

Molecular Systematics and Morphological Analyses of the Subgenus *Setihenricia* (Echinodermata: Asteroidea: *Henricia*) from Japan

Daiki Wakita^{1,4}, Toshihiko Fujita², and Hiroshi Kajihara³

¹Division of Life Science, Graduate School of Life Science, Hokkaido University,
Sapporo, Hokkaido 060-0810, Japan
E-mail: dwntzrd@gmail.com

²Department of Zoology, National Museum of Nature and Science,
Tsukuba, Ibaraki 305-0005, Japan

³Department of Biological Sciences, Faculty of Science, Hokkaido University,
Sapporo, Hokkaido 060-0810, Japan

⁴Corresponding author

(Received 9 December 2017; Accepted 14 May 2019)

The subgenus *Setihenricia* Chichvarkhin and Chichvarkhina, 2017 currently consists of 32 out of 94 species of sea stars in the genus *Henricia* Gray, 1840. However, only 10 have been molecularly ascertained as to the phylogenetic position. Based on seven mitochondrial gene markers (16S rRNA, COI, tRNA-Ala, tRNA-Leu, tRNA-Asn, tRNA-Gln, and tRNA-Pro), we performed molecular analyses to assess the monophyly of *Setihenricia* using 16 species of *Henricia* from Japanese waters, as well as other congeners for which sequences were available in public databases. The monophyletic *Setihenricia* was only weakly recovered because the support values were low, but we assume this taxon may be valid in consideration of a morphological key. *Henricia kinkasana* Hayashi, 1940 is herein identified as a new member of *Setihenricia*, based on molecular and morphological evidence. Sequences in the public databases previously identified as *H. compacta* (Sladen, 1889) and *H. obesa* (Sladen, 1889) were nested in the *Setihenricia* clade; these two species are also likely members of *Setihenricia*. Our morphological assessment yielded three findings: (1) the number of the rows of spines on each inferomarginal plate, being three or more, appears to be a useful diagnostic character for *Setihenricia*, as has been suggested in previous studies, (2) the degree of spine tapering turned out to be less relevant in diagnosing the subgenus than previously supposed, (3) two newly introduced characters—(i) the width of adambulacral plates and (ii) the ratio of two measurements in ambulacral plates to the length of interradius—may be effective to distinguish members of this apparent *Henricia* subclade from other members of the genus.

Key Words: Echinasteridae, phylogeny, RAxML, MrBayes, taxonomic character, scanning electron microscopy (SEM), Pacific, Sea of Japan, East China Sea.

Introduction

At present, 94 species of sea stars in the genus *Henricia* Gray, 1840 are known worldwide, which account for 68% of the total species in the family Echinasteridae (Chichvarkhin 2017a, b; Chichvarkhin and Chichvarkhina 2017; Mah and Hansson 2017). *Henricia* has been recognized as a major taxon of sea stars along the coast of the North Pacific (Fisher 1911; Verrill 1914; Clark and Jewett 2010). From Japanese waters, Uchida (1928) described one species and Hayashi (1940) described or redescribed 15 species, one subspecies, and two forms of *Henricia*. The larval distribution is spatially limited due to breeding habits, causing many local variations (Hayashi 1973). Because the morphology in many species of *Henricia* intergrades almost continuously with one another, it is difficult to define species boundaries (Fisher 1911; Madsen 1987; Clark and Downey 1992; Mah and Blake 2012; Chichvarkhin 2017b). Numerous subspecies and varieties have been established, while a number of named taxa have turned out to be synonymous (Fisher 1910,

1911, 1928, 1930; Verrill 1914; Mortensen 1925; Uchida 1928; Hayashi 1940; D'jakonov 1958; Madsen 1987; Clark and Downey 1992; Clark and Jewett 2010; Eernisse *et al.* 2010).

Within *Henricia*, Chichvarkhin and Chichvarkhina (2017) proposed the new subgenus *Setihenricia*, including 31 species based on morphological characters. Using 16S rRNA (16S), cytochrome *c* oxidase subunit I (COI), and 28S rRNA sequences, Lopes *et al.* (2016) showed that *Henricia* formed a clade along with *Echinaster* (*Echinaster*) Müller and Troschel, 1840, which altogether was sister to *Echinaster* (*Othilia*) Gray, 1840. Of 31 *Setihenricia* species listed by Chichvarkhin and Chichvarkhina (2017), Lopes *et al.*'s (2016) analysis included only one, *H. sanguinolenta* (Müller, 1776). Chichvarkhin (2017b) showed that *Setihenricia* formed a clade with high nodal support in a molecular phylogeny of 16S in terms of six species of *Setihenricia*—*H. djakonovi* Chichvarkhin, 2017a, *H. hayashii* D'jakonov, 1961, *H. lineata* Clark and Jewett, 2010, *H. olga* Chichvarkhin, 2017b, *H. sanguinolenta*, and *H. uluudax* Clark and Jewett, 2010—and four of non-*Setihenricia*; the subgenus is

currently composed of 32 species, with the inclusion of *H. olga*. Knott *et al.* (2018) performed phylogenetic analyses of the two groups formerly referred to as the ‘*pertusa* group’ and ‘*perforata* group’ based on 16S and COI. Chichvarkhin and Chichvarkhina (2017) implied that the ‘*pertusa* group’ represents *Setihenricia*. The ‘*pertusa* group’ also formed a highly supported clade in Knott *et al.*'s (2018) phylogeny,

which included specimens morphologically identifiable as *H. hedingi* Madsen, 1987, *H. lisa* Clark, 1949, *H. pertusa* (Müller, 1776), *H. sanguinolenta*, and *H. spongiosa* (Fabricius, 1780), five of the 31 species originally listed as members of *Setihenricia* (Chichvarkhin and Chichvarkhina 2017). In summary, the monophyly of *Setihenricia* has been investigated in terms of only 10 out of 32 species. One pur-

Table 1. List of species and DDBJ/EMBL/GenBank accession numbers used in the analyses. Where applicable, museum catalogue number and locality of each voucher specimen used in this study are also provided.

