# A new quadrannulate species of Orobdella (Hirudinida,Arhynchobdellida, Orobdellidae) from western Honshu, Japan 

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#### Abstract

A new quadrannulate species of Orobdella Oka, 1895, Orobdella naraharaetmagarum sp. n., from the mountainous region of western Honshu, Japan is described. Orobdella naraharaetmagarum is a small species with a body length of less than 5 cm . Phylogenetic analyses using nuclear $18 S$ rRNA and histone H 3 , as well as mitochondrial cytochrome $c$ oxidase subunit $\mathrm{I}, \mathrm{tRNA}^{\mathrm{Crs}}, \mathrm{tRNA}^{\text {Mer }}, 12 \mathrm{~S}$ rRNA, $\mathrm{RRNA}{ }^{\mathrm{Val}}$, $16 S$ rRNA, $\mathrm{tRNA}{ }^{\text {teu }}$ and NADH dehydrogenase subunit 1 markers indicated that the present new species is the sister species of the quadrannulate Orobdella esulcata Nakano, 2010. Furthermore, mitochondrial DNA genealogy within $O$. naraharaetmagarum demonstrated that this new species is divided into eastern and western lineages.


## Keywords

Hirudinea, Orobdella, new species, gastroporous, molecular phylogeny, Japan

## Introduction

The terrestrial macrophagous leech genus Orobdella Oka, 1895 contains 12 species that are distributed throughout the Japanese Archipelago, Korean Peninsula, and Taiwan (Nakano 2014, Nakano and Lai 2012, Nakano and Seo 2014). These 12 species are split into three groups based on their mid-body somite annulation: seven species in the quadrannulate (four annuli) group, four in the sexannulate (six annuli) group, and one octannulate (eight annuli) species.

Orobdella leeches had been considered large species, with body lengths reaching to 10 cm or greater (Sawyer 1986). In recent years, however, small mature leeches belonging to this genus have been discovered in Japan and described as new species: Orobdella koikei Nakano, 2012b from Hokkaido, and Orobdella masaakikuroiwai Nakano, 2014 from central Honshu. The bodies of mature individuals of these two species are shorter than 4 cm . Both species possess mid-body somites that are quadrannulate. Nakano (2014) suggested that differences in the body lengths of mature leeches might enable more than one species of Orobdella to coexist in the same region.

Additional small Orobdella leeches were collected from Chugoku District, western Honshu, Japan. The body lengths of the specimens were less than 5 cm . Nevertheless, a few individuals were regarded as mature leeches because they possessed an obvious clitellum. These specimens are described here as a new species. In addition, the phylogenetic position of this new species was estimated using nuclear 18S rRNA and histone H3, as well as mitochondrial cytochrome $c$ oxidase subunit $\mathrm{I}, \mathrm{tRNA}{ }^{\mathrm{Cys}}$, $\mathrm{tRNA}^{\mathrm{Met}}, 12 \mathrm{~S}$ rRNA, tRNA ${ }^{\text {Val }}, 16 \mathrm{~S}$ rRNA, tRNA ${ }^{\text {Leu }}$, and NADH dehydrogenase subunit 1 sequence data.

## Materials and methods

## Sampling and morphological examination

Leeches were collected from five localities in Chugoku district, western Honshu, Japan (Fig. 1). When possible, elevation and geographical coordinates for localities were obtained using a Garmin eTrex GPS unit.

Almost all of the specimens were relaxed by the gradual addition of absolute ethanol (EtOH) to freshwater. For DNA extraction, botryoidal tissue was removed from the posterior part of the body around the caudal sucker of every specimen, and then preserved in absolute EtOH. The remainder of the body was fixed in $10 \%$ formalin and preserved in $70 \% \mathrm{EtOH}$. Four measurements were taken: body length (BL) from the anterior margin of the oral sucker to the posterior margin of the caudal sucker, maximum body width (BW), caudal sucker length (CL) from the anterior to the posterior margin of the sucker and caudal sucker width (CW) from the right to the left margin of the sucker. Examination, dissection, and drawing of the specimens were conducted using a stereoscopic microscope with a drawing tube (Leica M125). Specimens used in this study have been deposited in the Zoological Collection of Kyoto University (KUZ).


Figure I. Map showing the collection localities of the specimens examined in this study. Open circle (5) indicates the type locality of the new species, Orobdella naraharaetmagarum sp. n., and closed circles (I-4) indicate additional localities.

The numbering convention is based on Moore (1927): body somites are denoted by Roman numerals, and the annuli in each somite are given alphanumeric designations.

## PCR and DNA sequencing

The extraction of genomic DNA from botryoidal tissues preserved in absolute EtOH followed Nakano (2012b). Primer sets for the PCR and cycle sequencing (CS) reactions used in this study were as follows: for 18 S rRNA, A and L (PCR and CS), C and Y (PCR and CS), as well as O and B (PCR and CS) (Apakupakul et al. 1999); for histone H3 (H3), H3aF and H3bR (PCR and CS) (Colgan et al. 1998); for cytochrome $c$ oxidase subunit I (COI), LCO 1490 and HCO 2198 (PCR and CS) (Folmer et al. 1994), and LCO-in (Nakano 2012b) and HCO-outout (Nakano 2012a) (PCR and CS), or LCOinerpo2 (5'-GCTATTACAATATTACTTACAGATCG-3'; this study) and HCO-out (Nakano 2012b) (PCR and CS); for $\mathrm{tRNA}^{\mathrm{Cys}}$, $\mathrm{tRNA}^{\mathrm{Met}}, 12 \mathrm{~S}$ rRNA, $\mathrm{tRNA}^{\mathrm{Val}}$ and 16 S rRNA (tRNA ${ }^{\text {Cys }}-16 \mathrm{~S}$ ), 12 SA-out and $12 \mathrm{SB}-\mathrm{in}$ (PCR and CS), and 12SA-in and 12SBout (Nakano 2012b) or 12SB-outin (5'-AAAGGTACGAATATATTTAC-3'; this study) (PCR and CS); for $\mathrm{tRNA}{ }^{\text {Leu }}$ and NADH dehydrogenase subunit 1 (ND1) ( $\mathrm{tRNA}{ }^{\text {Leu }}-$

