Cryptic diversity of the subfamily Calaphidinae (Hemiptera: Aphididae) revealed by comprehensive DNA barcoding

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\textbf{Abstract}

Aphids are a species rich group comprising many important pests. However, species identification can be very difficult for aphids due to their morphological ambiguity. DNA barcoding has been widely adopted for rapid and reliable species identification as well as cryptic species detection. In this study, we investigated cryptic diversity in the subfamily Calaphidinae (Hemiptera: Aphididae) based on 899 sequences of cytochrome c oxidase I (COI) for 115 morphospecies (78 species collected in this study and sequences of 73 species downloaded from Genbank). Among these 115 morphospecies, DNA barcoding results of 90 (78.3\%) species were identical to results of morphological identification. However, 25 (21.7\%) morphospecies showed discrepancies between DNA barcoding and traditional taxonomy. Among these 25 discordances, a total of 15 cryptic species were identified from 12 morphospecies. We also found three morphologically distinct species pairs that sharing DNA barcoding. Based on molecular operational taxonomic unit (MOTU) estimation, we discussed on species delimitation threshold value for these taxa. Our findings confirm that Calaphidinae has high cryptic diversity even though aphids are relatively well-studied.

\textbf{Introduction}

Detecting cryptic species is essential for precise species diversity estimation [1]. With the availability of DNA barcoding methods, recognition of cryptic species has been increased over the past decades [1]. To date, DNA sequences of approximately 10\% of all described species (ca. 160,000) are deposited in open access databases such as Barcode of Life Data Systems (BOLD) and the National Center for Biotechnology Information (NCBI) [2–3]. More recently, some researchers have suggested that DNA-sequenced-based taxonomic description alone is not
only enough for species identification but also can be used as an alternative to classical taxon-
omy [4]. Indeed, DNA barcoding has been contributed to resolve morphological ambiguity in
various taxa [5–9] with identification accuracy rate of 97% [10–13]. Therefore, DNA barcoding
enables the detection of cryptic species and reassessment of species diversity.

Aphids are small and soft-bodied plant sap sucking insects. Over 5,000 of aphid species
have been described worldwide [14], including many important pests in agriculture, forestry,
and species of quarantine importance [15–19]. However, the lack of taxonomically informative
characters and morphological plasticity of aphids make species recognition challenging [20].
Notably, in the complex life cycle of aphids, multiple morphs within one species can occur due
to seasonal changes [21]. Numerous biotic factors such as host-plant relationships [22–23],
natural enemies [24], ant attendance [25], maternal effects [26], endosymbionts [27], and
infectious microorganisms [28] can affect intraspecific plasticity of aphids. Various abiotic fac-
tors, such as climate, temperature and photoperiod [29] also have effects on their intraspecific
plasticity. Conversely, extremely similar or even morphologically indistinguishable species can
occur among aphids [30–32]. The utility of DNA barcoding for aphid species identification
has been demonstrated at family level [7, 13, 30]. It is also useful for some subfamilies such as
Eriosomatinae [33], Greenideinae [34], and Lachninae [35]. According to these studies, DNA
barcoding can provide rapid and reliable identification results for aphids. However, no assess-
ment has been attempted for the subfamily Calaphidinae.

The subfamily Calaphidinae (Hemiptera: Aphididae) is the second largest subfamily in fam-
ily Aphididae. About 398 valid species belonging to 59 genera have been described in the
world [14, 36–37]. Most calaphidine aphids feed on woody angiosperms belonging to 16 plant
families such as Betulaceae, Fagaceae, Juglandaceae, Lythraceae, Myricaceae, and Ulmaceae
although some species feed on herbaceous plants belonging to Fabaceae and Poaceae [37–38].
Many aphid species belonging to this subfamily are economically important pests, causing
injury and transmitting viral diseases to cultivated plants such as leguminous crops, fruit, and
landscape trees [39–40]. For example, Therioaphis trifolii (Monell, 1882), Melanocallis carefo-
liae (Davis, 1910) and Monellia caryella (Fitch, 1855) are notorious aphid pest have caused
large agricultural economic losses [41–43]. Some aphid pest, such as Sarucallis kahawaluok-
lani (Kirkaldy, 1907), Shivaphis celti Das, 1918 and Tinocallis spp. have been dispersed from
their geographic origins to different continents [18, 44–46]. However, assessing calaphidine
species can be difficult and time consuming since their considerable morphological variation
based on seasonal changes and various biotic factors [25, 37]. The application of DNA barcod-
ing would assist in rapid and accurate identification of species in this subfamily. It can also aid
the detection of cryptic diversity.

In this study, we provided the first comprehensive assessment of DNA barcodes for the sub-
family Calaphidinae. A total of 501 Cytochrome oxidase I (COI) sequences of 78 morphospecies
collected in Korea and other countries from 2001 to 2015 were analyzed. The objectives of this
study were i) to clarify delimiting species boundaries in morphologically ambiguous taxa, ii) to
test the effectiveness of DNA barcoding in this taxa, and ultimately iii) to detect hidden species
diversity.

Materials and methods

Ethics statement

No permission was required for sampling at the sites studied. No endangered or protected spe-
cies are included in this study.
Taxon sampling

A total of 501 aphid individuals of 78 species were collected in Asia: Korea (382 specimens of 52 species), China (20 specimens of 8 species), Japan (23 specimens of 8 species) and Laos (2 specimens of 1 species); Europe: Czech Republic (15 specimens of 7 species), Poland (16 specimens of 7 species) and UK (5 specimens of 1 species); North America: USA (29 specimens of 14 species) and Oceania: New Zealand (9 specimens of 4 species) from 2001 to 2015 (S1 Table). Each specimen was preserved in 95–99% ethanol at -20˚C for genomic DNA extraction.