Species	Catalogue number	Locality	16S	tRNA-cluster	COI	Source
<i>Henricia alexeyi</i> Chichvarkhin and Chichvarkhina, 2017	—	—	KY464042	—	—	Chichvarkhin (2017b)
<i>Henricia compacta</i> (Sladen, 1889)	—	—	KT268112	—	KT268147	Lopes <i>et al.</i> (2016)
<i>Henricia djakonovi</i> Chichvarkhin, 2017a	—	—	KY464038	—	—	Chichvarkhin (2017b)
<i>Henricia granulifera</i> D'jakonov, 1958	—	—	KY744469	—	—	Chichvarkhin (2017b)
<i>Henricia hayashii</i> D'jakonov, 1961	NSMT E-12501	off Iwate	LC333121	LC336753	LC336732	present study
<i>Henricia hayashii</i> D'jakonov, 1961	—	—	KY934074	—	—	Chichvarkhin (2017b)
<i>Henricia hedingi</i> Madsen, 1987 ^a	—	—	KY853323	—	KY853274	Knott <i>et al.</i> (2018)
<i>Henricia kinkasana</i> Hayashi, 1940	NSMT E-12498	off Miyagi	LC333120	LC336752	LC336731	present study
<i>Henricia lineata</i> Clark and Jewett, 2010	—	—	MF133322	—	—	Chichvarkhin (2017b)
<i>Henricia lisa</i> Clark, 1949 ^a	—	—	KY853377	—	KY853275	Knott <i>et al.</i> (2018)
<i>Henricia nipponica</i> Uchida, 1928	NSMT E-12511	Muroran, Hokkaido	LC333122	LC336754	LC336733	present study
<i>Henricia nipponica</i> Uchida, 1928	NSMT E-12508	Oshoro, Hokkaido	LC333123	LC336755	LC336734	present study
<i>Henricia nipponica</i> Uchida, 1928	—	—	D63737	—	—	Wada <i>et al.</i> (1996)
<i>Henricia obesa</i> (Sladen, 1889)	—	—	KT268113	—	KT268148	Lopes <i>et al.</i> (2016)
<i>Henricia oculata</i> (Pennant, 1777)	—	—	KT268116	—	KT268151	Lopes <i>et al.</i> (2016)
<i>Henricia oculata</i> (Pennant, 1777)	—	—	AY652500	—	—	Chichvarkhin (2017b)
<i>Henricia oculata</i> (Pennant, 1777)	—	—	KY853395	—	KY853317	Knott <i>et al.</i> (2018)
<i>Henricia ohshimai</i> Hayashi, 1935	NSMT E-12500	Misaki, Kanagawa	LC333125	LC336757	LC336736	present study
<i>Henricia ohshimai</i> Hayashi, 1935	NSMT E-12507	Oshoro, Hokkaido	LC333124	LC336756	LC336735	present study
<i>Henricia ohshimai</i> Hayashi, 1935	—	—	AB084571	—	—	Matsubara <i>et al.</i> (2004)
<i>Henricia olga</i> Chichvarkhin, 2017b	—	—	KY934079	—	—	Chichvarkhin (2017b)
<i>Henricia pachyderma</i> Hayashi, 1940	—	—	D63739	—	—	Wada <i>et al.</i> (1996)
<i>Henricia pachyderma</i> Hayashi, 1940	—	—	KX610476	—	—	Chichvarkhin (2017b)
<i>Henricia perforata</i> (Müller, 1776)	—	—	KY853333	—	KY853302	Knott <i>et al.</i> (2018)
<i>Henricia pertusa</i> (Müller, 1776)	—	—	KY853337	—	KY853286	Knott <i>et al.</i> (2018)
<i>Henricia regularis</i> Hayashi, 1940	NSMT E-12506	Iki Island, Nagasaki	LC333128	LC336760	LC336739	present study
<i>Henricia reniessa</i> Hayashi, 1940	NSMT E-12505	off Iwanai, Hokkaido	LC333129	LC336761	LC336740	present study
<i>Henricia reticulata</i> Hayashi, 1940	NSMT E-12513	Akkeshi, Hokkaido	LC333126	LC336758	LC336737	present study
<i>Henricia sanguinolenta</i> (Müller, 1776)	—	—	AY652499	—	HM542200	Lopes <i>et al.</i> (2016)
<i>Henricia sanguinolenta</i> (Müller, 1776)	—	—	KT268115	—	—	Chichvarkhin (2017b)
<i>Henricia sanguinolenta</i> (Müller, 1776) ^a	—	—	KY853338	—	KY853253	Knott <i>et al.</i> (2018)
<i>Henricia spongiosa</i> (Fabricius, 1780) ^a	—	—	KY853393	—	KY853268	Knott <i>et al.</i> (2018)
<i>Henricia tumida</i> Verrill, 1909	NSMT E-12510	Akkeshi, Hokkaido	LC333136	LC336768	LC336747	present study
<i>Henricia tumida</i> Verrill, 1909	—	—	AF290033	—	AF290035	Foltz and Rocha-Olivares (unpublished)
<i>Henricia uluudax</i> Clark and Jewett, 2010	—	—	KY934075	—	—	Chichvarkhin (2017b)
<i>Henricia</i> sp. 1 (Akkeshi)	NSMT E-12509	Akkeshi, Hokkaido	LC333133	LC336765	LC336744	present study
<i>Henricia</i> sp. 2 (Abashiri 1)	NSMT E-12502	Abashiri, Hokkaido	LC333131	LC336763	LC336742	present study
<i>Henricia</i> sp. 3 (Abashiri 2)	NSMT E-12503	Abashiri, Hokkaido	LC333132	LC336764	LC336743	present study
<i>Henricia</i> sp. 4 (Fukushima)	NSMT E-12504	off Fukushima	LC333130	LC336762	LC336741	present study
<i>Henricia</i> sp. 5 (Okinawa)	NSMT E-12497	Okinawa Island, Okinawa	LC333127	LC336759	LC336738	present study
<i>Henricia</i> sp. 6 (Miyako)	NSMT E-12496	Miyako, Okinawa	LC333134	LC336766	LC336745	present study
<i>Henricia</i> sp. 7 (Nagannu)	NSMT E-12499	Nagannu Island, Okinawa	LC333135	LC336767	LC336746	present study
<i>Henricia</i> sp. 8 (Okinoerabu)	NSMT E-12495	Okinoerabu Island, Kagoshima	LC333119	LC336751	LC336730	present study
<i>Henricia</i> sp. 9 ^b	—	—	KY853329	—	KY853310	Knott <i>et al.</i> (2018)
Outgroup						
<i>Echinaster callosus</i> Marenzeller, 1895	NSMT E-6709	Aka Island, Okinawa	LC333116	LC336748	—	present study
<i>Echinaster callosus</i> Marenzeller, 1895	—	—	KT268086	—	KT268121	Lopes <i>et al.</i> (2016)
<i>Echinaster luzonicus</i> (Gray, 1840)	NSMT E-12512	Iriomote Island, Okinawa	LC333117	LC336749	—	present study
<i>Echinaster luzonicus</i> (Gray, 1840)	—	—	KT268102	—	KT268137	Lopes <i>et al.</i> (2016)
<i>Echinaster sepositus</i> (Retzius, 1783)	NSMT E-12494	Iheya Island, Okinawa	LC333118	LC336750	LC336729	present study
<i>Echinaster sepositus</i> (Retzius, 1783)	—	—	KT268108	—	KT268146	Lopes <i>et al.</i> (2016)

^aFollowing “initial identification” in Knott *et al.* (2018). ^bReferred to as “*Henricia perforata* group” or “*Henricia* sp., clade D” in Knott *et al.* (2018).

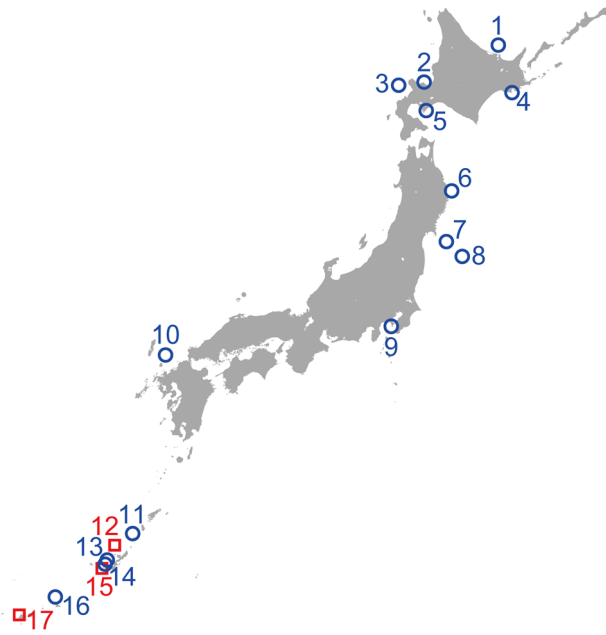


Fig. 1. Sampling locations of *Henricia* (circle) and *Echinaster* (square) specimens around Japan. (1) Off Abashiri, Hokkaido; *H. sp.* 2 (Abashiri 1), *H. sp.* 3 (Abashiri 2); (2) Oshoro, Hokkaido; *H. nipponica* (Oshoro), *H. ohshimai* (Oshoro); (3) Off Iwanai, Hokkaido; *H. reniessa*; (4) Akkeshi, Hokkaido; *H. reticulata*, *H. tumida*, *H. sp.* 1 (Akkeshi); (5) Muroran, Hokkaido; *H. nipponica* (Muroran); (6) Off Iwate; *H. hayashii*; (7) Off Miyagi; *H. kinkasana*; (8) Off Fukushima; *H. sp.* 4 (Fukushima); (9) Misaki, Kanagawa; *H. ohshimai* (Misaki); (10) Iki Island, Nagasaki; *H. regularis*; (11) Okinoerabu Island, Kagoshima; *H. sp.* 8 (Okinoerabu); (12) Iheya Island, Okinawa; *E. sepositus*; (13) Okinawa Island, Okinawa; *H. sp.* 5 (Okinawa); (14) Nagannu Island, Okinawa; *H. sp.* 7 (Nagannu); (15) Aka Island, Okinawa; *E. callosus*; (16) Miyako Island, Okinawa; *H. sp.* 6 (Miyako); (17) Iriomote Island, Okinawa; *E. luzonicus*.

