A new subterranean species of pseudocrangonyctid amphipod, *Pseudocrangonyx daejeonensis* sp. n. is described from the interstitial waters in Daejeon, Korea. *Pseudocrangonyx daejeonensis* sp. n. is distinguished from three morphologically similar congeners, *P. coreanus* Uéno, 1966, *P. febras* Sidorov, 2009, and *P. gudariensis* Tomikawa & Sato, 2016, by the characteristics of antenna 1, antenna 2, mandible, gnathopod 2, pleopods, uropods 1–2, and telson. Molecular phylogenetic analyses based on nuclear 28S rRNA and histone H3, and mitochondrial cytochrome *c* oxidase subunit I and 16S rRNA genes revealed that *P. daejeonensis* is a sister species of the unnamed *Pseudocrangonyx* sp. 3 inhabiting central Japan.

Keywords
Crangonyctoidea, Korean Peninsula, interstitial water, molecular phylogeny

Introduction
Amphipod species of the genus *Pseudocrangonyx* Akatsuka & Komai, 1922 have been known from subterranean waters and springs in Japan, the Korean Peninsula, Eastern China, and the Far East of Russia (Sidorov and Holsinger 2007; Tomikawa et al. 2016; Zhao and Hou 2017). Among the 22 known species of *Pseudocrangonyx*, only two...
species were recorded in Korean waters (Uéno 1966): *P. asiaticus* Uéno, 1934 and *P. coreanus* Uéno, 1966. When Uéno (1966) described *P. coreanus* based on specimens collected from the Korean Peninsula, he clearly stated that morphological variations in the antennae, maxilla 1, uropod 3, and telson were observed among six populations of this species. Because recent systematic studies of *Pseudocrangonyx* in other regions have shown high species diversity within this genus (e.g., Tomikawa et al. 2016), it is highly possible that the true species diversity of *Pseudocrangonyx* amphipods inhabiting the Korean Peninsula remains under-estimated.

Recently, unidentified specimens of *Pseudocrangonyx* were collected during field surveys of interstitial invertebrates in Korea by the first author. In this paper, we describe and illustrate this amphipod as a new species. In addition, the phylogenetic position of the new species was estimated using nuclear 28S rRNA and histone H3, and mitochondrial cytochrome c oxidase subunit I (COI) and 16S rRNA sequence data.

**Materials and methods**

**Sampling**

*Pseudocrangonyx* specimens were collected from interstitial water in Heukseok-dong, Seo-gu, Daejeon, South Korea (Fig. 1) using a 50 μm fine-mesh net. Specimens were pumped up with 80–100 L of interstitial water at 1–1.5 m beneath hyporheic zones using a core (Lee and Park 2016). All specimens were immediately preserved in 95% ethanol. The specimens are deposited in the collection of the Nakdonggang National Institute of Biological Resources, Korea (NNIBR) and in the Zoological Collection of Kyoto University (KUZ).

**Morphological observation**

The specimens were dissected in 70 % ethanol and mounted in gum-chloral medium on glass slides under a stereomicroscope (Model SZX-7; Olympus, Tokyo, Japan). Specimens were examined using a Nikon Eclipse Ni light microscope (Nikon, Tokyo, Japan) and illustrated with the aid of a drawing tube. The body length from the tip of the rostrum to the base of the telson was measured along the dorsal curvature to the nearest 0.1 mm. The nomenclature of the setal patterns on the mandibular palp follows that of Stock (1974).

**Molecular phylogenetic analyses**

Methods of the genomic DNA extraction, PCR and DNA sequencing were performed following Tomikawa et al. (2016). Accordingly, nine DNA sequences of nuclear 28S
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Figure 1. Map showing the collection locality of the specimens examined in this study.

rRNA, histone H3, COI and 16S rRNA from three Korean *Pseudocrangonyx* specimens were newly obtained in this study, and deposited into the International Nucleotide Sequence Database Collaboration (INSDC) through the DNA Data Bank of Japan (Table 1).

