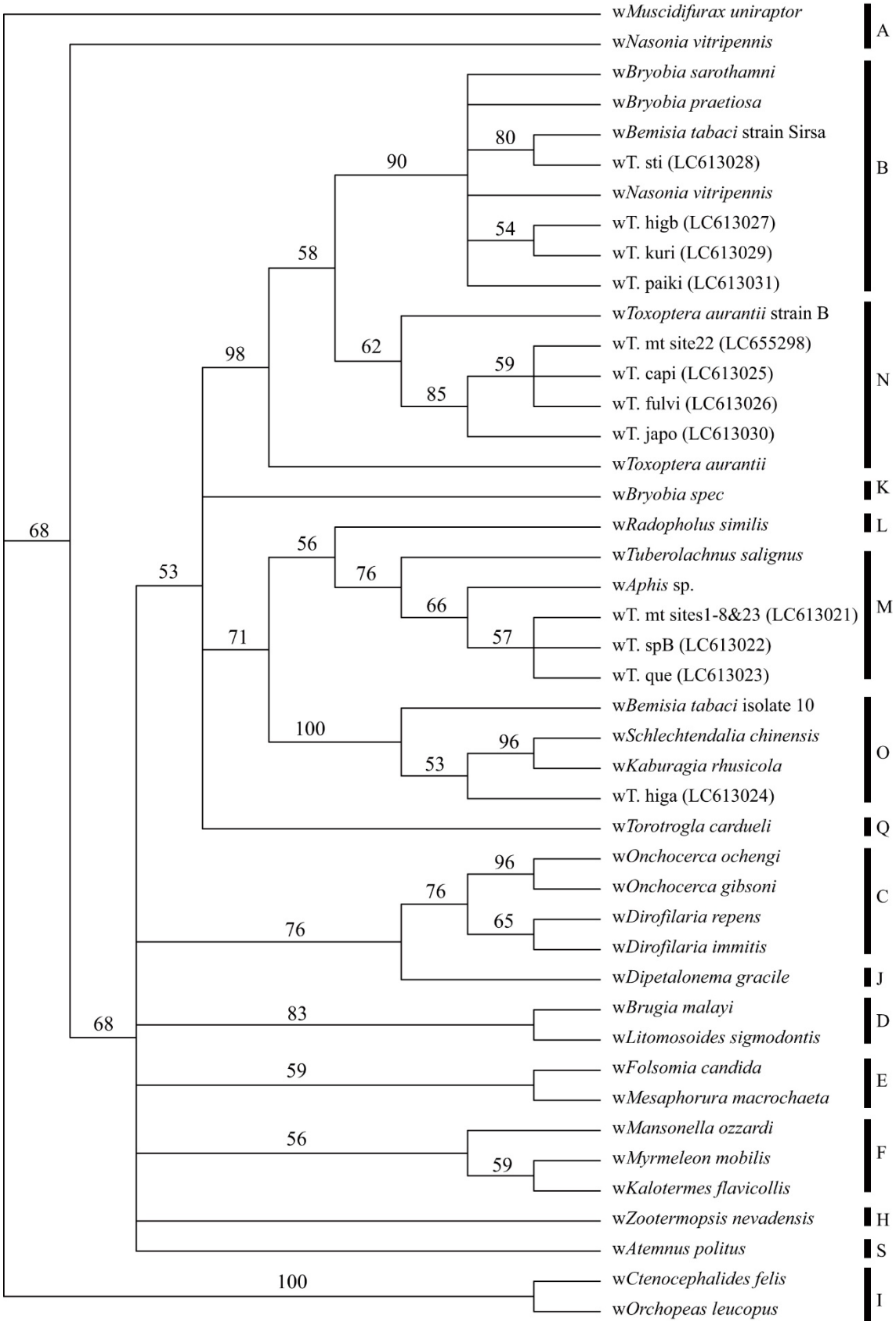

contrastwol
contrastant
1.0
0.5
0.0
-0.5
-0.2
0.0
0.2
0.4
0.6
0.8
1.0
$y = 0.557x$
($F_{1,21} = 13.7$, $P = 0.00134$)

Fig. 3


wMuscidifurax uniraptor
wNasonia vitripennis
wBryobia sarothamni
wBryobia praetiosa
wBemisia tabaci strain Sirsa
wT. sti (LC613028)
wNasonia vitripennis
wT. higb (LC613027)
wT. kuri (LC613029)
wT. paiki (LC613031)
wToxoptera aurantii strain B
wT. mt site22 (LC655298)
wT. capi (LC613025)
wT. fulvi (LC613026)
wT. japo (LC613030)
wToxoptera aurantii
wBryobia spec
wRadopholus similis
wTuberolachnus salignus
wAphis sp.
wT. mt sites1-8&23 (LC613021)
wT. spB (LC613022)
wT. que (LC613023)
wBemisia tabaci isolate 10
wSchlechtendalia chinensis
wKaburagia rhusicola
wT. higa (LC613024)
wTorotrogla cardueli
wOnchocerca ochengi
wOnchocerca gibsoni
wDirofilaria repens
wDirofilaria immitis
wDipetalonema gracile
wBrugia malayi
wLitomosoides sigmodontis
wFolsomia candida
wMesaphorura macrochaeta
wMansonella ozzardi
wMyrmeleon mobilis
wKalotermes flavicollis
wZootermopsis nevadensis
wAtemnus politus
wCtenocephalides felis
wOrchopeas leucopus
A
B
N
K
L
M
O
Q
C
J
D
E
F
H
S
I
80
90
54
58
62
59
85
98
68
56
76
66
57
71
100
96
53
96
76
65
76
68
83
59
56
59
100