Table 2. List of primers used in the present study.

Gene	Primer	Sequence (5' to 3')	Source
16S	16Sar	cgctgtttatcaaaaacat	Palumbi (1996)
	16Sbr	ccggtctgaactcagatcacgt	Palumbi (1996)
tRNA-cluster	Pat2	cttgaaggcttttagttgattaac	Hart <i>et al.</i> (1997)
	ECOLc	ccaatatcctgtgttagtaga	Smith <i>et al.</i> (1993)
COI	LCOech1aF1	tttttctactaaacacaaggatattgg	Layton <i>et al.</i> (2016)
	HCO2198	taaacttcagggtgacaaaataca	Folmer <i>et al.</i> (1994)

Table 3. Best partition scheme and nucleotide substitution models using AICc for maximum likelihood (ML) and BIC for Bayesian inference (BI) in PartitionFinder (Lanfear *et al.* 2012).

Subset partitions	Best model (ML)	Best model (BI)
16S	GTR+I+G	HKY+I+G
tRNA-Ala, tRNA-Leu, tRNA-Gln, tRNA-Pro	GTR+I+G	HKY+G
tRNA-Asn, COI 1st codon	GTR+I+G	K80+I
COI 2nd codon	GTR+G	F81
COI 3rd codon	GTR+I+G	HKY+I+G

pose of this study is to see if the monophyly of *Setihenricia* is still recovered even when additional OTUs from Japanese waters are included in an analysis. Our new OTUs include *H. nipponica* Uchida, 1928, *H. regularis* Hayashi, 1940, *H. reniessa* Hayashi, 1940, and *H. tumida* Verrill, 1909, which were listed as members of *Setihenricia* (Chichvarkhin and Chichvarkhina 2017) but have not been placed on a molecular phylogenetic context. We also test by morphological and molecular data if the three species *H. kinkasana* Hayashi, 1940, *H. ohshimai* Hayashi, 1935, and *H. reticulata* Hayashi, 1940 belong to *Setihenricia*, although they were not origi-

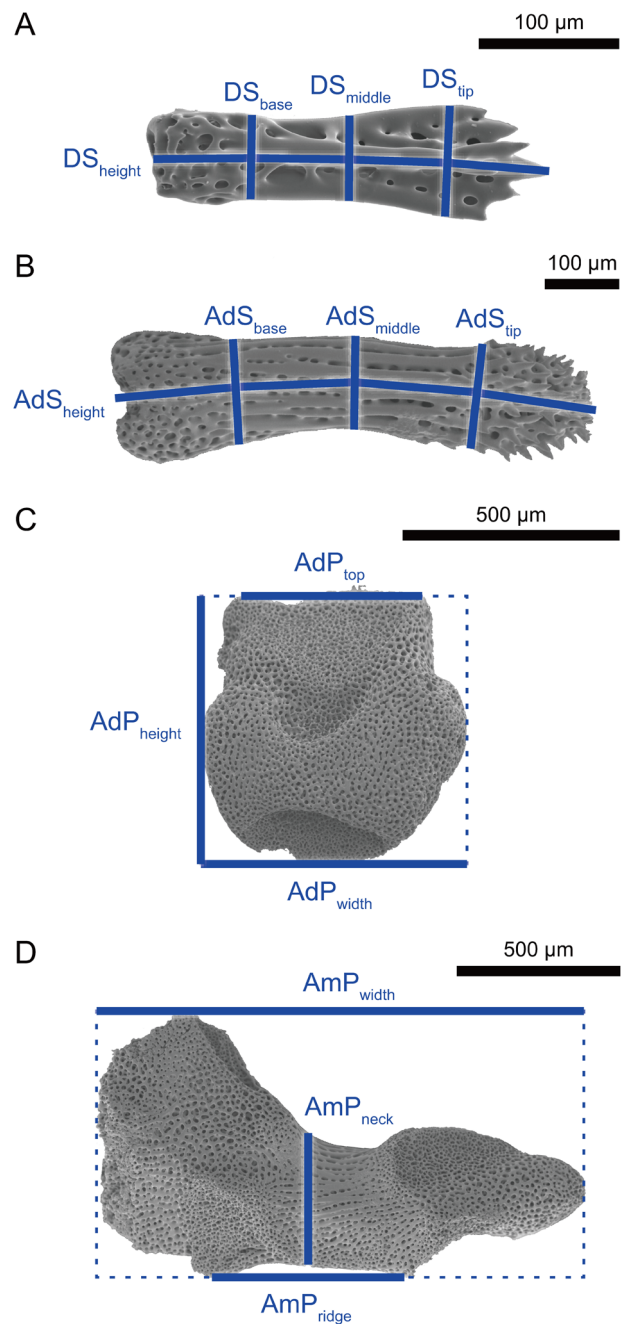


Fig. 2. Measurements (morphometrics) of four kinds of ossicles of *Henricia nipponica* (Oshoro): A, dorsal spine (DS); B, adambulacral spine (AdS); C, adambulacral plate (AdP), proximal view; D, ambulacral plate (AmP), proximal view.

Table 4. Abbreviations and definitions of the morphological characters used in this study.

Abbreviation	Measurement
Macroscopy	
R	length from the center of the mouth to the tip of the longest arm (in mm)
r	length from the center of the mouth to the midpoint of the margin of the longest interradus (in mm)
DS (dorsal spine) ^a	
DS _{height}	height (in μm)
DS _{base}	width at the quarter from the base (in μm)
DS _{middle}	width at the middle (in μm)
DS _{tip}	width at the quarter from the tip (in μm)
AdS (adambulacral spine) ^b	
AdS _{height}	height (in μm)
AdS _{base}	width at the quarter from the base (in μm)
AdS _{middle}	width at the middle (in μm)
AdS _{tip}	width at the quarter from the tip (in μm)
AdP (adambulacral plate) ^c	
AdP _{height}	height of the minimum circumscribed rectangle viewed proximally (in μm) ^d
AdP _{width}	width of the minimum circumscribed rectangle viewed proximally (in μm) ^d
AdP _{top}	length between the two aboral ridges viewed proximally (in μm)
AmP (ambulacral plate) ^e	
AmP _{width}	width of the circumscribed rectangle, the bottom edge of which is parallel to the line segment that connects the two oral ridges, viewed distro-proximally (in μm)
AmP _{ridge}	length between the two oral ridges (in μm)
AmP _{neck}	length of the line segment that runs on the plate and also crosses at the right angles the midpoint of the line segment connecting the two oral ridges (in μm)

^aMeasured for 2–10 spines at the aboral base of arms from each specimen. ^bMeasured for 2–5 spines which are nearest to furrow spines within the range from about the fourth to tenth positions in the adambulacral plate series from each specimen. ^cMeasured for 2–8 plates within the range from about the fourth to tenth positions in the adambulacral plate series from each specimen. ^dMeasured for a rectangle formed by tangents to an adambulacral plate viewed distro-proximally, with the upper tangent passing between two aboral ridges. ^eMeasured for 3–6 plates within the range from about the fourth to tenth positions in the ambulacral plate series from each specimen.

nally included in the subgenus. Assessment of whether *Setihenricia* is monophyletic would in turn resolve whether this subgenus should be accepted as a valid taxon.

Chichvarkhin and Chichvarkhina (2017) defined *Setihenricia* by (1) apical thorns of dorsal spines form crown, with the thorns pointed upwards or arranged on a club-shaped apex; (2) spines almost never tapering distally; (3) no thorns on spine shaft; (4) spines in inferomarginal brush-like pseudopaxillae in three or more rows; and (5) spines not covered by thick skin membrane. Among these characters, only (4) is quantitative, whereas objective assessment can be difficult as to the characters (1), (2), (3), and (5). The skeleton of a sea star comprises a variety of ossicles (Mortensen 1925; Blake 1987; Gale 1987); there might be some quantitative characters that were not mentioned in the diagnosis of *Setihenricia* but can effectively characterize the subgenus. On the assumption that this taxon is valid, the other purpose of our study is to explore whether some quantitative morphological characters can distinguish *Setihenricia* species from non-*Setihenricia* species. For this purpose, we measured several parts of four kinds of ossicles. We also attempted to quantify the degree of spine tapering to assess the diagnostic usefulness of the character (2).