The OTU set for phylogenetic analyses was almost identical to that used in the previous phylogenetic analyses in Tomikawa et al. (2016) with the DNA sequences of
Table 1. Samples used for the phylogenetic analyses. The information on the vouchers is accompanied by the collection locality and the INSDC accession numbers. Sequences marked with an asterisk were obtained for the first time in the present study. Acronyms: IZCAS, Institute of Zoology, Chinese Academy of Sciences; NNIBR, Nakdonggang National Institute of Biological Resources; NSMT, National Museum of Nature and Science, Tokyo.

| Species                        | Voucher or isolate #       | Locality or country | INSDC #         |
|--------------------------------|----------------------------|---------------------|-----------------|
|                                |                            |                     | 28S  | Histone H3 | COI       | 16S       |
| Pseudocrangonyx                |                            |                     |      |            |           |           |
| P. daejeonensis sp. n.         | NNIBRIV1 (Holotype)        | Daejeon, Korea      | LC322136*      | LC322138*  | LC322137*  | LC322135*  |
| P. daejeonensis sp. n.         | NNIBRIV2 (Paratype)        | Daejeon, Korea      | LC322143*      |            |           |           |
| P. daejeonensis sp. n.         | NNIBRIV3 (Paratype)        | Daejeon, Korea      | LC322140*      | LC322142*  | LC322141*  | LC322139*  |
| P. guadariensis                | NSMT-Cr 24605             | Aomori, Japan       | LC171498       | LC171500   | LC171499   | LC171497   |
| P. yezonis                     | G1280                      | Hokkaido, Japan     | LC171518       | LC171520   | LC171519   | LC171517   |
| P. yezonis                     | G1279                      | Akita, Japan        | LC171514       | LC171516   | LC171515   | LC171513   |
| Pseudocrangonyx sp. 1          | G400                       | Iwate, Japan        |                 |            | LC171479   |           |
| Pseudocrangonyx sp. 1          | G1281                      | Iwate, Japan        |                 |            | LC171521   |           |
| Pseudocrangonyx sp. 2          | G1283                      | Okayama, Japan      | LC171525       | LC171527   | LC171526   | LC171524   |
| Pseudocrangonyx sp. 2          | G1277                      | Yamaguchi, Japan    | LC171506       | LC171508   | LC171507   | LC171505   |
| Pseudocrangonyx sp. 2          | G1278                      | Yamaguchi, Japan    | LC171510       | LC171512   | LC171511   | LC171509   |
| Pseudocrangonyx sp. 3          | G404                       | Shiga, Japan        | LC171488       | LC171489   |             |           |
| Pseudocrangonyx sp. 3          | G405                       | Shiga, Japan        | LC171491       | LC171493   | LC171492   | LC171490   |
| Pseudocrangonyx sp. 3          | G406                       | Shiga, Japan        | LC171495       | LC171496   |             | LC171494   |
| Pseudocrangonyx sp. 4          | G1282                      | Shiga, Japan        | LC171523       |             |             | LC171522   |
| Pseudocrangonyx sp. 5          | G402                       | Shimane, Japan      | LC171485       | LC171487   | LC171486   | LC171484   |
| Pseudocrangonyx sp. 5          | G401                       | Shimane, Japan      | LC171481       | LC171483   | LC171482   | LC171480   |
| Pseudocrangonyx sp. 5          | G1271                      | Kagawa, Japan       | LC171502       | LC171504   | LC171503   | LC171501   |
| Pseudocrangonyx sp. 5          | G1295                      | Kochi, Japan        | LC171533       | LC171535   | LC171534   | LC171532   |
| Pseudocrangonyx sp. 5          | G1296                      | Kochi, Japan        | LC171537       | LC171539   | LC171538   | LC171536   |
| Pseudocrangonyx sp. 5          | G1294                      | Ehime, Japan        | LC171529       | LC171531   | LC171530   | LC171528   |
| Pseudocrangonyx sp. 6          | G1297                      | Gifu, Japan         | LC171541       | LC171543   | LC171542   | LC171540   |
| P. holsingeri                  |                            | Russian Far East    | KJ871679       |             |             | KF153111   |
| P. korkishkoorum               | B1                         | Russian Far East    | KJ871678       |             |             | KF153107   |
| P. korkishkoorum               | B2                         | Russian Far East    | KF153108       |             |             |           |
| P. korkishkoorum               | B3                         | Russian Far East    | KF153109       |             |             |           |
| P. korkishkoorum               | N1                         | Russian Far East    | KJ871676       |             |             | KF153105   |
| P. korkishkoorum               | N2                         | Russian Far East    | KJ871677       |             |             | KF153106   |
| P. kseniae                     |                            | Russian Far East    | KJ871675       |             |             | KF153115   |
| P. susanaensis                 |                            | Russian Far East    | KF153113       |             |             |           |
| P. sympatricus                 |                            | Russian Far East    | KF153112       |             |             |           |
| P. tianovi                     |                            | Russian Far East    | KJ871674       |             |             | KF153110   |
| P. elegantulus                 |                            | IZCAS I-A1602-2     | China           |             |             | KY436646   |
| Outgroup                       |                            |                     |                 |             |             | KY436647   |
| Cryomortygius thingvallensis   |                            |                     |                 |             |             | HQ286009   |
| Eocrangonyx primoryensis       |                            |                     |                 |             |             | HQ286011   |
| Crangonyx floridanus           | G1322                      | Chiba, Japan        | LC171549       |             | LC171550   | LC171548   |
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P. elegantulus Hou in Zhao and Hou (2017) (Table 1). The alignments of H3 and COI were trivial, as no indels were observed. 28S and 16S sequences were aligned using MAFFT v. 7.310 (Katoh and Standley 2013). The lengths of the 28S, H3, COI and 16S were 1357, 328, 658, and 430 bp, respectively. The concatenated sequences thus yielded 2773 bp of alignment positions. Phylogenetic trees were constructed using maximum likelihood (ML) and Bayesian inference (BI). The ML phylogeny was constructed using RAxML v. 8.2.8 (Stamatakis 2014) with the substitution model set as GTRCAT, immediately after nonparametric bootstrapping (BS) conducted with 1000 replicates. The best-fit partition scheme was identified with Akaike information criterion using PartitionFinder v. 2.1.1 (Lanfear et al. 2017) with the “greedy” algorithm (Lanfear et al. 2012): 28S/1st and 2nd positions of H3/H3 3rd position/COI 1st position/COI 2nd position/COI 3rd position/16S. BI and Bayesian posterior probabilities (PPs) were estimated using MrBayes v. 3.2.6 (Ronquist et al. 2012). The best-fit partition scheme and models for each partition were selected based on the Bayesian information criterion using PartitionFinder with the “greedy” algorithm: for 28S, GTR+G; for H3 1st and 2nd position, JC+I; for H3 3rd position, K80+G; for COI 1st position, SYM+I+G; for COI 2nd position, F81+I; for COI 3rd position, GTR+I+G; and GTR+I+G for 16S. Two independent runs of four Markov chains were conducted for 20 million generations, and the tree was sampled every 100 generations. The parameter estimates and convergence were checked using Tracer v. 1.6.0 (Rambaut and Drummond 2013), and the first 50001 trees were discarded based on the results.