Species identification

501 individuals were mounted in Canada balsam following the method of Blackman & Eastop [16] and Martin [47]. Measurements for each specimen were taken from digital images by using image analysis software (Active measure ver. 3.0.3 from Mitani Co. Ltd, Japan). Digital images were taken by a digital camera attached to a microscope (Leica 400B, Leica Microsystems, Germany). All slide specimens were deposited in the College of Agriculture and Life sciences, Seoul National University (CALS SNU), the Republic of Korea.

DNA extraction and DNA barcoding

Genomic DNA was extracted from each sample selected from each colony by using the DNeasy Blood & Tissue kit (Qiagen, Dusseldorf, Germany) according to the modified manufacturer’s protocols. To confirm morphological features, we used a nondestructive method: each whole-bodied specimen was put into a mixture of 90μl of ATL buffer and 10μl of proteinase K incubated without pulverization. After 24 h incubation, 90μl AL buffer was added and incubated for another 10 min. The solution was gently pipetted into a mini spin column leaving the cuticle of the specimen which was slide mounted.

A 658 bp of COI gene region, generally called as ‘barcoding region’ was amplified using a universal primer set: LCO1490 5’–GGTCAACAAATCATAAAGATATTGG–3’ and HCO2198 5’–TAAACTTCAGGGTGACCAAATCA–3’ [48]. Polymerase chain reaction (PCR) was conducted with AccuPower PCR PreMix (Bioneer, Daejeon, Korea) in 20 ml reaction mixtures under the following conditions: initial denaturation at 94˚C for 3 min; followed by 35 cycles at 94˚C for 30s, an annealing temperature of 45.2˚C for 30s, an extension at 72˚C for 1min; and the final extension at 72˚C for 5min. All PCR products were assessed 1.5% agarose gel electrophoresis. Successfully amplified samples were purified using a QIAquick PCR purification kit (Qiagen, Inc.), and then sequenced directly using an automated sequencer (ABI PrismH 3730 XL DNA Analyzer) at Macrogen Inc. (Seoul, Korea).

Molecular analyses

All sequences to be analyzed were initially assembled and examined using Seqman pro ver. 7.1.0 (DNA star, Inc., Madison, Wisconsin, USA). Poor quality sequences with ambiguous peaks were removed. We used the molecular identification criteria of putative orthologues and paralogues according to Moulton et al. [49] and Fontaneto et al. [50], to prevent misleading by nuclear mitochondrial pseudogenes (Numts) and heteroplasmy. A total of 501 COI sequences of 78 species including previously unknown sequences of 42 species were newly generated for the molecular analyses. Additionally, 398 COI sequences of 73 species were downloaded from Genbank using keyword ‘COI’ and ‘Calaphidinae’ (S2 Table). As a result, the final dataset consisted of 899 sequences of 115 species (S1 and S2 Tables).
These data was aligned using online utility MAFFT ver. 7 alignment package [51] and MEGA 6 [52]. In this step, we removed uncertain anterior and posterior regions were removed. Finally, ≥ 546 bp was used for analyses. For the aligned dataset, a neighbor-joining analysis was conducted using MEGA 6 based on Kimura-2-Parameter (K2P) model [53], the best for species level analysis, particularly for those with low distances [54]. Intra- and inter-specific distances in different taxonomic levels were calculated using pairwise distance method based on the K2P model [53] using MEGA 6.

To infer species delimitation criteria based on a partial COI gene in this subfamily, we performed molecular operational taxonomic units (MOTUs) estimation by using two effective tools to delimit molecular species. First, Automatic Barcode Gap Discovery (ABGD) analysis was conducted to automatically delimit sequences into hypothetical molecular species [55] (http://wwwabi.snv.jussieu.fr/public/abgd) by contrasting inter- and intra-specific distances. Standard settings were used with two values of relative gap width ($X = 1$ and $X = 1.5$) based on Kimura K80 model. Additionally, Bayesian Poisson Tree Processes (bPTP) analysis as implemented on the Exelixis Lab web-server (http://species.h-its.org/ptp/) was performed. This method delimits species based on the phylogenetic species concept [56]. Compared to the generalized mixed Yule coalescent (GMYC) model, bPTP model is a more robust and simpler method [57]. The required rooted phylogenetic input tree was drawn using RAxML [58] with GTR+G+I substitution model.

**Results**

**Genetic variation of morphospecies**

A total of 501 COI sequences (≥ 546 bp) from 78 morphospecies belonging to 36 genera of four subtribes, Calaphidina, Monaphidina, Myzocallidina and Panaphidina were newly generated in this study (S1 Table). All sequences are deposited in Genbank (KY306805-KY307305). Results of genetic divergence at different taxonomic levels are summarized in Table 1.

The overall mean distance was 13.2% for the final dataset of 899 sequences of 115 species. The mean interspecific distance ranged from 4.19% to 13.3% at genus level. The mean genetic distance between genera ranged from 11.92% to 14.51% at subtribe level.