Materials and Methods

Specimens

In total, 18 specimens of *Henricia* (representing eight nominal species and eight unidentified ones) from Japanese waters were used in this study (Table 1, Fig. 1). Species identification followed Hayashi (1940) and D'jakonov (1961). Of the 18, six were freshly collected and preserved in 99% ethanol, while 12 ethanol-preserved specimens had been deposited in the National Museum of Nature and Science, Tsukuba, Japan (NSMT).

Molecular phylogeny

Total DNA was extracted from tube feet of ethanol-preserved specimens using silica (Boom *et al.* 1990) with some modifications. Three regions in the mitochondrial genome were amplified by polymerase chain reaction (PCR): a fragment of 16S, a segment comprising five tRNA genes (tRNA-cluster: tRNA-Ala, tRNA-Leu, tRNA-Asn, tRNA-Gln, and tRNA-Pro), and a fragment of COI, using primers listed in Table 2. PCR was performed by a thermal cycler, 2720 Thermal Cycler (Applied Biosystems), in a 10 μl reaction volume containing 2 μl of template DNA and 8 μl of premix made with 5.55 μl deionized water, 1 μl Ex *Taq* Buffer (TaKaRa Bio), 0.8 μl dNTP (each 2.5 mM), 0.3 μl each primer (each 10 μM), and 0.05 μl TaKaRa Ex *Taq* (5 U/ μl , TaKaRa Bio).

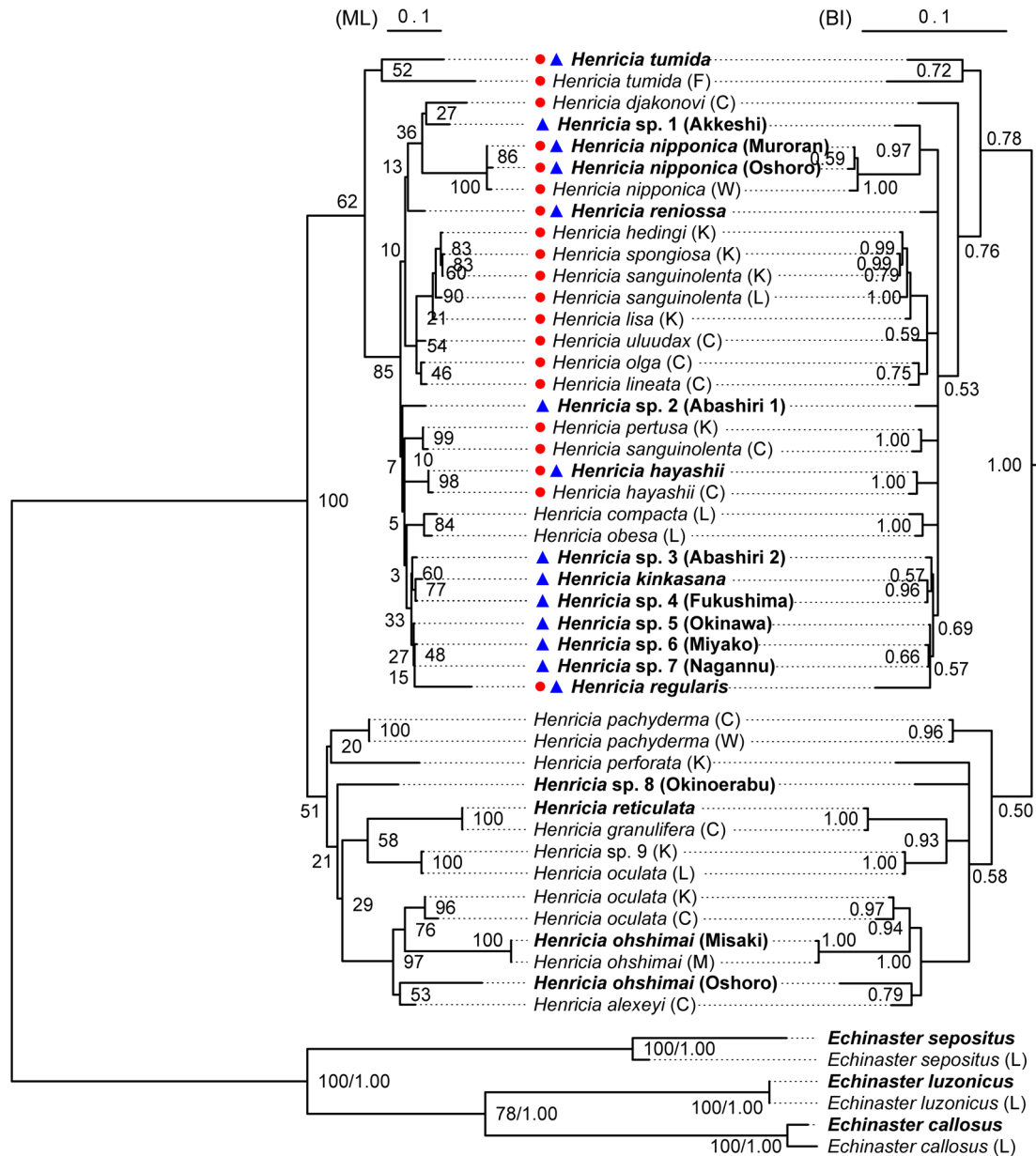


Fig. 3. Phylogenetic trees obtained from a concatenated dataset with a total length of 1,277 bp, consisting of seven mitochondrial genes (16S, tRNA-Ala, tRNA-Leu, tRNA-Asn, tRNA-Gln, tRNA-Pro, and COI). The trees were built based on maximum likelihood (ML, left) and Bayesian inference (BI, right). Values at nodes indicate bootstrap scores from ML and posterior probabilities from BI. Outgroups are only shown in the ML tree with both the support values. Scale bars indicate the number of nucleotide substitutions per site. OTUs sequenced in this study are in bold face. Each letter in parentheses after non-bold OTUs denotes the source: C, Chichvarkhin (2017b); F, Foltz and Rocha-Olivares (unpublished); K, Knott *et al.* (2018); L, Lopes *et al.* (2016); M, Matsubara *et al.* (2004); W, Wada *et al.* (1996). Circles indicate species listed as *Setihenricia* in Chichvarkhin and Chichvarkhina (2017). Triangles indicate species morphologically identified as *Setihenricia* in this study (see Fig. 4A).

Amplification consisted of an initial denaturation (1 min, 95°C); followed by 30 cycles of denaturation (30 sec, 95°C), annealing (30 sec, 50°C for 16S, 41°C for tRNA-cluster, 47°C for COI), and elongation (1 min, 72°C); ended by a final elongation (7 min, 72°C). The PCR product was purified using silica (Boom *et al.* 1990). Both strands were sequenced with a BigDye® Terminator ver. 3.1 Cycle Sequencing Kit (Applied Biosystems) following the manufacturer's protocol, using the same primer set as the initial PCR amplification (Table 2). Sequencing was performed with an ABI Prism 3730 DNA Analyzer (Applied Biosystems).