ND1), LDN3000 and HND1932 (PCR and CS) (Light and Siddall 1999). The PCR reactions and DNA sequencing were performed using the modified method mentioned in Nakano (2012a). The PCR reactions were performed using a GeneAmp PCR System 2700 and a GeneAmp PCR System 9700 (Applied Biosystems) as well as a T100 Thermal Cycler (Bio-Rad). The PCR mixtures were heated to $94^{\circ} \mathrm{C}$ for 5 min , followed by 35 cycles at $94^{\circ} \mathrm{C}(10$ s each $), 52^{\circ} \mathrm{C}$ for 18 S and $\mathrm{H} 3,60^{\circ} \mathrm{C}$, and $44^{\circ} \mathrm{C}$, respectively, for the anterior, and posterior parts of $\mathrm{tRNA}^{\mathrm{Cys}}-16 \mathrm{~S}$ or $42^{\circ} \mathrm{C}$ for COI and $t R N A^{\text {Leu }}-\mathrm{ND} 1$ (20 s), and $72^{\circ} \mathrm{C}(42 \mathrm{~s} \mathrm{each})$, and a final extension at $72^{\circ} \mathrm{C}$ for 6 min . The sequencing mixtures were heated $96^{\circ} \mathrm{C}$ for 2 min , followed by 40 cycles at $96^{\circ} \mathrm{C}(10 \mathrm{~s}$ each $), 50^{\circ} \mathrm{C}$ ( 5 s each) and $60{ }^{\circ} \mathrm{C}$ ( 48 s each). The obtained sequences were edited using DNA BASER (Heracle Biosoft S.R.L.). The DNA sequences listed in Table 1 were newly obtained in this study and were deposited with the International Nucleotide Sequence Database Collaboration (INSDC) through the DNA Data Bank of Japan (DDBJ).

## Molecular phylogenetic and genetic distance analyses

Eighty published sequences were obtained from the INSDC for use in molecular phylogenetic analyses (Table 1). In addition to 12 known Orobdella species, the following four erpobdelliform species were used as outgroup taxa: Erpobdella japonica Pawłowski, 1962 (Erpobdellidae), Gastrostomobdella monticola Moore, 1929 (Gastrostomobdellidae), Mimobdella japonica Blanchard, 1897, and Odontobdella blanchardi (Oka, 1910) (both Salifidae).

The phylogenetic position of the newly identified Orobdella species within the genus was estimated based on $18 \mathrm{~S}, \mathrm{H} 3, \mathrm{COI}, \mathrm{tRNA}^{\mathrm{Cys}}-16 \mathrm{~S}$, and $\mathrm{tRNA}{ }^{\text {Leu }}-\mathrm{ND} 1$ sequences. The alignments of H 3 and COI were trivial, as no indels were observed. 18 S , $\mathrm{tRNA}{ }^{\mathrm{Cys}}-16 \mathrm{~S}$, and $\mathrm{tRNA}{ }^{\text {Leu }}-\mathrm{ND} 1$ were aligned using MAFFT v. 7.245 L-INS-i (Katoh and Standley 2013). The lengths of the 18S, H3, COI, tRNA ${ }^{\text {Cys }}-16 S$, and tRNA ${ }^{\text {Leu }}-N D 1$ sequences were $1,844,328,1,267,1,120$, and 633 bp , respectively. The concatenated sequences yielded $5,192 \mathrm{bp}$ of aligned positions.

Phylogenetic trees were constructed using maximum likelihood (ML) and Bayesian inference (BI). ML phylogenies were constructed using RAxML v. 8.1.5 (Stamatakis 2014) with the substitution model set as GTRCAT, immediately after nonparametric bootstrapping (Felsenstein 1985) conducted with 1,000 replicates. The best-fit partitioning scheme for the ML analyses was identified with the Akaike information criterion (Akaike 1974) using PartitionFinder v. 1.1.1 (Lanfear et al. 2012) with the "greedy" algorithm: 18S/the $1^{\text {st }}$ and $2^{\text {nd }}$ positions of $\mathrm{H} 3 /$ the $3^{\text {rd }}$ position of $\mathrm{H} 3 /$ the $1^{\text {st }}$ position of $\mathrm{COI} /$ the $2^{\text {nd }}$ position of $\mathrm{COI} /$ the $3^{\text {rd }}$ positions of COI and $\mathrm{ND} 1 /$ the $1^{\text {st }}$ position of ND1/the $2^{\text {nd }}$ position of ND2/12S/16S/tRNA ${ }^{\text {Cys }}, \mathrm{tRNA}^{\mathrm{Met}}, \mathrm{tRNA}^{\mathrm{Val}}$ and $\mathrm{tRNA}{ }^{\text {Leu }}$.