Intraspecific genetic distance was calculated for the 100 of 115 morphospecies. For the remaining 15 species, intraspecific distances could not be calculated because there was only one individual representing each species. The mean intraspecific distance ranged from 0 to 6.0% in each species. Among these 100 morphospecies, 78 species showed very low to moderate genetic divergences (below 1.5%, Fig 1). Another 4 species showed ambiguous intraspecific distances ranging from 1.7% to 1.9% (Fig 1). However, the remaining 18 species showed relatively high intraspecific distances ranging from 2.9% to 16.6% (Fig 1, Table 2). Such a high level of intraspecific distance indicate that there might be potential cryptic species and/or mis-identified sequences in these of 18 species (Table 2).

| Table 1. Genetic divergences in different taxonomic level within Calaphidinae. |
|-------------------------|--------|-----------------|-----------------|
| Comparison within       | Mean (%) | Minimum (%) | Maximum (%) |
| Species                 | 0.7     | 0              | 16.6           |
| Genus                   | 21.1    | 0              | 9.5            |
| Subtribe                | 13.4    | 3.4            | 22.8           |
| Tribe                   | 13.8    | 3.4            | 23.1           |

https://doi.org/10.1371/journal.pone.0176582.t001
MOTUs estimation

The number of MOTUs determined by ABGD differed slightly depending on the value of the relative gap width (Fig 2). When $p$ value was set at 0.0129, the number of MOTUs was 133 at relative gap width $X = 1$ or $X = 1.5$ (Fig 2). Based on ABGD result, 16 species were divided into 2–4 inner-groups in each species group (Fig 3). Three morphologically distinct species pairs: i) *Pterocallis alnijaponicae* and *P. nigrostriata*, ii) *Tiliaphis pseudoshinae* and *T. shinae* and iii) *Tuberculatus* (*Oriantuberculoides*) *capitatus* and *T. (O.) fangi* were clustered together as a single MOTU, respectively (Table 3).

The bPTP model recognized a total of 136 MOTUs except for outgroup based on the maximum likelihood method. Among 115 morphospecies, 19 species were subdivided into 2–4 inner-groups in each species group (Fig 3). This result was similar not only based on the number of MOTUs, but also based on MOTU compositions obtained from ABGD (Fig 3). Overall, the bPTP model tended to be more sensitive to MOTU delimitation. ABGD and bPTP resulted
in different estimates for 8 morphospecies (Fig 3). For example, 4 morphospecies, *Takecallis arundinariae*, *Tuberculatus* (*Nippocallis* *) kuricola*, *Tuberculatus* (*Arakawana*) *stigmatus* and *Myzocallis* (*Lineomyzocallis*) *bellus*, were subdivided into two groups each in the bPTP results. However, no subdivision was detected for these 4 species based on ABGD results (Fig 3). On the contrary, *Tiliaphis shinae* and *T. pseudoshinae* pair was clustered together as a single MOTU in the bPTP model (Fig 3).

### Neighbor joining analysis: Case of species delimitation

The Neighbor joining tree (NJ tree) was derived for these 899 COI sequences of the 115 species (S1 Fig). For majority of cases, sequence clusters in the NJ tree showed high congruence with morphological identification results. Practically, 90 species (78.3%) of all species could be clearly identified by COI sequence. However, for the remaining 25 species (21.7%), discrepancies between morphology and barcode based identification were detected. DNA barcoding analyses revealed the following: i) 15 cryptic species from 12 morphospecies, ii) six possible cryptic or potential misidentified sequences in the Genbank, iii) three morphologically distinct species pairs that sharing a single MOTU, and iv) four species with ambiguous inter- and intra-specific distances. Detailed results for these four cases are described as follows.

#### Case I: Discovering cryptic species

A total of 15 cryptic species from 12 morphospecies were found based on original description and other information of each morphospecies such as host-plant association and distributional information. Because in any case, it is impossible to identify DNA barcode for type materials. For each case, comparison between original and cryptic species was discussed.

A total of 66 individuals of 5 species belonging to genus *Calaphis* were analyzed in this study. DNA barcoding detected 2 cryptic species in *Calaphis flava* (Fig 4). Between group 1 and group 2, intergroup divergence was 2.9–4.0% (Fig 4). Group 3 consisted of 7 Canadian

| Subtribe       | Species               | No. of subgroups | Max. Intraspecific distance |
|----------------|-----------------------|------------------|-----------------------------|
| Calaphidina    | *Calaphis flava*      | 3                | 6.1%                        |
| Monaphidina    | *Monaphis antennata*  | 2                | 2.9%                        |
| Myzocallidina  | *Myzocallis boerneri*  | 2                | 8.2%                        |
|                | *Myzocallis coryli*   | 2                | 3.4%                        |
|                | *Tuberculatus annulatus* | 2             | 5.5%                        |
|                | *Tuberculatus higuchi* | 4                | 4.1%                        |
|                | *Tuberculatus indicus* | 2                | 2.9%                        |
|                | *Tuberculatus kashiwae* | 2             | 5.5%                        |
|                | *Tuberculatus punctata* | 2             | 6.6%                        |
|                | *Tuberculatus quercicus* | 2            | 8.2%                        |
|                | *Tuberculatus quercicola* | 2            | 6.0%                        |
|                | *Tuberculatus yokoyamai* | 2            | 4.1%                        |
| Panaphidina    | *Eucallipterus tiliae* | 2                | 2.9%                        |
|                | *Mesocallis corylicola* | 2               | 7.1%                        |
|                | *Shivaphis celti*     | 2                | 16.6%                       |
|                | *Takecallis arundicolens* | 3            | 10.4%                       |
|                | *Therioaphis trifoli*  | 2                | 1.9%                        |
|                | *Tiliaphis shinae*    | 2                | 10.3%                       |
|                | *Tinocallis zelkowae*  | 2                | 6.6%                        |

https://doi.org/10.1371/journal.pone.0176582.t002
Fig 2. Number of MOTUs by the prior intraspecific divergence using ABGD with two values of relative gap width. (A) $X = 1$. (B) $X = 1.5$.

https://doi.org/10.1371/journal.pone.0176582.g002
Fig 3. Maximum likelihood COI gene tree with delimited MOTUs by ABDG and PTP analyses. (Red S) Subdivided morphospecies. (Green C) combined morphologically different species. (Blue D) discordant between ABDG and PTP results.

https://doi.org/10.1371/journal.pone.0176582.g003
and 1 American individual was distinct from group 1 and group 2 with 4.5–6.1% of genetic divergence (Fig 4). This species was originally distributed throughout Europe and East Asia. Now it is regarded as a widespread species found in South Africa, Australia and North America. Thus, further studies are needed to compare European individuals of *C. flavus* in the future study to investigate whether this species is a real cosmopolitan species or a species complex.