Chromatogram and sequence data were checked with MEGA ver. 6.06 (Tamura *et al.* 2013), eliminating primer sequences. COI sequences of *Echinaster callosus* Marenzeller, 1895 and *Echinaster luzonicus* (Gray, 1840) were not obtained in our study. Twenty-nine sequences of 16S and 14 sequences of COI were obtained from GenBank (Table 1). The 16S and tRNA-cluster sequences were aligned using MAFFT ver. 7 (Katoh and Standley 2013) with default parameters. The aligned tRNA-cluster sequences were divided into five gene sequences (tRNA-Ala, tRNA-Leu, tRNA-Asn, tRNA-Gln, and tRNA-Pro) based on the feature data in

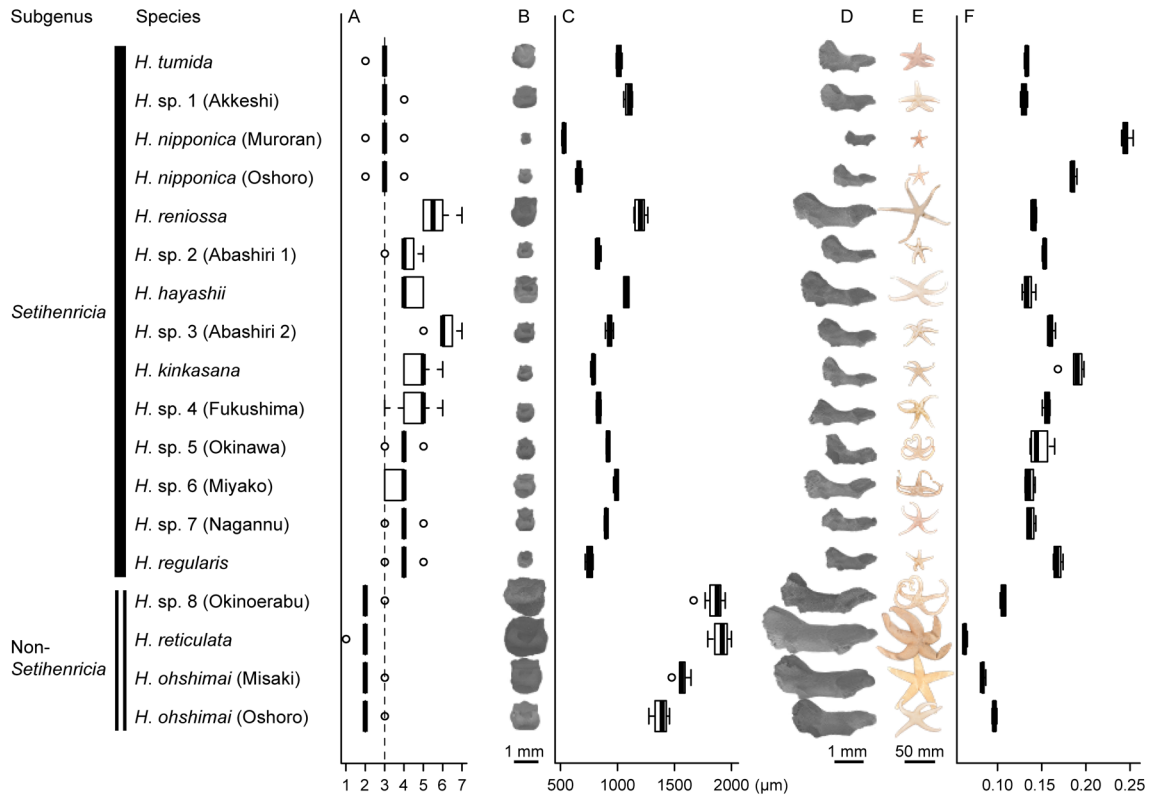


Fig. 4. Morphological characters which might be useful to distinguish *Setihenricia* species from non-*Setihenricia* species. A, number of rows of spines on each inferomarginal plate; B, SEM images of AdP; C, AdP_{width}; D, SEM images of AmP; E, photographs of entire body; F, ratios of AmP_{width}/AmP_{neck/r}. See Table 4 for abbreviations. Circles on boxplots (A, C, and F) indicate outliers. Specimens that show three (indicated by a vertical dashed line on A) or more rows of the spines can be identified as *Setihenricia* (Chichvarkhin and Chichvarkhina 2017). Enlarged images of B, D, and E are available in the figshare repository (Wakita *et al.* 2019).

GenBank (AY245501; Knott *et al.* 2003). The last nucleotide position in the tRNA-Ala was eliminated because it overlapped with the first position of the tRNA-Leu. The 16S, each of the tRNA, and COI sequences were trimmed using Gblocks ver. 0.91b (Talavera and Castresana 2007) with the most stringent option. The COI sequences contained no indels. The 16S, tRNA-cluster, and COI sequences were concatenated with Kakusan ver. 4 (Tanabe 2011). The final dataset had a total length of 1,277 bp, consisting of 16S (451 bp), tRNA-Ala (65 bp), tRNA-Leu (70 bp), tRNA-Asn (72 bp), tRNA-Gln (70 bp), tRNA-Pro (67 bp), and COI (482 bp).

Outgroups were chosen from *Echinaster* (*Echinaster*), which is sister to *Henricia* based on Lopes *et al.*'s (2016) analysis. Specifically, the selected outgroups were *Echinaster callosus*, *E. luzonicus*, and *E. sepositus* (Retzius, 1783) (Table 1). The best partition schemes and nucleotide substitution models were estimated using AICc for maximum likelihood (ML) and BIC for Bayesian inference (BI) in PartitionFinder ver. 1.1.1 (Lanfear *et al.* 2012) with the “greedy” scheme (Table 3). The concatenated sequences were analyzed with the ML and BI methods. ML phylogenetic tree was estimated by using RAxML ver. 8.2.10 (Stamatakis 2014) based on the GTR+I+G model. Bootstrap support was calculated from 1,000 replicates. BI phylogenetic tree was estimated with MrBayes ver. 3.2.6 (Ronquist *et al.* 2012) based on the Markov chain Monte Carlo algorithm (MCMC) for 1,000,000 generations in two runs, each with four chains,

sampled with a frequency of every 100 generations. The convergence in MCMC was checked with Tracer ver. 1.6 (Rambaut *et al.* 2014); burn-in parameter was set to exclude the first 25% of the trees. The best ML tree and the 50% majority-rule consensus BI tree were visualized and manipulated in FigTree ver. 1.4.2 (Rambaut 2014).

Morphology

The number of the rows of spines on each inferomarginal plate was counted under a stereomicroscope; 20 plates were randomly selected within the range from about the fourth to tenth positions in the plate series of at least two arms. For observation of ossicles by scanning electron microscopy (SEM), body walls were cut and put in an undiluted solution of commercial bleach for an hour. Dorsal spines (DS; also referred to as aboral or abactinal spines) were extracted from the aboral walls at the base of an arm. Adambulacral spines (AdS), adambulacral plates (AdP), and ambulacral plates (AmP) were extracted from an ambulacral area at the base of an arm, ranging from about the fourth to tenth plate series. Ossicles of these four kinds were sub-sampled from each specimen, coated with gold using Ion Sputter E-1045 (Hitachi; 180 sec, 15 mA), and observed with S-3000N scanning electron microscope (Hitachi; 10.1 or 15.0 kV). Several parts in the ossicles were measured with ImageJ (Abramoff *et al.* 2004) (Fig. 2A–D, Table 4).

Some measurements were combined into ratios. The de-

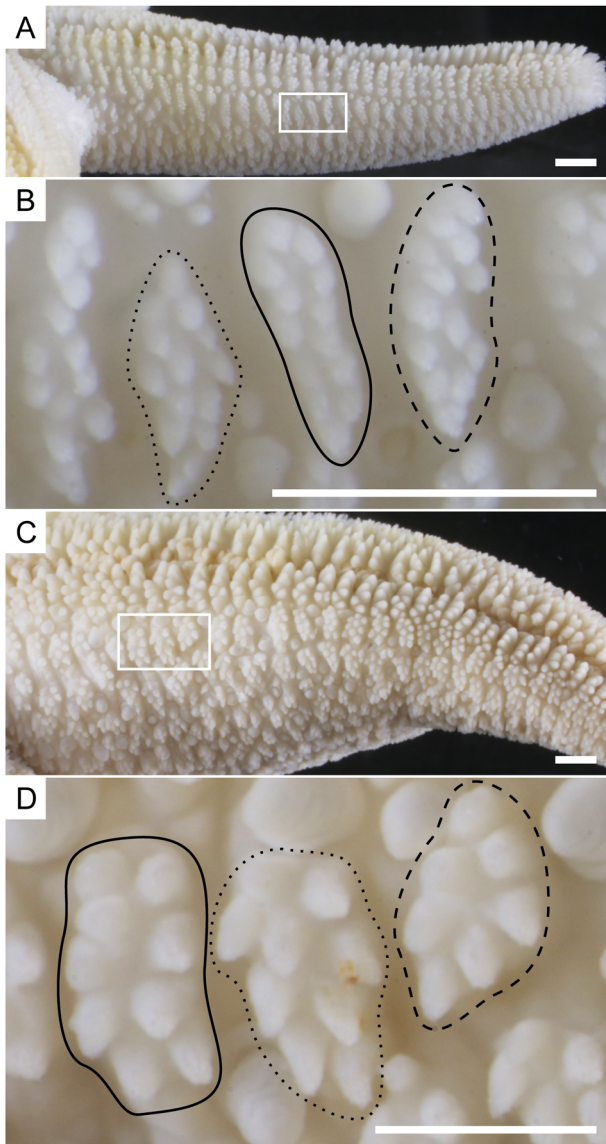


Fig. 5. Side views of the arms of *Henricia nipponica* (Oshoro) (A, B) and *Henricia tumida* (C, D). A, C, Lower magnification; B, D, higher magnification of the rectangle on A and C, respectively. The number of the rows of spines on the inferomarginal plate can be interpreted as either two (solid line), two or three (dashed line), or three (dotted line). Scale bars indicate 1 mm.