BI and Bayesian posterior probabilities (BPPs) were estimated using MrBayes v . 3.2.5 (Ronquist et al. 2012). The best-fit partition scheme and models for each partition were selected based on the Bayesian information criterion (Schwarz 1978) using PartitionFinder with the "greedy" algorithm: for 18 S and the $1^{\text {st }}$ position of H3, $\mathrm{K} 80+\mathrm{I}$; for the $2^{\text {nd }}$ position of H 3 , JC69; for the $3^{\text {rd }}$ position of H 3 , HKY85; for the
Table I. Samples used for the phylogenetic analyses. The information on the vouchers is accompanied by the collection locality numbers for Orobdella naraharaetmagarum sp. n. (see Fig. 1) and the INSDC accession numbers. Sequences marked with an asterisk were obtained for the first time in the present study. Acronyms: KUZ, the Zoological Collection of Kyoto University; UNIMAS, the Universiti Malaysia Sarawak.

| Species | Voucher (locality number) | 18S | Histone H3 | COI | tRNA ${ }^{\text {Cys }} \mathbf{- 1 6 S}$ | tRNA ${ }^{\text {Leu }}$-ND1 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Orobdella naraharaetmagarum sp. n. | KUZ Z1360 (4) |  |  | LC087131* | LC087130* | LC087132* |
| Orobdella naraharaetmagarum sp. n. | KUZ Z1380 (2) |  |  | LC087134* | LC087133* | LC087135* |
| Orobdella naraharaetmagarum sp. n. | KUZ Z1535 (1) |  |  | LC087137* | LC087136* | LC087138* |
| Orobdella naraharaetmagarum sp. n. | KUZ Z1582 Paratype (5) |  |  | LC087140* | LC087139* | LC087141* |
| Orobdella naraharaetmagarum sp. n. | KUZ Z1652 Holotype (5) | LC087143* | LC087145* | LC087144* | LC087142* | LC087146* |
| Orobdella naraharaetmagarum sp. n. | KUZ Z1654 Paratype (5) |  |  | LC087148* | LC087147* | LC087149* |
| Orobdella naraharaetmagarum sp. n. | KUZ Z1655 (3) |  |  | LC087151* | LC087150* | LC087152* |
| Orobdella dolichopharynx Nakano, 2011b | KUZ Z120 Holotype | AB663665 | AB698876 | AB679680 | AB679681 | AB828558 |
| Orobdella esulcata Nakano, 2010 | KUZ Z29 Holotype | AB663655 | AB698873 | AB679664 | AB679665 | AB828555 |
| Orobdella ijimai Oka, 1895 | KUZ Z110 Topotype | AB663659 | AB698877 | AB679672 | AB679673 | AB828559 |
| Orobdella kawakatsuorum Richardson, 1975 | KUZ Z167 Topotype | AB663661 | AB698878 | AB679704 | AB679705 | AB828561 |
| Orobdella ketagalan Nakano \& Lai, 2012 | KUZ Z208 Holotype | AB704785 | AB704786 | AB704787 | AB828582 | AB828563 |
| Orobdella koikei Nakano, 2012b | KUZ Z156 Holotype | AB698883 | AB698882 | AB679688 | AB679689 | AB828560 |
| Orobdella masaakikuroiwai Nakano, 2014 | KUZ Z694 Holotype | AB938003 | AB938013 | AB938006 | AB937997 | AB938016 |
| Orobdella mononoke Nakano, 2012a | KUZ Z224 Holotype | AB698868 | AB698869 | AB698866 | AB698867 | AB828564 |
| Orobdella octonaria Oka, 1895 | KUZ Z181 Topotype | AB698870 | AB698871 | AB679708 | AB679709 | AB828562 |
| Orobdella shimadae Nakano, 2011b | KUZ Z128 Holotype | AB663663 | AB698875 | AB679676 | AB679677 | AB828557 |
| Orobdella tsushimensis Nakano, 2011a | KUZ Z134 Holotype | AB663653 | AB698872 | AB679662 | AB679663 | AB828554 |
| Orobdella whitmani Oka, 1895 | KUZ Z45 Topotype | AB663657 | AB698874 | AB679668 | AB679669 | AB828556 |
| Erpobdella japonica Pawłowski, 1962 | KUZ Z178 | AB663648 | AB698879 | AB679654 | AB679655 | AB828542 |
| Gastrostomobdella monticola Moore, 1929 | UNIMAS/A3/BH01/10 | AB663649 | AB698880 | AB679656 | AB679657 | AB828543 |
| Mimobdella japonica Blanchard, 1897 | KUZ Z179 | AB663650 | AB698881 | AB679658 | AB679659 | AB828544 |
| Odontobdella blanchardi (Oka, 1910) | KUZ Z180 | AB663651 | AB938012 | AB938004 | AB937995 | AB938014 |

$1^{\text {st }}$ position of COI, GTR+G; for the $2^{\text {nd }}$ positions of COI and ND1, HKY85+I; for the $3^{\text {rd }}$ positions of COI and ND1 plus 16 S , HKY85 $+\mathrm{I}+\mathrm{G}$; and for the $1^{\text {st }}$ position of ND1, and $12 S, \mathrm{tRNA}^{\mathrm{Cys}}$, $\mathrm{tRNA} A^{\mathrm{Met}}$, $\mathrm{tRNA}{ }^{\mathrm{Val}}$ and $\mathrm{tRNA}^{\text {Leu }}$, GTR $+\mathrm{I}+\mathrm{G}$. Two independent runs of four Markov chains were conducted for 12 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 2009) and the first 30,001 trees were discarded based on these results.