A total of 23 individuals of *Eucallipterus tiliae* were collected from Europe: Czech Republic, France, Italy and Poland, North America: Canada and USA, and Oceania: New Zealand. ABGD, bPTP (Fig 3) and NJ tree revealed 1 cryptic species (group 2) of *E. tiliae* with about 2.9% of intergroup genetic divergence (Fig 5). Morphologically, individuals in group 1 and 2 are very similar. However, cryptic species (group 2) is distinguishable from typical *E. tiliae* (group 1) by having longer length of 3rd–5th antennal segment and 2nd tarsal segment.

| Subtribe     | Species 1              | Species 2              | Genetic distance |
|--------------|------------------------|------------------------|------------------|
| Myzocallidina | *Tuberclatus capitatus* | *Tuberclatus fangi*    | 0.5–1.3%         |
| Panaphidina  | *Pterocallis aijaponiae* | *Pterocallis nigrostriata* | 0.5%             |
|              | *Tiliaphis pseudoshinae* | *Tiliaphis shinae*     | 1.7–1.9%         |

Table 3. Case of sharing low genetic distances between morphologically distinct species pairs.

https://doi.org/10.1371/journal.pone.0176582.t003

Fig 4. Neighbor-joining tree of COI partial gene sequences of *Calaphis* spp. (66 sequences of 5 morphospecies).

https://doi.org/10.1371/journal.pone.0176582.g004
In genus *Eucallipterus*, only two species have been described in the world. *E. tiliae* is a common species widely distributed throughout Europe across central Asia, and South-Africa. It has also been introduced into North America and New Zealand [37]. Our results suggest that this species might be a species complex rather than a cosmopolitan species.

ABGD and bPTP analyses on 45 individuals of four Korean species of *Mesocallis* disclosed one cryptic species in *M. corylicola* (Figs 3 and 6). *M. corylicola* (group 1) and cryptic species (group 2) showed 5.5–7.1% of genetic distance (Fig 6). In contrast to such a high level of genetic distance, the cryptic species and *M. corylicola* are superficially similar in morphology. Compared to *M. corylicola*, the cryptic species has a shorter siphunculi and longer length of ultimate rostral segment (S3 Fig). Host-plant preference appeared to differ between the two species. Most *M. corylicola* was collected on *Corylus sieboldiana* while most cryptic species were collected on *C. heterophylla* (S1 Table). According to the original description, original species was collected on *Corylus sieboldiana* [59]. Thus, the two groups have different COI sequences, host-plant preference, and morphology.

Eight individuals of *Monaphis antennata* collected from Canada, Korea, and Poland were analyzed with ABGD, bPTP (Fig 7), and NJ tree (Fig 7). Korean individuals formed a group (group 1) distinct from Polish and Canadian individuals with 2.4–2.9% intergroup genetic divergence (Fig 7). *Monaphis* is a monotypic genus originally described from Europe. This species lives solitarily on *Betula* spp. In Korea, *Monaphis* is extremely rare. It has only been collected on *Betula schmidtii*. In Europe and Japan, it has been collected on *B. pendula* [60–61], *B. maximowicziana* and *B. platyphylla var. japonica* [59]. Although we could not perform morphological comparisons on subgroups, the Korean cluster seems to be a distinct species based on the molecular divergence level.

We analyzed 58 individuals of five *Takecallis* spp. collected from East Asia: China, Japan, and Korea, Europe: Czech Republic, France, and Italy, and North America: Canada and USA. According to both ABGD and bPTP analyses, *T. arundicolens* were separated into three groups (Figs 3 and 8). Genetic divergence between group 1 and group 2+3 ranged from 7.6% to 10.4%. Genetic divergence between group 2 and group 3 was 2.5%. In each group, morphological differences were only detected between alatoid nymphs. Alatoid nymphs in group 1 could be distinguished by a dark colored cauda with short dorsal abdominal setae (S4 Fig). Group 2 and group 3 shared similar morphology. However, group 3 could be distinguished from group 2 by shorter siphunculi with long filiform setae on the body (S6 Fig 4). *T. arundicolens* is one of common bamboo feeding species. Originally, this species was described from East Asia. It

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**Fig 5.** Neighbor-joining tree of COI partial gene sequences of *Eucallipterus* spp. (23 sequences).

https://doi.org/10.1371/journal.pone.0176582.g005
has been introduced into Europe and North America. Our results indicate that European populations (group 2) might be species distinct from Asian species (group 1 and group 3).

A total of 48 individuals of nine Tinocallis spp. collected from East Asia: Korea and North America: Canada and USA were analyzed. Both ABGD and bPTP analyses recognized two subgroups (group 1 and group 2) among 22 individuals of T. zelkowae collected from Korea (Fig 3). These groups showed genetic divergence of 5.5–6.5% (Fig 9). Morphologically,
individuals of group 1 have longer ultimate rostral segments than those of group 2 with pigmented dorsal abdominal elevations (S5 Fig). Group 1 is relatively rare. It was only collected on native Zelkova serrata (var. latifolia) growing in Korean mountain areas. T. zelkowae of the group 2 is one of the most common species dwelling on Zelkova trees in urban area (mostly the Japanese species Zelkova serrata (var. japonica)). Results of this study indicate that group 1 might be species separated from group 2 with different host-plant association.