Taxonomy

Family Pseudocrangonyctidae Holsinger, 1989
Genus Pseudocrangonyx Akatsuka & Komai, 1922

Pseudocrangonyx daejeonensis sp. n.
http://zoobank.org/ECC7F708-DD43-4A48-9458-B6DA59265796
Figs 2–9

Material examined. Holotype: Female (NNIBRIV1, 3.8 mm), Heukseok-dong (36°15.65’N, 127°20.43’E), Daejeon, Korea, collected by Lee CW on 31 May 2017. Paratypes: 1 male (NNIBRIV2, 2.7 mm), 1 female (NNIBRIV3, 2.3 mm), 3 females (KUZ Z1924), data same as for holotype.

Etymology. The specific name is an adjective derived from the type locality name of the new species.

Description. Female [NNIBRIV1, 3.8mm]. Head (Fig. 2) with short dorsal setae; rostrum reduced; lateral cephalic lobe rounded; antennal sinus shallow with rounded angle; eyes absent. Dorsal margin of pleonites 1–3 and urosomites 1–2 with setae (Fig. 2). Ventral margin of urosomite 1 without setae (Fig. 2). Ventral margin of epimeral plate 1 with seta, posterovertral corner rounded with seta (Fig. 2); ventral
and posterior margins of plate 2 each with seta, posteroventral corner rounded with seta (Fig. 2); ventral and posterior margins of plate 3 each with 2 setae, posteroventral corner rounded with seta (Fig. 2).