gree of tapering of DS and AdS was represented by DS_{base}/DS_{tip} (Fig. 2A) and AdS_{base}/AdS_{tip} (Fig. 2B), the ratios of the width at the quarter point from the base to the width at the quarter point from the tip in spines. The values of these ratios should be less than one if the spines are “almost never tapering distally” as stated in the diagnosis of *Setihenricia* (Chichvarkhin and Chichvarkhina 2017).

Results

The ML and BI trees moderately resembled each other in terms of topology, where *Henricia* was monophyletic with maximum support values (100/1.00; bootstrap score from ML and posterior probability from BI, respectively; Fig. 3).

All the OTUs of *Henricia* were principally divided into two clades with low support; the first clade (62/0.78) included *H. compacta* (Sladen, 1889), *H. djakonovi*, *H. hayashii*, *H. hedingi*, *H. kinkasana*, *H. lineata*, *H. lisa*, *H. nipponica*, *H. obesa* (Sladen, 1889), *H. olga*, *H. pertusa*, *H. regularis*, *H. reniessa*, *H. sanguinolenta*, *H. spongiosa*, *H. tumida*, and *H. uluudax*, as well as the unidentified *H. spp.* 1–7; the second clade (51/0.50) had *H. alexeyi* Chichvarkhin and Chichvarkhina, 2017, *H. granulifera* D’jakonov, 1958, *H. oculata* (Pennant, 1777), *H. ohshimai*, *H. pachyderma* Hayashi, 1940, *H. perforata* (Müller, 1776), *H. reticulata*, and *H. spp.* 8 and 9.

Specimens with three or more rows of spines on each inferomarginal plate, and thus morphologically identifiable as *Setihenricia*, were *H. hayashii*, *H. kinkasana*, *H. regularis*, *H. reniessa*, *H. spp.* 1–7 (Fig. 4A). We also identified *H. nipponica* (Murooran), *H. nipponica* (Oshoro), and *H. tumida* as *Setihenricia*, because they showed mostly three or more, but rarely two, rows of the spines (Fig. 5A–D). The other four specimens—*H. ohshimai* (Misaki), *H. ohshimai* (Oshoro), *H. reticulata*, and *H. sp.* 8 (Okinoerabu)—were not identified as *Setihenricia* because they consistently had fewer than three rows of the spines. Among the morphological characters examined in this study (see Appendix), AdP_{width} (Fig. 4B, C) and $AmP_{width}/AmP_{neck}/r$ (Fig. 4D–F) appeared to be useful to distinguish *Setihenricia* from the others. Values of the ratio DS_{base}/DS_{tip} often exceeded more than one in the specimens identified as *Setihenricia*, except for *H. hayashii* (Fig. 6A, B). Values of the ratio AdS_{base}/AdS_{tip} were also more than one in the *Setihenricia* specimens, except for *H. hayashii*, *H. nipponica* (Oshoro), *H. tumida*, and *H. sp.* 3 (Abashiri 2) (Fig. 6C, D).

Discussion

In our dataset, 25 species names were applied to the OTUs included in the analyses. Of the 25, 14 had been originally listed as *Setihenricia* by Chichvarkhin and Chichvarkhina (2017). The 14 species are: *H. djakonovi*, *H. hayashii*, *H. hedingi*, *H. lineata*, *H. lisa*, *H. nipponica*, *H. olga*, *H. pertusa*, *H. regularis*, *H. reniessa*, *H. sanguinolenta*, *H. spongiosa*, *H. tumida*, and *H. uluudax*. Although our best tree from the analyses based on 16S, tRNA-cluster, and COI sequences included a monophyletic grouping of these species, its support values were low (62% bootstrap score in ML and 0.78 posterior probability in BI; Fig. 3). Thus the monophyly of *Setihenricia* was not strongly indicated, which suggests that we cannot positively accept the validity of this taxon. The lack of support was likely due to weak evidence for *H. tumida* grouping with other *Setihenricia*. Within the *Setihenricia* grouping, the ML and BI trees differed in terms of the position of *H. djakonovi*. To obtain a more robust phylogenetic estimate of the genus *Henricia*, other markers such as 18S and 28S rRNA genes and more complete taxon sampling should be sought in future analyses.

Assuming that *Setihenricia* may be a valid taxon, one of the characters that can define the subgenus appears to be

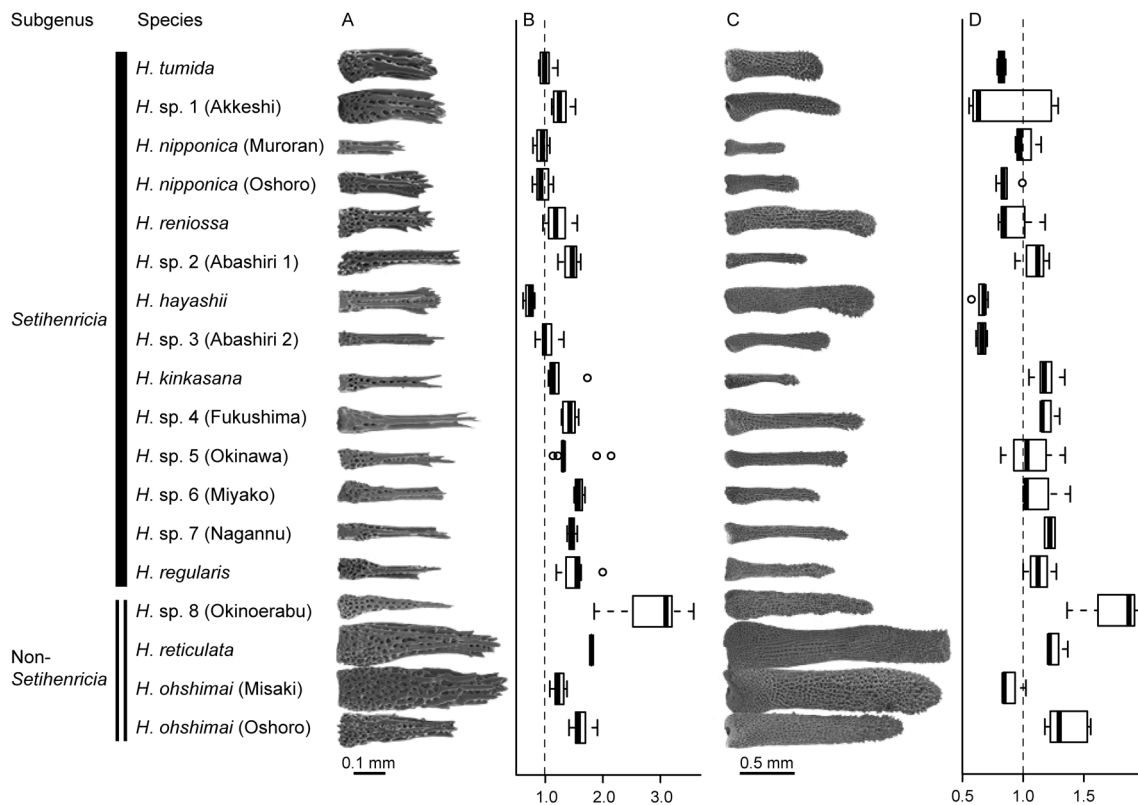


Fig. 6. Morphological characters representing the degree of tapering of spines. A, SEM images of DS; B, ratios of DS_{base}/DS_{tip} ; C, SEM images of AdS; D, ratios of AdS_{base}/AdS_{tip} . See Table 4 for abbreviations. Circles on boxplots (B and D) indicate outliers. On B and D, a value above one (indicated by a vertical dashed line) can be interpreted that the spine is tapered. Enlarged images of A and C are available in the figshare repository (Wakita *et al.* 2019).