In genus Tuberculatus, a total of seven cryptic species were discovered. Detailed results for each subgenus are provided as follows. A total of 69 individuals of two species belonging to the subgenus Acanthocallis collected from Japan and Korea were analyzed. Of 21 Korean individuals of Tuberculatus (Acanthocallis) quercicola, 16 individuals formed group 3, distinct from Japanese and remaining 5 Korean individuals of T. (A.) quercicola (group 2) with about 5.2% of intergroup genetic divergence (Figs 3 and 10). Individuals of group 3 were different from group 2 only by having shorter setae on 3rd antennal segment with more setae on 4th–5th antennal segments (S6 Fig). Watanabe et al. [62] have reported that Acanthocallis species tend to have high host specificity. It has been shown that Japanese T. (A.) quercicola and T. (A.) macrotuberculatus have distinct host-plant associations with Quercus mongolica spp. crispula and Q. dentate, respectively [62]. Likewise, three Korean species: T. (A.) quercicola in group 2, T. (A.) macrotuberculatus in group 1, and cryptic species in group 3 showed distinct host-plant association with Q. mongolica, Q. dentate, and Q. aliena, respectively. Therefore, different host associations between species can be used for species identification in this group.

A total of 26 individuals of four species belonging to subgenus Acanthotuberculatus were analyzed. As shown in Figs 3 and 11, 14 specimens of Tuberculatus (Acanthotuberculatus) indicus were split into two subgroups with intergroup genetic divergence ranging from 2.4% to 2.9%. Nine specimens collected from Korea were identical to undescribed species Tuberculatus sp. E (Yao, unpublished, COI sequence Genbank accession no. AB861448). Tuberculatus (A.)
and undescribed species, *Tuberculatus* sp. IY-C [63], showed inter-specific distance of 4.5% to 5.1%. No morphological comparisons was undertaken.

DNA barcoding uncovered five cryptic species among 86 specimens of seven species in subgenus *Orientuberculoides* (Fig 3). *Tuberculatus* (*Orientuberculoides*) *higuchii* was subdivided...
into four subgroups, including previously detected species \( T. (O.) higuchii \) A (group 1) and \( T. (O.) higuchii \) B (group 3) [63] and two newly detected subgroups (Fig 12). For convenience, these newly detected subgroups were named as \( T. (O.) higuchii \) C (group 2) and \( T. (O.) higuchii \) D (group 4). Among those four subgroups, intergroup genetic divergence of 2.8% to 4.1% was observed (Fig 12). \( T. (O.) higuchii \) A has shorter 2nd–4th antennal segments in comparison with other subgroups (S7 Fig). \( T. (O.) higuchii \) B is distinct from other subgroups by having shorter siphunculi (S7 Fig). \( T. (O.) higuchii \) C has slightly longer second tarsal segments and cauda (S7 Fig). \( T. (O.) higuchii \) D is distinguished from others by having more secondary sensilla on the 3rd antennal segment (S7 Fig).

**Fig 10.** Neighbor-joining tree of COI partial gene sequences of subgenus Acanthocallis spp. (69 sequences of 2 morphospecies).

https://doi.org/10.1371/journal.pone.0176582.g010

**Fig 11.** Neighbor-joining tree of COI partial gene sequences of subgenus Acanthotuberculus spp. (26 sequences of 4 morphospecies).

https://doi.org/10.1371/journal.pone.0176582.g011
Fourteen specimens of *Tuberculatus (Orientuberculoides) kashiwae* collected from Japan and Korea were divided into two subgroups with intergroup genetic divergence of 5.1% to 5.5% (Fig 12), confirming earlier suggestion by Yao [63]. All Korean specimens were clustered together with *T. (O.) kashiwae B* [63]. Only two sequences from Japan formed group 1 (*T. (O.) kashiwae A*).

A total of 25 individuals of *Tuberculatus (Orientuberculoides) yokoyamai* collected from Japan and Korea were analyzed. Results are shown in Fig 12. ABGD and bPTP analyses detected two cryptic species within *T. (O.) yokoyamai*, showing genetic divergence of 3.5% to 4.1% (Fig 12). Thirteen specimens collected from Japan and Korea formed group 1 together
with *Tuberculatus* sp. D Yao (unpublished, COI sequence Genbank accession no. AB861455). Morphologically, group 1 was distinguished from group 2 by having longer length of 5th antennal segment, siphunculi, and cauda (S8 Fig). There was no host-plant difference between the two groups.

COI sequences of 19 specimens of three species in the subgenus *Tuberculoides* revealed two groups within *Tuberculatus* (*Tuberculoides*) *annulatus* (Figs 3 and 13) with genetic divergence of 5.1–5.5%. All 10 individuals of group 1 were collected from Europe: France, Poland and UK. Group 2 comprised of 7 North American specimens, 3 New Zealand specimens, and one French specimen. One French specimen in group 2 has been regarded as an outlier in a previous study of Coeur d’Acier et al. [64]. However, this haplotype was present in North America and New Zealand. Morphological features of the two subgroups (group 1 and group 2) of *T. (T.) annulatus* were compared. Group 2 is distinguishable from group 1 by shorter antennae and smaller cauda knob with longer setae (S9 Fig). Based on molecular and morphological differences, group 1 and group 2 might be distinct species. It has been widely assumed that *T. (T.) annulatus* is introduced into North America, South America, Australia, and New Zealand from Europe [38]. However, such assumption need to be examined in future studies.