Antenna 1 (Fig. 3A) 0.38 times as long as body length, peduncular articles 1 to 3 in length ratio of 1.0 : 0.5 : 0.4; accessory flagellum (Fig. 3B) 2-articulate, terminal article with 3 setae and 1 aesthetasc; primary flagellum 10-articulate, 1 aesthetasc on some articles. Antenna 2 (Fig. 3C) 0.58 times as long as antenna 1; flagellum 0.65 times as long as peduncular articles 4 and 5 combined, consisting of 4 articles; calceoli absent.

Upper lip (Fig. 3D) with rounded anterior margin, bearing fine setae. Mandibles (Fig. 3F, G, H) with left and right incisors with 5- and 4-dentate, respectively; left lacinia mobilis 4-dentate, right lacinia bifid, bearing many teeth; molar process triturative; accessory setal rows of left and right mandibles with 3- and 2- pectinate setae, respectively; palp 3-articulate, article 3 with 1 A-, 7 D-, and 3 E-setae. Lower lip (Fig. 3I) with broad outer lobes with fine setae, mandibular process of outer lobe rounded apically; inner lobes indistinct. Maxilla 1 (Fig. 3J) with inner and outer plates, and palp; inner plate subovate with 2 plumose setae; outer plate subrectangular with 7 serrate teeth apically; palp 2-articulate, longer than outer plate, article 2 with 2 apical robust setae. Maxilliped (Fig. 4A) with inner and outer plates, and palp; inner plate reaching base of palp article 1, with 2 apical robust setae; outer plate not exceeding end of palp article 1, with 2 plumose setae and some medial setae; palp 4-articulate, medial margin of article 2 lined with setae, article 4 with nail.

Gnathopod 1 (Fig. 4B) with subquadrate coxa, bearing setae on its anterodistal and posteroventral corners, width 1.9 times as long as depth; posterior margin of basis with 3 setae; posterodistal corner of carpus with slender setae, some weakly pectinate; propodus stout, subchelate, palmar margin with 3 medial and 3 lateral robust setae; posterior
Figure 3. *Pseudocrangonyx daejeonensis* sp. n., holotype, female (3.8 mm): A–D, F–K paratype female (2.3 mm): E. A antenna 1, lateral view B accessory flagellum of antenna 1, lateral view C antenna 2, medial view D upper lip, anterior view E upper lip, anterior view F left mandible, medial view G incisor, lacinia mobilis, and molar process of left mandible, medial view H incisor and lacinia mobilis of right mandible, medial view I lower lip, ventral view J maxilla 1, dorsal view K maxilla 2, ventral view.
Figure 4. *Pseudocrangonyx daejeonensis* sp. n., holotype, female (3.8 mm). 