three or more rows of spines on each inferomarginal plate (Fig. 4A). However, only two rows of spines were rarely observed in *H. nipponica* and *H. tumida* (Fig. 5). In such cases, some rows were arranged irregularly (Fig. 5B, D), making it difficult to assess the exact number of the rows. The “rows of spines” can be defined more strictly to exclude subjectivity, which, however, is beyond the scope of this study. Although the phylogenetic supports of *Setihenricia* were low, this taxon could be defined by this morphological character if we ignore the rare cases, so we here retain its validity for further discussions.

We found *H. kinkasana* represents a new member of *Setihenricia* based on its molecular phylogenetic position and morphological characters. In addition, *H. compacta* and *H. obesa* appear to belong to the subgenus as well, because the sequences in the public databases identified by Lopes *et al.* (2016) were nested among other *Setihenricia* species in our ML and BI trees (Fig. 3). These three (*H. compacta*, *H. kinkasana*, and *H. obesa*) were not originally included in *Setihenricia* by Chichvarkhin and Chichvarkhina (2017) probably because the number of the rows of spines on each inferomarginal plate—one of the diagnostic characters—had not been sufficiently described (Sladen 1889; Hayashi 1940; Rowe and Albertson 1986). Our specimen of *H. kinkasana* possessed four or more rows of the spines (Fig. 4A).

The degree of spine tapering appears to be less relevant in diagnosing *Setihenricia*. Chichvarkhin and Chichvarkhina

(2017) stated that spines are “almost never tapering distally” in *Setihenricia*. If this character is represented by the ratio of the width at the quarter point from the base to the width at the quarter point from the tip in dorsal spines (DS_{base}/DS_{tip} , *cf.* Fig. 2A) and in adambulacral spines (AdS_{base}/AdS_{tip} , *cf.* Fig. 2B), these ratios should take values of less than one. Nevertheless, out of the 14 specimens identified as *Setihenricia* in our study, 13 in DS_{base}/DS_{tip} and 10 in AdS_{base}/AdS_{tip} showed the values of more than one (Fig. 6B, D), indicating that their dorsal and adambulacral spines are indeed tapering distally (Fig. 6A, C). This tendency was also recognized in more apical regions represented by DS_{middle}/DS_{tip} and AdS_{middle}/AdS_{tip} (*cf.* Fig. 2A, B, and Appendix).

We introduced quantitative morphological characters in terms of dorsal spine (DS), adambulacral spine (AdS), adambulacral plate (AdP), and ambulacral plate (AmP) in the hope of distinguishing *Setihenricia* species from non-*Setihenricia* species. Most of the characters seemed to be useless for this purpose, except two, which exhibited potential usefulness as new taxonomic characters. One was the width of adambulacral plates (AdP_{width} , *cf.* Fig. 2C), which was smaller in the species identified or identifiable as *Setihenricia* (Fig. 4C). The other was the ratio of two measurements in ambulacral plates to a macroscopic measurement ($AmP_{width}/AmP_{neck/l}$, *cf.* Fig. 2D); it was greater in the *Setihenricia* specimens (Fig. 4F). The intra- and interspecific variation of these characters should be investigated with a large sample

size in future studies.

Acknowledgments

Our heartfelt appreciation goes to Dr. K. Kakui (Hokkaido University, HU) for deep discussion with tender care. We sincerely thank Dr. H. Fortunato and Dr. K. Wakeman (HU) for kindly having commented on an earlier version of the manuscript. Our colleagues have given me enormous helps. We are particularly grateful to Dr. S. Tomioka and Mr. A. Shibata (HU) for supporting our experiments and analyses. The precious samples have been borrowed from the National Museum of Nature and Science, Tsukuba, Japan. Taxonomic ideas and some samples of sea stars comprising our study have been provided by Mr. M. Arai and Mr. A. Ogawa (The University of Tokyo). We have received many favors of the staff from Oshoro Marine Station, Akkeshi Marine Station, Muroran Marine Station, and Iriomote Station for sampling. Dr. A. Chichvarkhin (Institute of Marine Biology, Vladivostok) motivated us for SEM work and provided us helpful ideas on species identification.