The phylogenetic relationships within the available Orobdella materials were estimated based on sequences of COI, $t \mathrm{RNA}^{\mathrm{Cys}}-16 \mathrm{~S}$ and $t \mathrm{RNA}^{\text {Leu }}-\mathrm{ND} 1 . \mathrm{tRNA}^{\mathrm{Cys}}-16 \mathrm{~S}$ and tRNA ${ }^{\text {Leu }}-N D 1$ were aligned using MAFFT L-INS-i. The lengths of the COI, $\mathrm{tRNA}^{\mathrm{Cys}}{ }_{-}$ $16 S$, and $t R N A{ }^{\text {Leu }}-N D 1$ sequences were $1,267,634$, and $1,107 \mathrm{bp}$, respectively. The concatenated sequences yielded $3,008 \mathrm{bp}$ of aligned positions. ML phylogenies were constructed in RAxML with the substitution model set as GTRCAT, immediately after nonparametric bootstrapping conducted with 1,000 replicates. The best-fit partitioning scheme was identified with the Akaike information criterion using PartitionFinder with the "greedy" algorithm: the $1^{\text {st }}$ position of $\mathrm{COI} /$ the $2^{\text {nd }}$ positions of COI and $\mathrm{ND} 1 /$ the $3^{\text {rd }}$ positions of COI and ND1/the $2^{\text {nd }}$ position of ND $1 /$ the $1^{\text {st }}$ position of ND1/tRNA${ }^{\text {Cys }}$, , $\mathrm{RNA}^{\text {Met }}, \mathrm{tRNA}^{\mathrm{Val}}$, $\mathrm{tRNA}{ }^{\text {Leu }} / 12 \mathrm{~S} / 16 \mathrm{~S}$. BI and BPPs were estimated using MrBayes. 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 the $1^{\text {st }}$ positions of COI and ND1, GTR $+\mathrm{I}+\mathrm{G}$; for the $2^{\text {nd }}$ positions of COI and ND1, F81+I; for the $3^{\text {rd }}$ positions of COI and ND1 plus $16 \mathrm{~S}, \mathrm{HKY}+\mathrm{G}$; tRNA ${ }^{\mathrm{Cys}}$, $\mathrm{tRNA}^{\text {Met }}$, 12 S , $\mathrm{tRNA}{ }^{\text {Val }}$ and $\mathrm{tRNA}^{\text {Leu }}$, GTR $+\mathrm{I}+\mathrm{G}$. Two independent runs of four Markov chains were conducted for 10 million generations and the tree was sampled every 100 generations. The parameter estimates and convergence were checked using Traced, and the first 25,001 trees were discarded based on these results.

Nodes with bootstrap support (BS) values higher than $70 \%$ were considered sufficiently resolved (Hillis and Bull 1993). Nodes with BPPs higher than $95 \%$ were considered statistically significant (Leaché and Reeder 2002).

Pairwise comparisons of uncorrected $p$-distances for seven COI sequences (1,266 bp) obtained from specimens of the studied species and Orobdella esulcata Nakano, 2010 were calculated using MEGA6.06 (Tamura et al. 2013).

## Taxonomy

Family Orobdellidae Nakano et al., 2012
http://zoobank.org/5F5BABE8-BD26-4FC7-9593-F73E62E26122

Genus Orobdella Oka, 1895
http://zoobank.org/FA8333ED-8C17-41FD-AFC1-62A4F98D4AC1

## Orobdella narabaraetmagarum sp. n.

http://zoobank.org/5A831984-50F6-433A-A058-ED2ECFF2DFDC
Figs 2-5

Diagnosis. Body length of mature individual less than 5 cm . Somite IV uniannulate, somites VIII-XXV quadrannulate. Male gonopore in middle of XI b6, female gonopore in middle of XIII a1, behind gastropore, gonopores separated by $1 / 2+4+1 / 2$ annuli. Clitellum in XI b5 to XIII a2. Pharynx reaching to XIII b5/b6-XIII/XIV. Gastropore conspicuous, in middle of XIII a1. Gastroporal duct bulbous, slightly winding at junction with gastropore. Paired epididymides in XIV b6-XV b5 to XVIII b6-XX a2/b5, occupying 16-20 annuli (four to five somites). Atrial cornua developed, ellipsoid or ovate.

Type materials. Holotype. KUZ Z1652, dissected, collected from under a rock along a mountain trail at Mt. Kanmuriyama, Hatsukaichi, Hiroshima Pref., Japan ( $34.47325^{\circ}, 132.10362^{\circ}$; Elev. 757 m; locality number 5), by TN on 25 April 2015. Paratypes. Two specimens from near the type locality, along a forest road, "Japan National Route 488", Hatsukaichi, both dissected: KUZZ1582, under a rock ( $34.50118^{\circ}$, $132.08933^{\circ}$; Elev. 790 m ; locality number 5), by Yoshiko Yamane on 10 August 2014, and KUZ Z1654, under a rotten tree (34.50182 ${ }^{\circ}$, $132.08961^{\circ}$; Elev. 791 m ; locality number 5), by TN on 16 June 2015. For locality numbers, see Fig. 1.

Additional materials. In total four specimens were examined, all dissected. Three specimens collected from Hiroshima Pref., Japan: KUZ Z1360, from Hosomi, Kitahiroshima ( $34.685^{\circ}, 132.292^{\circ}$; Elev. 470 m ; locality number 4), by Yukiko Narahara on 9 July 2011; KUZ Z1380, from Mt. Azumayama, Hiwacho, Shobara (35.0639º, $133.0268^{\circ}$; Elev. 1010 m; locality number 2), by Ayane Maga on 3 October 2011; and KUZ Z1655, from under a rock along a mountain trail at Mt. Izaiyama, Hiwacho-Mitsugaichi, Shobara $\left(35.00143^{\circ}, 133.04640^{\circ}\right.$; Elev. 906 m ; locality number 3), by TN on 17 June 2015. KUZ Z1535, collected from under a rock along a mountain trail at Mt. Iimoriyama, Noigura, Kotoura, Tottori Pref., Japan (35.37603, $133.59953^{\circ}$; Elev. 619 m ; locality number 1), by TN on 11 December 2013. For locality numbers, see Fig. 1.


Figure 2. Orobdella naraharaetmagarum sp. n., holotype, KUZ Z1652. A Dorsal and B ventral views. Scale bar: 5 mm .