A maxilliped, dorsal view

B gnathopod 1, medial view

C palmar margin of propodus and dactylus of gnathopod 1, medial view

D gnathopod 2, medial view

E palmar margin of propodus and dactylus of gnathopod 2, medial view.
margin of dactylus dentate (Fig. 4C). Gnathopod 2 (Fig. 4D) with subquadrate coxa, bearing setae on its anterodistal and posteroventral corners, width 1.6 times as long as depth; posterior margin of basis with 4 setae; posterodistal corner of carpus with slender setae, some weakly pectinate; propodus stout, subchelate, palmar margin with 7 medial and 2 lateral robust setae; posterior margin of dactylus dentate (Fig. 4E). Pereopod 3 (Fig. 5A) with subquadrate coxa bearing setae on its anterodistal and posteroventral corners, width 1.6 times as long as depth; anterior and posterior margins of basis with 2 and 4 setae, respectively; merus, carpus, and propodus in length ratio of 1.0 : 0.9 : 0.8; posterior margin and submargin of dactylus each with seta (Fig. 5B). Pereopod 4 (Fig. 5C) with subquadrate coxa bearing setae on its anterodistal corner, width 1.7 times as long as depth; anterior and posterior margins of basis each with 4 setae; merus, carpus, and propodus in length ratio of 1.0 : 0.7 : 0.8; posterior margin and submargin of dactylus each with seta (Fig. 5D). Pereopod 5 (Fig. 6A) with weakly bilobed coxa bearing setae on anterior and posterior lobes, width 1.7 times as long as depth; anterior and posterior margins of basis with 5 and 6 setae, respectively; merus, carpus, and propodus in length ratio of 1.0 : 0.7 : 0.8; anterior margin of dactylus with 2 setae (Fig. 6B). Pereopod 6 (Fig. 6C) with coxa bearing concave lower margin, marginally bare; anterior and posterior margins of basis with 5 and 3 setae, respectively; merus, carpus, and propodus in length ratio of 1.0 : 0.8 : 0.9; anterior margin of dactylus with 2 setae (Fig. 6D). Pereopod 7 (Fig. 6E) with subtriangular coxa, bearing seta on posteroproximal corner; anterior and posterior margins of basis with 3 and 4 setae, respectively; merus, carpus, and propodus in length ratio of 1.0 : 0.8 : 1.1; anterior margin of dactylus with 2 setae (Fig. 6F).

Coxal gills (Figs 4D, 5A, 6A, C) on gnathopod 2 and pereopods 3–6; sternal gills absent. Brood plates (Figs 4D, 5A, C, 6A) slender, with numerous setae, on gnathopod 2 and pereopods 3–5.

Peduncle of pleopod 1 (Fig. 7A) with 1 outer marginal and 1 outerdistal seta; peduncle of pleopod 2 (Fig. 7C) with outerdistal seta; peduncle of pleopod 3 (Fig. 7D) lacking marginal and distal setae. Pleopods 1–3 with paired retinacula (Fig. 7B), and lacking bifid setae (clothes-pin setae) on inner basal margin of inner ramus; inner ramus of pleopods 1–3 3-, 3-, and 2-articulate (Fig. 7A, C, D); outer ramus of pleopods 1–3 4-, 3-, and 2-articulate (Fig. 7A, C, D).

Uropod 1 (Fig. 7E) with basofacial seta on peduncle; inner ramus 0.87 times as long as peduncle, inner margin of former with 2 robust setae, outer margin bare, basal part with slender seta; outer ramus 0.63 times as long as inner, marginally bare. Uropod 2 (Fig. 7F) with inner and outer rami; inner ramus 1.10 times as long as peduncle, its inner margin with robust seta, outer margin without setae; outer ramus 0.68 times as long as inner ramus, marginally bare. Uropod 3 (Fig. 7G) with peduncle 0.34 times as long as outer ramus, with 1 robust and 1 slender setae; inner ramus absent; outer ramus 2-articulate, proximal article with robust setae, terminal article 0.32 times as long as proximal article, with 3 distal setae. Telson (Fig. 7H) length 1.3 times as long as wide, cleft for 0.08 times of length, each telson lobe with 2 lateral penicillate setae, 1 apical robust and 1 apical short setae.
Male [NNIBRIV2, 2.7 mm]. Antenna 1 (Fig. 8A, B) 0.46 times as long as body length, primary flagellum 7-articulate, 1 aesthetasc on some articles. Antenna 2 (Fig. 8C) 0.57 times as long as antenna 1; flagellum 0.72 times as long as peduncular articles 4 and 5 combined, consisting of 4 articles, first 2 of which with calceoli (Fig. 8D).

Gnathopod 1 (Fig. 8E) with coxa width 1.84 times as long as depth; palmar margin with 3 medial and 3 lateral robust setae (Fig. 8F). Gnathopod 2 (Fig. 8G) with coxa width 1.66 times as long as depth; palmar margin with 3 medial and 4 lateral robust setae (Fig. 8H).

Uropod 1 (Fig. 9A) with robust seta on inner margin of inner ramus; outer ramus 0.62 times as long as inner. Uropod 2 (Fig. 9B) with 2 serrate and 4 simple robust setae and slender seta at distal part. Uropod 3 (Fig. 9C) with peduncle 0.32 times as long as outer ramus; terminal article of outer ramus 0.5 times as long as proximal article.