**Case II: Discovering possible cryptic or misidentified sequences in NCBI.** A total of 398 COI sequences of 73 species were downloaded from Genbank and used in this study. The following statements are only suggestions since specimen morphology could not be examined in this study.

In genus *Myzocallis*, a total of three possible discrepancies compared to current species concepts were identified. Detailed results of each subgenus are described as follows. We analyzed 26 sequences of four species belonging to subgenus *Myzocallis* collected from Europe: Czech Republic, France, Italy and Poland; North America: Canada and USA; and Oceania: New Zealand. ABGD and bPTP analyses revealed two subgroups in *Myzocallis* (*Myzocallis*) *coryli*. Group 1 and group 2 showed intergroup genetic divergence of 2.9–3.4% (Fig 14). Group 1 of *M. (M.) coryli* formed a sister group of *M. (M.) carpini* with genetic divergence of 2.7–3.1%. Group 1 mostly comprised of North American specimens except for one French specimen (Genabank accession no. KF639545) while all specimens of group 2 were from France. Coeur d’Acier et al. [64] have reported that *M. (M.) coryli* show exceptionally high intraspecific divergence due to the outlier. However, the outlier was the most common haplotype of *M. (M.)
*coryli* in our dataset. *M. (M.) coryli* is widely known as a cosmopolitan species. However, our results suggested that this species might be a species complex.

*Myzocallis* (*Myzocallis*) *boerneri* specimens collected from New Zealand and Italy showed genetic divergence of 8.4% (Fig 14). This species was originally described from Europe and introduced into New Zealand [65]. Assuming species identification of Italian *M. (M.) boerneri* is correct. Our results suggested that European and New Zealand populations are distinct species.

*COI* sequences of two *Neosymydobius* species, *Myzocallis* (*Neosymydobius*) *asclepiadis* and *M. (N.) punctata*, revealed possible misidentified sequences of *M. (N.) punctata* (Fig 15). As a result of ABGD and bPTP analyses, *M. (N.) punctata* was subdivided into two subgroups (Fig 3). Four sequences of *M. (N.) punctata* (Genbank accession no. EU701771, KR030925, KR043984, and KR037948) were completely identical to those of *M. (N.) asclepiadis* rather than to *M. (N.) punctata* with 100% support value (Fig 15). Our results suggested that 4 individuals (Genbank accession no. EU701771, KR030925, KR043984, and KR037948) of *M. (N.) punctata* might be misidentification of *M. (N.) asclepiadis*.

Among 22 specimens of five *Shivaphis* spp., 7 individuals of *Shivaphis celti* showed extremely high levels of intraspecific distance (16.1% to 16.6%) due to a single sequence (Genbank accession no. JQ920934) (Fig 16). Such a high intraspecific distance suggests that this specimen might be a misidentification or a cryptic species. Subsequent morphological re-examination of the voucher specimen for this sequence (Genbank accession no. JQ920934) is needed.

A total of 20 specimens of four *Tiliaphis* species were analyzed. We found a possible misidentified sequence which was identified as *Tiliaphis shinae* (Genbank accession no. GU978821) (Fig 17). Subsequently, we re-examined the voucher specimen of this sequence and found that this specimen was in fact *T. coreana*.

Three sequences of *Tuberculatus* (*Tuberculatus*) *querceus* were divided into two subgroups with divergence of 8.2% (Fig 18). This species is distributed through Europe to South-western
Asia. It is also introduced into Canada [30, 38]. However, our results suggested that European and Canadian specimens might be distinct species.

**Case III: Low genetic distances between morphologically distinct species.** Morphologically, *Pterocallis alnijaponicae* and *P. nigrostriata* are easily distinguishable species (Fig 19). As
shown in Fig 19, *P. nigrostriata* has 5–6 long and conspicuous dorsal abdominal tubercles and forewing with unique marginal patch while *P. alnijaponicae* has 3–4 short dorsal abdominal tubercles and forewing without marginal patches. Unexpectedly, low genetic divergence (0.5%) between *P. alnijaponicae* and *P. nigrostriata* was found (Fig 20). ABGD and bPTP analyses also supported these results by combining *P. alnijaponicae* and *P. nigrostriata* as a single MOTU (Fig 3). Such discrepancies between morphological and DNA barcoding results raise a question about the validity of these species.

*Tiliaphis pseudoshinae* and *T. shinae* are morphologically similar. However, *T. pseudoshinae* can be distinguished from *T. shinae* by having longer ultimate rostral segment, shorter
antenna, and less secondary sensoria on the 3rd antennal segment (Fig 21). Between these two species, sequence divergence of 2.1% to 2.5% was observed (Fig 17). Based on ABGD analysis, T. pseudoshinae and T. shinae were separated into distinct MOTUs. However, bPTP analysis grouped these two species as a single MOTU (Fig 3). Thus, comparing more specimens with multiple generic markers is needed for better delimitation of these species.

Four individuals of Tuberculatus (Orientuberculoides) capitatus and nine individuals of T. (O.) fangi were analyzed. T. (O.) capitatus and to T. (O.) fangi can be distinguished by having different shapes and lengths of setae (Fig 22). However, their genetic divergence was only 0.5% to 1.5% (Fig 12). In fact, different shapes, arrangement, and lengths of hairs are often can be
Fig 21. Alate vivipara of *Tiliaphis pseudoshinae* (A, C) and *T. shinae* (B, D). (A-B) antenna. (C-D) ultimate rostral segment (scale bars 0.1mm).

https://doi.org/10.1371/journal.pone.0176582.g021

Fig 22. Alate vivipara of *Tuberculaphis* (Orientuberculoides) *capitatus* (A, C, E, G) and *T. (O.) fangi* (B, D, F, H). (A-B) head. (C-D) setae on thorax. (E-F) abdominal dorsal tubercles. (G-H) 4th abdominal marginal tubercle (scale bars, 0.1mm).

https://doi.org/10.1371/journal.pone.0176582.g022
critical characteristics for aphid species delimitation. A review of the variations within and between these species is still required.