$a 1=a 2=a 3$ (Fig. 3A, B). Somites VIII - XXV quadrannulate, $a 1=a 2=b 5=b 6$ (Fig. 3A-E). Somite XXVI dorsally triannulate, $a 1>a 2<a 3$, $a 3$ with slight furrow; ventrally biannulate, $(a 1+a 2)>a 3$, $(a 1+a 2)$ with slight furrow; $(a 1+a 2)$ being ventrally last complete annulus (Fig. 3C, D). Somite XXVII uniannulate with slight dorsal furrow; anus behind it with no post-anal annulus (Fig. 3C).

Male gonopore in middle of XI b6 (Fig. 3E). Female gonopore slightly anterior to middle of XIII a1, inconspicuous, located posterior to gastropore (Fig. 3E, F). Gonopores separated by $1 / 2+4+1 / 2$ annuli (Fig. 3E).

Anterior ganglionic mass in VI a2 and a3. Ganglion VII in a2. Ganglia VIII-X, of each somite, in a2. Ganglia XI and XII, of each somite, in a2 (Fig. 4A). Ganglion XIII in a2 and b5 (Fig. 4A). Ganglion XIV in a2 (Fig. 4A). Ganglia XV-XX, of each somite, in a1 and a2 (Fig. 4A). Ganglia XXI-XXIII, of each somite, in a2. Ganglion XXIV in a1. Ganglion XXV in b6 of XXIV and al of XXV. Ganglion XXVI in b5 and b 6 of XXV. Posterior ganglionic mass in $(\mathrm{a} 1+\mathrm{a} 2)$ of XXVI.

Eyes in three pairs, first pair dorsally on posterior margin of II, second and third pairs dorsolaterally on posterior margin of $\mathrm{V}(\mathrm{a} 1+\mathrm{a} 2)$ (Fig. 3A). Papillae numerous, minute, hardly visible, one row on every annulus.

Nephridiopores in 17 pairs, one each situated ventrally at posterior margin of al of each somite in VIII-XXIV (Fig. 3B, D, E).

Pharynx agnathous, euthylaematous, reaching to XIII b5/b6 (Fig. 3G). Crop tubular, acecate, reaching to XIX b5/b6. Gastropore conspicuous, ventral, slightly anterior to middle of XIII a1 (Fig. 3E, F). Gastroporal duct bulbous, slightly winding at junction with gastropore, joining with crop in XIV al (Fig. 3G). Intestine tubular, acecate, reaching to XXIV a1. Rectum tubular, thin-walled, straight.

Testisacs multiple (Fig. 4A); on right side, in XVIII b6 to XXIV a2, in total app. 28 testisacs, 1 in XVIII, 3 in XIX, 5 in XX, 4 in XXI, 6 in XXII, 7 in XXIII, 2 in XXIV; on left side, in XIX a1 to XXV a2, in total app. 27 testisacs, 5 in XIX, 6 in XX, 4 in XXI, 4 in XXII, 4 in XXIII, 3 in XXIV, 1 in XXV. Paired epididymides; right epididymis in XIV b6 to XVIII b6, occupying 17 annuli; left epididymis in XIV/XV to XVIII b6, occupying 16 annuli (Fig. 4A). Ejaculatory bulbs absent. Paired ejaculatory ducts; right duct in XI a2/b5 to XIV b6; left duct in XI a2/b5 to XIV/XV; coiled in position posterior to ovisacs; each duct crossing ventrally beneath each ovisac, then curved in position anterior to ovisacs; each widening from respective junction with epididymis, narrowing at junction with atrial cornua, then turning sharply inward toward atrial cornua without pre-atrial loop (Fig. 4A). Pair of muscular atrial cornua ellipsoid, in XI b5 and b6 (Fig. 4A-D). Atrium short, muscular, globular in XI b5 and b6 (Fig. 4B-D). Penis sheath and penis absent.

Paired ovisacs globular; right ovisac in XIII a2 and b5; left ovisac in XIII a1 and a2 (Fig. 4A, E). Oviducts, thin-walled, left oviduct crossing ventrally beneath nerve cord; both oviducts converging into common oviduct in XIII a2 (Fig. 4A, E). Common oviduct thin-walled, short, directly descending to female gonopore (Fig. 4E).

Variations. BL 48.2 (KUZ Z1654)-33.0 (KUZ Z1360) mm, BW 4.4 (KUZ Z1535)-2.7 (KUZ Z1380) mm, CL 2.5 (KUZ Z1654)-1.2 (KUZ Z1380) mm, CW