**Variation.** Peduncle of pleopod 1 with or without seta on outer margin.

**Distribution.** This species is known only from the type locality.

**Molecular phylogenetic position.** The BI tree (mean ln $L = −14039.10$; Fig. 10) for estimating the phylogenetic position of the new species had an identical topology to that of the ML tree (ln $L = −14504.12$; not shown). *Pseudocrangonyx daejeonensis* belonged to a well-supported clade (BS = 98 %, PP = 0.99) containing the three phylogroups known from the western parts of Honshu and Shikoku, i.e., *Pseudocrangonyx* spp. 3–5. The new
Figure 6. *Pseudocrangonyx daejeonensis* sp. n., holotype, female (3.8 mm). **A** pereopod 5, medial view **B** dactylus of pereopod 5, medial view **C** pereopod 6, medial view **D** dactylus of pereopod 6, medial view **E** pereopod 7, lateral view **F** dactylus of pereopod 7, lateral view.
Figure 7. *Pseudocrangonyx daejeonensis* sp. n., holotype, female (3.8 mm). **A** pleopod 1, anterior view **B** retinacula on peduncle of pleopod 1, anterior view **C** pleopod 2, anterior view **D** pleopod 3, anterior view **E** uropod 1, dorsal view **F** uropod 2, ventral view **G** uropod 3, dorsal view **H** telson, ventral view. Plumose setae on pleopodous rami omitted.
Figure 8. *Pseudocrangonyx daejeonensis* sp. n., paratype, male (2.7 mm). **A** antenna 1, lateral view **B** accessory flagellum of antenna 1, medial view **C** antenna 2, lateral view **D** calceolus of antenna 2, medial view **E** gnathopod 1, medial view **F** palmar margin of propodus and dactylus of gnathopod 1, medial view **G** gnathopod 2, lateral view **H** palmar margin of propodus and dactylus of gnathopod 2, lateral view.
species formed a clade (BS = 91 %, PP = 0.99) with *Pseudocrangonyx* sp. 3 inhabiting the eastern part of Shiga Prefecture, Japan. Monophyly of the present specimens of *P. daejeonensis* was fully-supported (BS = 99 %, PP = 1.0).

**Remarks.** *Pseudocrangonyx daejeonensis* is morphologically similar to *P. coreanus* in having 1) relatively small body size (smaller than 6 mm), 2) eyes completely absent, 3) carpus of gnathopod 2 without serrate robust setae on posterodistal corner, 4) outer margin or outer distal corner of pleopods 1 and 2 with setae, 5) inner basal margin of inner ramus of pleopods without bifid setae, and 6) small number of articles (less than 5) of rami of pleopods. However, the former is distinguished from the latter by the following features (features of *P. coreanus* in parentheses): 1) antenna 1 shorter (longer) than 0.4 times as long as body length, 2) antenna 2 of female without calceoli (with calceoli), 3) uropod 1 not exceeding (slightly exceeding) tip of uropod 2, and 4) outer ramus of uropod 2 without robust seta (with robust seta).

*Pseudocrangonyx daejeonensis* is also similar to *P. febras* Sidorov, 2009 and *P. gudariensis* Tomikawa and Sato in Tomikawa et al. (2016) in having 1) relatively smaller body size, 2) eyes completely absent, and 3) urosomite 1 without basal setae. However, *P. daejeonensis* is distinguished from these two species by the following features: from *P. febras* (features of *P. febras* in parentheses), 1) antenna 1 shorter than 0.4 times as long as body length (longer than 0.7 times), 2) peduncular article 2 of antenna 1 0.5 (0.7)
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Figure 10. Bayesian inference tree for 2773 bp of nuclear 28S rRNA, plus histone H3 and mitochondrial COI and 16S rRNA markers. Numbers on nodes represent bootstrap values for maximum likelihood and Bayesian posterior probabilities. Specimen numbers are shown in Table 1.