**Case IV: Ambiguous genetic distances.** In this study, six individuals of three species in subgenus *Lineomyzocallis* belonging to genus *Myzocallis* were analyzed. DNA barcoding showed 1.7% intraspecific distance between two Canadian and one American specimen of *Myzocallis (L.)* *bellus* (Fig 23). ABGD analysis detected them as a single MOTU. However, bPTP analysis separated these Canadian and American specimens as distinct MOTUs. This situation requires further morphological and molecular analysis.

A total of 57 specimens belonging to 3 species of *Therioaphis* were analyzed in this study. As shown in Fig 24, 43 specimens of *Therioaphis (Pterocallidium) trifolii* were subdivided into two subgroups with intergroup barcode divergence of 1.9% (Fig 24). Group 1 comprised of American, Canadian, and Korean specimens while group 2 only contained Canadian specimens. However, both groups formed a single MOTU according to both ABGD and bPTP analyses. There are four subspecies, *T. (P.)* *trifolii albae*, *T. (P.)* *trifolii maculata*, *T. (P.)* *trifolii trifolii* and *T. (P.)* *trifolii ventromaculata* in of *T. (P.)* *trifolii* [14, 38]. Detailed molecular studies have not been conducted on this group of taxa. Thus, comparing worldwide samples of *T. (P.)* *trifolii* spp. is needed in future studies.

Within *Tuberculatus (Arakawana) stigmatus*, two distinct subgroups were detected based on NJ tree and bPTP analyses with genetic divergence of 1.5% to 1.9% (Figs 3 and 25). Individuals in group 1 and group 2 showed distinct morphological features (S10 Fig). Group 2 has shorter siphunculi and small abdominal marginal tubercles without marginal tubercle on the 5th abdominal segment (S10 Fig). Based on their morphological differences, group 1 and group 2 might be distinct species despite the relatively low divergence level between these two subgroups.

Among 38 individuals of *Tuberculatus (Nippocallis) kuricola*, two subgroups (group 1 and group 2) were detected based on NJ tree and bPTP analysis (Fig 3). Genetic divergence between the two groups was 1.9% (Fig 26). Morphologically, the two groups are superficially similar. However, group 2 can be recognized by having 3–4 setae on each dorsal abdominal tubercle (rather than 2 in group 1) with a shorter ultimate rostral segment (S11 Fig). Takahashi (1936) has described subspecies *T. (N.)* *kuricola cantoensis*. However, it is currently unclear whether *T. (N.)* *kuricola cantoensis* and *T. (N.)* *kuricola kuricola* correspond to these two genetically divided subgroups.
Discussion

This study demonstrates that DNA barcoding can be used to reliably identify aphid species in the subfamily Calaphidinae. DNA barcoding of 115 morphospecies (899 sequences) revealed 25 discordances between DNA barcoding results and morphology results. These conflicts involved 18 cases of exceptionally high intraspecific distances, three morphologically distinct species pairs with low genetic distances and four cases of ambiguous intraspecific distances. Except for four undeterminable cases, a total of 15 cryptic species were identified from 12 morphospecies. Among these cases, slight morphological differences were detected in seven species complexes. In the most cases, morphological differences were due to different lengths of various body parts rather than a different shapes or numerical characters. Slight length differences...
are easy to be overlooked. They can be obscured by intraspecific variations. From these results, we can infer the presence of cryptic diversity in Calaphidinae.

In this study, distinct host-plant associations were revealed in three species complexes: *Mesocallis corylicola*, *Tinocallis* (*Tinocallis*) *zelkowae* and *Tuberculatus* (*Acanthocallis*) *quercicola*. Differences in host-plant associations are often critical for aphid species identification. Host-plant shifts and subsequent genetic differentiation can lead to speciation (ecological species concept) [66–69]. These ecological differences between cryptic species emphasize the necessity to reexamine different host-plant associated populations in a single species.

We also confirmed that species with a wide distribution could have more possibilities to include cryptic species. Some species assessed in this study demonstrated this assumption. For example, *Calaphis flava*, which has been known as invasive species with cosmopolitan distribution, included subgroups with each restricted collection area. However, most previous DNA barcoding studies in aphids targeted samples only collected within restricted area [7, 35, 64]. Our results suggest that multiregional sampling would be very important to recognize cryptic diversity.

However, we also found deep intraspecific divergence within species complexes without obvious host-plant and geographic difference. In fact, groups within *Takecallis arundicolens* and *Tuberculatus* (*Nippocallis*) *kuricola* were found together within the same individual colony, respectively. In particular, viviparous females of *T. arundicolens* (group 1 and group 3) were not distinguishable by morphology. Considering that both bamboo and chestnut trees are urban landscape plants, the phenomenon (genetically distinct groups occur in the same colony) might indicate that human mediated transportation might have played a role in subsequent colony merging.

We found three morphologically distinct species pairs with relatively low genetic distances: *Pterocallis alnijaponicae* and *P. nigrostriata*, *Tiliaphis pseudoshinae* and *T. shinae* and *Tuberculatus* (*Orientuberculoides*) *capitatus* and *T. (O.) fangi*. This could be due to many reasons such as rapid radiation, balanced polymorphism and introgressive hybridization [70]. In the three DNA barcode sharing species pairs, it was common that each pair of species was collected together on the same host-plants and geographical regions. Considering that, there can be opportunities of hybridization between species pair during sexual reproduction on the same
host-plant. Such phenomenon has been frequently reported in several taxa of aphids [71–73]. However, conducting additional ecological and molecular research is needed to verify whether barcode sharing is due to hybridization.