Figure 4. Orobdella naraharaetmagarum sp. n., holotype, KUZ Z1652. A Dorsal view of reproductive system including ventral nervous system. B Dorsal (including positions of ganglia XI and XII), C lateral, and $\mathbf{D}$ ventral views of male atrium. E Dorsal view of female reproductive system including position of ganglion XIII. Scale bars: $2 \mathrm{~mm}(\mathbf{A}), 0.5 \mathrm{~mm}(\mathbf{B}-\mathbf{D})$ and $0.25 \mathrm{~mm}(\mathbf{E})$. Abbreviations: ac, atrial cornua; at, atrium; cod, common oviduct; ed, ejaculatory duct; ep, epididymis; gp, gastropore; od, oviduct; ov, ovisac; and ts, testisac.
3.0 (KUZ Z1654)-1.8 (KUZ Z1360, Z1380) mm. Somite XXVI triannulate, a1 = a2 < a3 (KUZ Z1380, Z1582); KUZ Z1360, Z1535, Z1654, Z1655, a3 with slight dorsal furrow. Somite XXVII uniannulate or biannulate (KUZ Z1360). Male gonopore generally in middle of XI b6, rarely slightly anterior or posterior to middle of XI b6. Female gonopore in middle of XIII a1, slightly anterior or posterior to middle of XIII a1. X b5 and XIII a2, respectively, being first and last annuli of clitellum. Eyes generally three pairs; KUZ Z1654, Z1655, first pair dorsally on anterior margin of III; KUZ Z1582, multiple eyes detected, one eye on left dorsal of II/III, one small eyespot on right dorsal of III, one small eyespot on left dorsal of III/IV, one small eyespot on right dorsal of IV, and two pairs of eyes dorsolaterally on posterior margin of V (al + a2). Pharynx reaching to XIII b5/b6-XIII/XIV. Crop reaching to XIX b5/b6-XX a1. Gastropore in middle of XIII a1, slightly anterior or posterior to middle of XIII a1. Gastroporal duct generally bulbous; KUZ Z1360, Z1582, tubular. Intestine reaching to XXIII/XXIV-XXIV/XXV. Testisacs multiple; right side app. 11-24 sacs in XIX b6XX b5 to XXIV b5-b6; left side app. 11-23 sacs in XIX b6-XX a2 to XXIII b6-XXV a1. Paired epididymides; right epididymis in XV a1-XV b5 to XIX a2-XX a2/b5, occupying 17-20 annuli; left epididymis in XV a1-XV b5 to XIX b5-XX a2, occupying 17-20 annuli. Paired ejaculatory ducts, curved, loosely curved, or straight in position posterior to ovisacs. Atrial cornua ovate, fusiform, or ellipsoid in XI b5 and b6; KUZ Z1535 in XI b5-XII a. Atrium generally in XI b5 and b6; KUZ Z1535, Z1582 in XI b6. Paired ovisacs generally in XIII a2 and b5; KUZ Z1535, undeveloped, in XIII a2. Right or left oviduct crossing ventrally beneath nerve cord; KUZ Z1380 both oviducts converging into common oviduct in XIII a1/a2.

Coloration. In life, dorsal surface bluish gray (Fig. 5A, B), or gray; ventral surface reddish white or ash gray; clitellum, when obvious, whitish gray (Fig. 5B). Color faded in preservative; KUZ Z1535 with one dorsal black line from VIII b5 to XXVI a2.

Distribution. This species was primarily collected in Hiroshima Prefecture in Chugoku District, Honshu, Japan, and in Tottori Prefecture. The lowest elevation among the localities was 470 m , and the highest was 1010 m . The locality data for this species suggested that it is distributed in mountainous regions in Chugoku District, Honshu, Japan.

Natural history. This species was generally found curled up under rocks or rotten trees in moist mountainous habitats (Fig. 5C, D). Oligochaete worms were sometimes observed in the digestive tract during specimen dissection. Therefore, this species is an earthworm-eater, as are the other known Orobdella leeches.

A mature leech with an obvious clitellum was collected on 16 June. Moreover, the holotype, which appeared to have a clitellum (Fig. 5A), was collected on 25 April. These findings suggest that the reproductive season of the new species begins in May and then continues at least to mid-to-late June.

Remarks. The new species unambiguously belongs to Orobdella as it has all the generic diagnostic characteristics defined in Nakano et al. (2012): post-anal annulus absent; pharynx agnathous, euthylaematous; gastropore in XIII; gastroporal duct lying on female organ; gonopores separated by more than one full somite; testisacs multiple;


Figure 5. Orobdella naraharaetmagarum sp. n., holotype, KUZ Z1652 (A, D) and paratype, KUZ Z1654 (B). Habitat of holotype (C). A and B Dorsal views of live animals. D Live animal found curled up under a stone at the type locality. Scale bars: 5 mm .
male atrium in XI without penis sheath and penis; ovisacs globular in XIII; female median reproductive system essentially lacking.

The specimens were small (up to 48 mm ). However, one leech (KUZ Z1654) was determined to be mature because it possessed an obvious clitellum. The holotype, which had a body length of 40 mm , also possessed a slightly developed clitellum and developed testisacs. Two specimens (KUZ Z1360, Z1582) possessed a tubular gastroporal duct. This tubular gastroporal duct was thought to be observed in immature individuals because these two specimens had undeveloped, undetectable testisacs.

Taxonomic studies (Nakano 2010, 2012b, 2014, Nakano and Gongalsky 2014, Nakano and Lai 2012, Nakano and Seo 2014) indicate that the new species differs from the seven other quadrannulate species (i.e., O. esulcata, O. kawakatsuorum Richardson, 1975, O. ketagalan Nakano \& Lai, 2012, O. koikei, O. masaakikuroiwai, O. tsushimensis Nakano, 2011a, and O. whitmani Oka, 1895) by the following combination of characteristics (Table 2): body length less than 5 cm , IV uniannulate, gonopores separated by $1 / 2+4+1 / 2$ annuli, XXV quadrannulate, gastroporal duct bulbous, epididymides in XIV to XX, atrial cornua developed, ellipsoid or ovate. Among the above seven quadrannulate species, two species, $O$. koikei and $O$. masaakikuroiwai, are known to have body lengths shorter than 5 cm . Orobdella naraharaetmagarum can be distinguished from these two species by the annulation of XXV and the length of the epididymides.