times as long as article 1, 3) palp article 2 of mandible with 3 (7) setae, 4) carpus of male gnathopod 2 without serrate robust setae on posterodistal corner (with serrate robust setae), 5) fewer articles of pleopodal rami, up to 4 (more, up to 6), 6) inner ramus of uropod 1 with 2 inner marginal robust setae (5 inner and 3 outer marginal robust setae), 7) outer ramus of uropod 1 without setae (with 2 robust setae), and 8) inner ramus of uropod 2 with inner robust seta (3–4 inner and 2–3 outer marginal robust setae); from *P. gudariensis* (features of *P. gudariensis* in parentheses), 1) basal part of inner ramus of uropod 1 with 1 slender setae (with 3 slender setae), 2) outer ramus of uropod 1 without setae (with 2 setae), 3) inner margin of inner ramus uropod 2 with 1 robust setae (with 4 robust setae), and 4) telson lobe with 1 robust seta apically (with 2 robust setae apically).

Although the phylogenetic position of *P. coreanus* remains uncertain, the results of the previous molecular phylogenetic studies (Tomikawa et al. 2016; Zhao and Hou 2017) and our phylogenetic analyses showed that *P. daejeonensis* and the two morphologically similar species, *P. febras* and *P. gudariensis*, did not form a monophyletic
lineage with large genetic divergences. Because these three species inhabit interstitial waters, not subterranean habitats, morphological similarities observed among them may reflect their similar habitat preferences.

The phylogenetic position of *P. daejeonensis* also sheds light onto the complex faunistic relationships between the *Pseudocrangonyx* species inhabiting the Japanese Archipelago and those inhabiting the Far Eastern continental area. Common ancestors of the Japanese *Pseudocrangonyx* species were considered to have migrated from the continental part to the Japanese Archipelago (Sidorov and Holsinger 2007). Previous systematic studies revealed that the *Pseudocrangonyx* amphipods distributed in northern Japan and the western tip of Honshu, Japan, i.e., *P. yezonis* and *Pseudocrangonyx* sp. 2, are phylogenetically close to the continental species (Sidorov and Holsinger 2007; Tomikawa et al. 2016; Zhao and Hou 2017). As *P. daejeonensis* formed a well-supported clade with *Pseudocrangonyx* sp. 3, which is indigenous to the central part of Honshu, their phylogenetic relationship suggested that the species diversity of the Japanese *Pseudocrangonyx* has been increased as a result of multiple continental-origins. It is also feasible that *P. daejeonensis* diverged from a common ancestor indigenous to the Japanese Archipelago. To clarify the biogeographical history of *Pseudocrangonyx* amphipods, further faunistic surveys along with molecular phylogenetic analyses are essentially needed.

The uncorrected $p$-distance of 15.0 % for COI, calculated using MEGA7.0.16 (Kumar et al. 2016) between *P. daejeonensis* and *Pseudocrangonyx* sp. 3 is equivalent to sequence divergence thresholds for discriminating amphipod species (Witt et al. 2006; Rock et al. 2007; Hou et al. 2009). The former is distinguished from the latter in having the following features (features of *Pseudocrangonyx* sp. 3 in parentheses): 1) outer distal corner of peduncle of pleopod 3 without seta (with seta), 2) each of inner and outer ramus of pleopod 3 2-articulate (3-articulate), 3) outer rami of uropods 1 and 2 without marginal robust setae (with marginal seta), and 4) robust setae on distal part of proximal article of uropod 3 short, not reaching tip of terminal article (long, exceeding tip of terminal article) (Tomikawa pers. observation).

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References

Akatsuka K, Komai T (1922) *Pseudocrangonyx*, a new genus of subterranean amphipods from Japan. Annotationes Zoologicae Japonenses 10: 119–126.

Holsinger JR (1989) Allocrangonyctidae and Pseudocrangonyctidae, two new families of Holarctic subterranean amphipod crustaceans (Gammaridea), with comments on their phylogenetic and zoogeographic relationships. Proceedings of the Biological Society of Washington 102: 947–959.

Hou Z, Li Z, Li S (2009) Identifying Chinese species of *Gammarus* (Crustacea: Amphipoda) using DNA barcoding. Current Zoology 55: 158–164.