In insects, species delimitation threshold value from 2 to 5% of divergence, depending on the group, is generally accepted as a means to estimate species boundaries. For example, 2% intraspecific divergence may indicate the existence of hidden species in Lepidoptera [11, 74]. However, within Diptera, this level of divergence only represents an intraspecific difference [75–77]. In the suborder Heteroptera, threshold value of 2.2% has been applied for DNA barcode based species delimitation [78–79]. In this study, about 2.5% of species delimitation value could be applied for most of species. However, there is clear limits applying certain species delimitation threshold value in every case. In this study, we found highly varied interspecific genetic variation for COI, from 0.5% to 20.1% between congeneric species of Calaphidinae. These results suggested that applying a single threshold value might not work for all members in this group. To get more accurate species delimitation results, combination of additional information such as different genetic markers, morphological characters, and ecological differences is required. For example, in the present study, morphological re-examination of two species, *Tuberculatus* (*Arakawana*) *stigmatus* and *Tuberculatus* (*Nippocallis*) *kuricola*, with intraspecific divergence of 1.9% resulted in the detection of morphological differences, suggesting that additional species are present.

Some aphid species are important pest of various crop and ornamental plants. They play a critical role in ecosystems. Although DNA barcoding has been carried out several times for aphids before, most studies have been mainly focused on the largest subfamily Aphidinae with less emphasis on other subfamilies [7, 13, 30, 64]. DNA barcodes produced in this study are of value in aiding in the identification of species of Calaphidinae. Further, remarkable cryptic diversity and suspicious cases such as barcode sharing species pairs are detected in this study. Our findings suggest that many more cryptic diversity are not yet been uncovered in aphids. For more accurate and higher resolution of possibly overlooked species diversity investigation, future studies should focus on well-designed sampling plan to reflect morphological, ecological and distributinal diversities within species.

**Supporting information**

S1 Fig. Neighbor-joinig tree for the 899 individuals of 115 morphospecies based on COI barcoding region.

(TIF)

S2 Fig. Alate vivipara of 2 subgroups of *Eucallipterus tiliae* group 1 (A-B) and group 2 (C-D). (A, C) antenna. (B, D) 2nd segment of hind tarsi (scale bars, 0.1mm).

(TIF)

S3 Fig. Alate vivipara of 2 subgroups of *Mesocallis corylicola* group 1 (A, C) and group 2 (B, D). (A-B) siphunculi. (C-D) ultimate rostral segment (scale bars, 0.05mm).

(TIF)

S4 Fig. Alatoid nymph of 3 subgroups of *Takecallis arundicolens* group 1 (A), group 2 (B) and group 3 (C). (A-C) body (scale bars, 0.5mm).

(TIF)

S5 Fig. Alate vivipara of 2 subgroups of *Tinocallis zelkowae* group 1 (A, C) and group 2 (B, D). (A-B) ultimate rostral segment. (C-D) abdomen.

(TIF)
S6 Fig. Alate vivipara of 2 subgroups of *Tuberculatus* (*Acanthocallis*) *quercicola* group 2 (A) and group 3 (B). (A-B) antenna (scale bars 0.1mm).

(TIF)

S7 Fig. Alate vivipara of 4 subgroups of *Tuberculatus* (*Orientuberculoides*) *higuchii* group 1 (A, E, I, M), group 2 (B, F, J, N), group 3 (C, G, K, O) and group 4 (D, H, L, P). (A-D) cauda. (E-H) siphunculi. (I-L) 2nd segment of hind tarsi. (M-P) antenna (scale bars 0.1mm).

(TIF)

S8 Fig. Alate vivipara of 2 subgroups of *Tuberculatus* (*Orientuberculoides*) *yokoyamai* group 1 (A, C, E) and group 2 (B, D, F). (A-B) antennae. (C-D) cauda. (E-F) siphunculi (scale bars 0.1mm).

(TIF)

S9 Fig. Alate vivipara of 2 subgroups of *Tuberculatus* (*Tuberculoides*) *annulatus* group 1 (A, C) and group 2 (B, D). (A-B) antenna. (C-D) cauda (scale bars, 0.1mm).

(TIF)

S10 Fig. Alate vivipara of 2 subgroups of *Tuberculatus* (*Arakawana*) *stigmatus*, group 1 (A, C, E) and group 2 (B, D, F). (A-B) abdomen. (C-D) siphunculi. (E-F) antenna.

(TIF)

S11 Fig. Alate vivipara of 2 subgroups of *Tuberculatus* (*Nippocallis*) *kuricola*, group 1 (A, C) and group 2 (B, D). (A-B) ultimate rostral segment. (C-D) abdomen.

(TIF)

S1 Table. Detailed collection information and Genbank accession numbers of species used in this study.

(XLSX)

S2 Table. Detailed collection information and Genbank accession numbers of sequences downloaded from Genbank.

(XLSX)

**Acknowledgments**

We are grateful to Hyoseok Lee for his great support and help particularly collecting samples with Yerim Lee. We thanks to Dr. Hyojoong Kim, Dr. Jaroslav Holman, Dr. Jong-Kook Jung, Dr. Lukasz Depa and Dr. Ram Duwal Keshari. This project was funded by the Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Education, Science (NRF-2015R1D1A1A01059031).

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Supervision: YL.

Validation: YL.

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