The new species is distinguishable from the four sexannulate species $O$. dolichopharynx Nakano, 2011b, O. ijimai Oka, 1895, O. mononoke Nakano, 2012a, and O. shimadae Nakano, 2011b and the octannulate species O. octonaria Oka, 1895, since O. naraharaetmagarum possesses mid-body somites that are quadrannulate.
Table 2. Comparisons of morphological characters between Orobdella naraharaetmagarum $\mathrm{sp} . \mathrm{n}$. and seven quadrannulate congeneric species.

| Character | O. narabaraetmagarum sp. n. | O. esulcata <br> Nakano, 2010 | O. kawakatsuorum Richardson, 1975 | O. ketagalan Nakano \& Lai, 2012 | $\begin{gathered} \text { O. koikei } \\ \text { Nakano, 2012b } \end{gathered}$ | O. masaakikuroivai Nakano, 2014 | O. tsushimensis Nakano, 2011a | O. whitmani Oka, 1895 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Body length of mature leech | less than 5 cm | up to approx. 10 cm | up to approx. 10 cm | $\begin{array}{\|c} \hline \text { up to approx. } \\ 10 \mathrm{~cm} \end{array}$ | less than 4 cm | less than 4 cm | up to approx. 10 cm | up to approx. 10 cm |
| $\begin{aligned} & \text { Annulation } \\ & \text { of IV } \end{aligned}$ | uniannulate | uniannulate | biannulate | uniannulate | uniannulate | uniannulate | uniannulate | uni- or <br> biannulate |
| Number of annuli between gonopores | $1 / 2+4+1 / 2$ | $2 / 3+4+1 / 3$ | 6 | $1 / 2+4+1 / 2$ | $1 / 2+4+1 / 2$ | $1 / 2+4+1 / 2$ | $1 / 2+5$ | $1 / 2+4+1 / 2$ |
| $\begin{gathered} \text { Annulation of } \\ \text { XXV } \\ \hline \end{gathered}$ | quadrannulate | quadrannulate | quadrannulate | quadrannulate | triannulate | quadrannulate | quadrannulate | quadrannulate |
| Gastroporal duct | bulbous | tubular, but bulbous at junction with gastropore | simple tubular | simple tubular | bulbous | bulbous | bulbous | bulbous |
| Epididymides | $\begin{gathered} \text { XV (posterior of XIV) } \\ \text { to XX } \\ \hline \end{gathered}$ | XVI to XX | XVI to XVII | absent | XV to XX | XVI to XVIII | XVII to XIX | XVI to XVIII |
| Atrial cornua | developed, ellipsoid or ovate | developed, ovate | undeveloped | undeveloped | developed, ovate | developed, ovate | developed, ovate | developed, ovate |

## Molecular phylogenies and genetic distances

The BI tree (Fig. 6) for estimating the phylogenetic position of the new species had an identical topology to that of the ML tree ( $\ln L=-24617.61$; not shown). The monophyly of Orobdella naraharaetmagarum and $O$. esulcata was strongly supported ( $\mathrm{BS}=$ $100 \%$, BPP = 1.0).

The ML tree ( $\ln L=-15057.02$ ) (Fig. 7) used to construct the phylogenetic relationships of the new species had an almost identical topology to that of the BI tree (not shown). The monophyly of the specimens identified as $O$. naraharaetmagarum was recovered $(\mathrm{BS}=100 \%, \mathrm{BPP}=1.0)$. This clade was divided into two subclades (hereafter lineages 1 and 2). The monophyly of lineage 1 was strongly supported ( $\mathrm{BS}=$ $99 \%$, BPP $=1.0$ ). Lineage 1 comprised three specimens: KUZ Z1380 (locality number 2 in Fig. 1), Z1535 (locality number 1), and Z1655 (locality number 3). KUZ Z1380 and Z1655 formed a monophyletic group $(\mathrm{BS}=99 \%, \mathrm{BPP}=0.99)$. The monophyletic lineage $2(\mathrm{BS}=100, \mathrm{BPP}=1.0)$ contained four individuals: KUZ Z1380 (locality number 4), Z1582, Z1650, and Z1652 (locality number 5). In the ML phylogeny, the three specimens KUZ Z1582, Z1650, and Z1652 formed a monophyletic lineage, although this relationship was not supported ( $\mathrm{BS}=33 \%$ ). The monophyly of KUZ Z1582 and Z1652 was also not fully supported $(B S=63 \%, B P P=0.76)$.

The pairwise COI uncorrected $p$-distance within $O$. naraharaetmagarum was $0.6-4.7 \%($ mean $=3.3 \%)($ Table 3). The genetic distance between lineages 1 and 2


Figure 6. Bayesian inference tree for $5,192 \mathrm{bp}$ of nuclear 18 S rRNA and histone H 3 and mitochondrial COI, tRNA ${ }^{\text {Cys }}$, tRNA ${ }^{\text {Met }}, 12 \mathrm{~S}$ rRNA, $\mathrm{tRNA}^{\mathrm{Val}}, 16 \mathrm{~S}$ rRNA, $\mathrm{tRNA}{ }^{\text {Leu }}$ and ND1 markers. Numbers on nodes represent bootstrap values for maximum likelihood and Bayesian posterior probabilities. A species name of Orobdella in red indicates a quadrannulate species; in green, sexannulate; and in blue, octannulate.


Figure 7. Maximum likelihood tree ( $\ln L=-15057.02$ ) for 3,008 bp of mitochondrial COI, tRNA ${ }^{\mathrm{Cys}}$, tRNA ${ }^{\text {Met }}, 12 \mathrm{~S}$ rRNA, tRNA ${ }^{\text {Val }}, 16 \mathrm{~S}$ rRNA, tRNA ${ }^{\text {Leu }}$ and ND1 markers. Numbers on nodes represent bootstrap values for maximum likelihood and Bayesian posterior probabilities.