Katoh K, Standley DM (2013) MAFFT multiple sequence alignment software version 7: improvements in performance and usability. Molecular Biology and Evolution 30: 772–780. https://doi.org/10.1093/molbev/mst010

Kumar S, Stecher G, Tamura K (2016) MEGA7: Molecular Evolutionary Genetics Analysis Version 7.0 for bigger datasets. Molecular Biology and Evolution 33: 1870–1874. https://doi.org/10.1093/molbev/msw054

Lanfear R, Calcott B, Ho SYM, Guindon S (2012) PartitionFinder: Combined selection of partitioning schemes and substitution models for phylogenetic analyses. Molecular Biology and Evolution 29: 1695–1701. https://doi.org/10.1093/molbev/mss020

Lanfear R, Frandsen PB, Wright AM, Senfeld T, Calcott B (2017) PartitionFinder 2: New methods for selecting partitioned models of evolution for molecular and morphological phylogenetic analyses. Molecular Biology and Evolution 34: 772–773. https://doi.org/10.1093/molbev/msw260

Lee CW, Park JG (2016) Distribution of meiobenthic arthropod communities in the hyporheic zone of Nakdonggang. Journal of Species Research 5: 157–165. https://doi.org/10.12651/JSR.2016.5.1.157

Rambaut A, Drummond AJ (2013) Tracer v. 1.6. http://tree.bio.ed.ac.uk/software/tracer/

Rock J, Ironside J, Potter T, Whiteley NM, Lunt DH (2007) Phylogeography and environmental diversification of a highly adaptable marine amphipod *Gammarus duebeni*. Heredity 99: 102–111. https://doi.org/10.1038/sj.hdy.6800971

Ronquist F, Teslenko M, van der Mark P, Ayres DL, Darling A, Höhna S, Larget B, Liu L, Suchard MA, Huelsenbeck JP (2012) MrBayes 3.2: Efficient Bayesian phylogenetic inference and model choice across a large model space. Systematic Biology 61: 539–542. https://doi.org/10.1093/sysbio/sys029

Sidorov DA (2009) New species of stygobiont amphipod (Crustacea: Amphipoda: Pseudocrangonyctidae) from Primorye, with description of female of *Pseudocrangonyx levanidovi* Birstein from the Khor River springs. Amurian Zoological Journal 1: 92–105.

Sidorov D, Holsinger JR (2007) *Procrangonyx stygoedincus*, a new species of subterranean amphipod (Pseudocrangonyctidae) from the Far East of Russia, with remarks on biogeographic relationships. Crustaceana 80: 417–430. https://doi.org/10.1163/156854007780440984

Stamatakis A (2014) RAxML version 8: a tool for phylogenetic analysis and post-analysis of large phylogenies. Bioinformatics 30: 1312–1313. https://doi.org/10.1093/bioinformatics/btu033
Stock JH (1974) The systematics of certain Ponto-Caspian Gammaridae (Crustacea, Amphipoda). Mitteilungen aus dem Hamburgischen Zoologischen Museum und Institut 70: 75–95.

Tomikawa K, Nakano T, Sato A, Onodera Y, Ohtaka A (2016) A molecular phylogeny of *Pseudocrangonyx* from Japan, including a new subterranean species (Crustacea, Amphipoda, Pseudocrangonyctidae). Zoosystematics and Evolution 92: 187–202. https://doi.org/10.3897/zse.92.10176

Uéno M (1934) Subterranean Crustacea from Kwantung. Annotationes Zoologicae Japonenses 14: 445–450.

Uéno M (1966) Results of the speleological survey in South Korea 1966 II. Gammarid Amphipoda found in subterranean waters of South Korea. Bulletin of the National Science Museum, Tokyo 9: 501–535.

Witt JDS, Threloff DL, Hebert PDN (2006) DNA barcoding reveals extraordinary cryptic diversity in an amphipod genus: implications for desert spring conservation. Molecular Ecology 15: 3037–3082. https://doi.org/10.1111/j.1365-294X.2006.02999.x

Zhao S, Hou Z (2017) A new subterranean species of *Pseudocrangonyx* from China with an identification key to all species of the genus (Crustacea, Amphipoda, Pseudocrangonyctidae). ZooKeys 647: 1–22. https://doi.org/10.3897/zookeys.647.11192