Table 3. Uncorrected $p$-distances for the 1266 bp for the COI sequences of Orobdella naraharaetmagar$u m \mathrm{sp} . \mathrm{n}$. specimens, with associated collection locality numbers (see Fig. 1).

| Specimen (locality number) | $\mathbf{1}$ | $\mathbf{2}$ | $\mathbf{3}$ | $\mathbf{4}$ | $\mathbf{5}$ | $\mathbf{6}$ | $\mathbf{7}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1: KUZ Z1535 (1) |  |  |  |  |  |  |  |
| 2: KUZ Z1380 (2) | 0.042 |  |  |  |  |  |  |
| 3: KUZ Z1655 (3) | 0.041 | 0.031 |  |  |  |  |  |
| 4: KUZ Z1360 (4) | 0.045 | 0.043 | 0.041 |  |  |  |  |
| 5: KUZ Z1654 (5) | 0.042 | 0.042 | 0.042 | 0.011 |  |  |  |
| 6: KUZ Z1582 (5) | 0.047 | 0.043 | 0.042 | 0.014 | 0.009 |  |  |
| 7: KUZ Z1652 (5) | 0.047 | 0.046 | 0.044 | 0.013 | 0.010 | 0.006 |  |

was $4.1-4.7 \%$ (mean $=4.4 \%$ ). The genetic divergences within lineages 1 and 2 were $3.1-4.2 \%$ (mean $=3.8 \%$ ) and $0.6-1.4 \%$ (mean $=1.1 \%$ ), respectively. In addition, that between $O$. naraharaetmagarum and $O$. esulcata was $8.4-9.1 \%$ (mean $=8.9 \%$ )

## Discussion

The obtained molecular phylogenies showed that the present specimens formed a wellsupported clade. In addition, the mean value of the COI uncorrected $p$-distance among the individuals was $4.4 \%$. This value indicated a clear gap between the present speci-
mens and the closest congener, Orobdella esulcata. Therefore, all of the specimens examined can be considered to belong to the same species, Orobdella naraharaetmagarum.

Although the precise phylogenetic position of $O$. tsushimensis from the Korean Peninsula and the adjacent islets could not be determined in the obtained phylogenies (see Fig. 6), they showed that the genus Orobdella comprises three clades: a Hokkaido lineage including $O$. kawakatsuorum and $O$. koikei; an eastern lineage consisting of four species, $O$. ijimai, $O$. masaakikuroiwai, $O$. octonaria, and $O$. whitmani, distributed in eastern Honshu; and a western lineage containing five previously described species, $O$. mononoke, $O$. dolichopharynx and $O$. shimadae from the Nansei Islands, O. ketagalan from Taiwan, and O. esulcata recorded in Kyushu, Japan. The present phylogenies demonstrated that $O$. naraharaetmagarum was a member of the last clade, and formed a monophyly with $O$. esulcata with strong support. Therefore, the range of the western lineage group covers the area from Chugoku District, at the western tip of Honshu, to Taiwan.

As indicated in Figure 7, Orobdella naraharaetmagarum was divided into eastern (lineage 1 ; locality numbers $1-3$ ) and western (lineage 2 ; locality numbers 4,5 ) phylogroups. The COI uncorrected $p$-distances within lineage 1 were higher than those within lineage 2. The calculated genetic divergences between the three specimens collected from Mt. Kanmuriyama (locality number 5; KUZ Z1582, Z1652, and Z1654) and one individual, KUZ Z1360, from Kitahiroshima (locality number 4) was $1.1-1.4 \%$. The geographic distance between these two collection localities is ca. 28 km . In comparison, the genetic distance between KUZ Z1380 collected on Mt. Azumayama (locality number 2) and KUZ Z1655 from Mt. Izaiyama (locality number 3) was greater than this value (3.1\%), although these two localities are separated only by ca. 7 km . These phylogenetic relationships and genetic divergences implied that leeches belonging to lineage 2 dispersed more recently and rapidly than those of lineage 1. Such discordance between the COI genetic divergences and geographical distances was also seen in the small species $O$. masaakikuroiwai (Nakano 2014). Further molecular phylogenetic studies will help to reveal the biogeographical history of the Orobdella leeches.

The phylogenetic position of $O$. naraharaetmagarum also indicated that the small size of the mature leeches evolved in parallel within Orobdella, as mentioned in Nakano (2014). According to the obtained molecular phylogenies and the phylogenetic trees from studies (Nakano 2012b, 2014), each of the three small species, O. koikei, O. masaakikuroiwai, and $O$. naraharaetmagarum may have diverged from a single large quadrannulate species. As with the other two small species (Nakano 2012b, 2014), O. naraharaetmagarum is also distributed sympatrically with undescribed large quadrannulate species in Chugoku District (Nakano, unpublished data). Therefore, further systematic studies should be carried out to reveal the species diversity and evolutionary history of the genus Orobdella.

## Acknowledgments

The author is grateful to Ayane Maga, Yukiko Narahara, and Yoshiko Yamane (Kyoto University) for providing specimens of the new species, to Professor Hidetoshi Nagamasu (The Kyoto University Museum) for his helpful advice on a specific name of the new species, and to one anonymous reviewer, Dr Bonnie Bain (Dixie State University) and Dr Fredric R. Govedich (Southern Utah University) for their constructive comments on this manuscript. A part of this study was financially supported by Grants for Biodiversity and Evolutionary Research of Global COE (A06) and for Excellent Graduate Schools, both from MEXT, Japan, to Kyoto University, and JSPS Grants-in-Aid for JSPS Fellows (\#15J00720) and Young Scientists (B) (\#26840127) to the author.

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[^0]:    Academic editor: F. Govedich | Received 30 September 2015 | Accepted 30 November 2015 | Published 14 January 2016
    http://zoobank.org/FCCE8F28-BBCE-4C47-8305-4C43EEFF1C01
    Citation: Nakano T (2016) A new quadrannulate species of Orobdella (Hirudinida, Arhynchobdellida, Orobdellidae) from western Honshu, Japan. ZooKeys 553: 33-51. doi: 10.3897/zookeys.553.6723