References

- Abràmoff, M. D., Magalhães, P. J., and Ram, S. J. 2004. Image processing with ImageJ. *Biophotonics International* 11: 36–42.
- Blake, D. B. 1987. A classification and phylogeny of post-Paleozoic sea stars (Asteroidea: Echinodermata). *Journal of Natural History* 21: 481–528.
- Boom, R., Sol, C. J. A., Salimans, M. M. M., Jansen, C. L., Wertheim-van Dillen, P. M. E., and van der Noordaa, J. 1990. Rapid and simple method for purification of nucleic acids. *Journal of Clinical Microbiology* 28: 495–503.
- Chichvarkhin, A. Y. 2017a. *Henricia djakonovi* sp. nov. (Echinodermata, Echinasteridae): a new sea star species from the Sea of Japan. *PeerJ* 5: e2863.
- Chichvarkhin, A. Y. 2017b. Sea star *Henricia spiculifera* (Clark, 1901) in the northwestern Pacific: one species or three? *PeerJ* 5: e3489.
- Chichvarkhin, A. Y. and Chichvarkhina, O. V. 2017. A new sea star species of the genus *Henricia* Gray, 1840 (Echinodermata, Asteroidea) from the northwestern Sea of Japan and description of a new subgenus. Pp. 202–209. *In*: Dautova, T. N., Sun, X., Song, S., and Adrianov, A. (Eds) *Life-supporting Asia-Pacific Marine Ecosystems*. Science Press, Beijing.
- Clark, A. M. and Downey, M. E. 1992. *Starfishes of the Atlantic*. Chapman and Hall, London, xxvi+794 pp., 113 pls.
- Clark, R. N. and Jewett, S. C. 2010. A new genus and thirteen new species of sea stars (Asteroidea: Echinasteridae) from the Aleutian Island Archipelago. *Zootaxa* 2571: 1–36.
- D'jakonov, A. M. 1958. [Echinoderms (Echinodermata), except holothurians, collected by Kuril-Sakhalin expedition in 1947–1949]. *Issledovaniya Dal'nevostochnykh Morei SSSR* 5: 271–357. [In Russian]
- D'jakonov, A. M. 1961. [Review of sea stars of the genus *Henricia* Gray from the north-western parts of Pacific Ocean]. *Issledovaniya Dal'nevostochnykh Morei SSSR* 7: 5–39. [In Russian]
- Eernisse, D. J., Strathmann, M. F., and Strathmann R. R. 2010. *Henricia pumila* sp. nov.: a brooding seastar (Asteroidea) from the coastal northeastern Pacific. *Zootaxa* 2329: 22–36.
- Fisher, W. K. 1910. New starfishes from the North Pacific, II: Spinulosa. *Zoologischer Anzeiger* 35: 568–574.
- Fisher, W. K. 1911. *Asteroidea of the North Pacific and Adjacent Waters, Part 1: Phanerozonia and Spinulosa*. U. S. Government Printing Office, Washington, D. C., vi+419 pp., 122 pls.
- Fisher, W. K. 1928. *Asteroidea of the North Pacific and Adjacent Waters, Part 2: Forcipulata (Part)*. U. S. Government Printing Office, Washington, D. C., iii+245 pp., 81 pls.
- Fisher, W. K. 1930. *Asteroidea of the North Pacific and Adjacent Waters, Part 3: Forcipulata (Concluded)*. U. S. Government Printing Office, Washington, D. C., iii+356 pp., 93 pls.
- Folmer, O., Black, M., Hoeh, W., Lutz, R., and Vrijenhoek, R. 1994. DNA primers for amplification of mitochondrial cytochrome *c* oxidase subunit I from diverse metazoan invertebrates. *Molecular Marine Biology and Biotechnology* 3: 294–299.
- Gale, A. S. 1987. Phylogeny and classification of the Asteroidea (Echinodermata). *Zoological Journal of the Linnean Society* 89: 107–132.
- Hart, M. W., Byrne, M., and Smith, M. J. 1997. Molecular phylogenetic analysis of life-history evolution in asterinid starfish. *Evolution* 51: 1848–1861.
- Hayashi, R. 1940. Contributions to the classification of the sea-stars of Japan, I: Spinulosa. *Journal of the Faculty of Science, Hokkaido Imperial University, Series 6, Zoology* 7: 107–204.
- Hayashi, R. 1973. *The Sea-stars of Sagami Bay*. Biological Laboratory, Imperial Household, Tokyo, xi+114 pp., 18 pls.
- Katoh, K. and Standley, D. M. 2013. MAFFT multiple sequence alignment software version 7: improvements in performance and usability. *Molecular Biology and Evolution* 30: 772–780.
- Knott, K. E., Balser, E. J., Jaeckle, W. B., and Wray, G. A. 2003. Identification of asteroid genera with species capable of larval cloning. *Biological Bulletin* 204: 246–255.
- Knott, K. E., Ringvold, H., and Blicher, M. E. 2018. Morphological and molecular analysis of *Henricia* Gray, 1840 (Asteroidea: Echinodermata) from the Northern Atlantic Ocean. *Zoological Journal of the Linnean Society* 182: 791–807.
- Lanfear, R., Calcott, B., Ho, S. Y., and Guindon, S. 2012. PartitionFinder: combined selection of partitioning schemes and substitution models for phylogenetic analyses. *Molecular Biology and Evolution* 29: 1695–1701.
- Layton, K. K., Corstorphine, E. A., and Hebert, P. D. 2016. Exploring Canadian echinoderm diversity through DNA barcodes. *PLOS ONE* 11: e0166118.
- Lopes, E. M., Pérez-Portela, R., Paiva, P. C., and Ventura, C. R. R. 2016. The molecular phylogeny of the sea star *Echinaster* (Asteroidea: Echinasteridae) provides insights for genus taxonomy. *Invertebrate Biology* 135: 235–244.
- Madsen, F. J. 1987. The *Henricia sanguinolenta* complex (Echinodermata: Asteroidea) of the Norwegian Sea and adjacent waters: a re-evaluation, with notes on related species. *Steenstrupia* 13: 201–268.
- Mah, C. L. and Blake, D. B. 2012. Global diversity and phylogeny of the Asteroidea (Echinodermata). *PLoS ONE* 7: e35644.
- Mah, C. L. and Hansson, H. 2017. Echinasteridae. *In*: Mah, C. L. (Ed.) *World Asteroidea Database*. Accessed through: World Register of Marine Species. Available at <http://www.marinespecies.org/aphia.php?p=taxdetails&id=123132> (17 November 2017).
- Matsubara, M., Komatsu, M., and Wada, H. 2004. Close relationship between *Asterina* and Solasteridae (Asteroidea) supported by both nuclear and mitochondrial gene molecular phylogenies. *Zoological Science* 21: 785–793.
- Mortensen, T. 1925. Papers from Dr. Th. Mortensen's Pacific expedition 1914–16, 29: echinoderms of New Zealand and the Auckland-Cambell Islands, 3–5: Asteroidea, Holothurioidea and Crinoidea.

- Videnskabelige Meddelelser fra Dansk Naturhistorisk Forening 79: 261–420.
- Palumbi, S. R. 1996. Nucleic acids II: the polymerase chain reaction. *Molecular Systematics* 2: 205–247.
- Rambaut, A. 2014. FigTree 1.4.2 software. Institute of Evolutionary Biology, University of Edinburgh. Available at <http://tree.bio.ed.ac.uk/software/figtree/> (22 July 2016).
- Rambaut, A., Suchard, M. A., Xie, W., and Drummond, A. J. 2014. Tracer v. 1.6. Institute of Evolutionary Biology, University of Edinburgh. Available at <http://tree.bio.ed.ac.uk/software/tracer/> (22 July 2016).
- Ronquist, F., Teslenko, M., van der Mark, P., Ayres, D. L., Darling, A., Höhna, S., Larget, B., Liu, L., Suchard, M. A., and Huelsenbeck, J. P. 2012. MrBayes 3.2: efficient Bayesian phylogenetic inference and model choice across a large model space. *Systematic Biology* 61: 539–542.
- Rowe, F. E. W. and Albertson, E. L. 1986. The echinoderm genus *Henricia* Gray, 1840 (Asteroidea: Echinasteridae) in southern and southeastern Australian waters, with the description of a new species. *Proceedings of the Linnean Society of New South Wales* 109: 183–194.
- Sladen, W. P. 1889. Report on the Asteroidea collected during the voyage of HMS Challenger during the years 1873–76. Pp. 1–893. In: Thomson, C. W. and Murray, J. (Eds) *Report on the Scientific Results of the Voyage of HMS Challenger During the Years 1873–76, Zoology, Volume 30*. Her Majesty's Government, London.
- Smith, M. J., Arndt, A., Gorski, S., and Fajber, E. 1993. The phylogeny of echinoderm classes based on mitochondrial gene arrangements. *Journal of Molecular Evolution* 36: 545–554.
- Stamatakis, A. 2014. RAxML version 8: a tool for phylogenetic analysis and post-analysis of large phylogenies. *Bioinformatics* 30: 1312–1313.
- Talavera, G. and Castresana, J. 2007. Improvement of phylogenies after removing divergent and ambiguously aligned blocks from protein sequence alignments. *Systematic Biology* 56: 564–577.
- Tamura, K., Stecher, G., Peterson, D., FilipSKI, A., and Kumar, S. 2013. MEGA6: molecular evolutionary genetics analysis version 6.0. *Molecular Biology and Evolution* 30: 2725–2729.
- Tanabe, A. S. 2011. Kakusan4 and Aminosan: two programs for comparing nonpartitioned, proportional and separate models for combined molecular phylogenetic analyses of multilocus sequence data. *Molecular Ecology Resources* 11: 914–921.
- Uchida, T. 1928. Report of the biological survey of Mutsu Bay, 11: starfishes of Mutsu Bay. *Scientific Reports of Tohoku Imperial University* 4: 785–803.
- Verrill, A. E. 1914. *Monograph of the Shallow-water Starfishes of the North Pacific Coast from the Arctic Ocean to California*. Smithsonian Institution, Washington, D. C., xii+408 pp., 110 pls.
- Wada, H., Komatsu, M., and Satoh, N. 1996. Mitochondrial rDNA phylogeny of the Asteroidea suggests the primitiveness of the Paxillo-sida. *Molecular Phylogenetics and Evolution* 6: 97–106.
- Wakita, D., Fujita, T., and Kajihara, H. 2019. Supplementary figures of “Molecular Systematics and Morphological Analyses of the Subgenus *Setihenricia* (Echinodermata: Asteroidea: *Henricia*) from Japan”. figshare. Available at <https://doi.org/10.6084/m9.figshare.7570676.v1> (10 January 2019).

Appendix

Measured values of morphological characters used in this study. Combined ratios are not shown except for DS_{base}/DS_{tip} , AdS_{base}/AdS_{tip} , and $AmP_{width}/AmP_{neck}/r$ (see Figs 6B, 6D, and 4F, respectively) as well as DS_{middle}/DS_{tip} and AdS_{middle}/AdS_{tip} . For each kind of ossicles, values in the same column of each specimen were obtained from the same ossicle. See Table 4 for abbreviations.

Species	Number of rows of spines on each inferomarginal plate																		R	r	
<i>Henricia tumida</i>	3	3	2	3	3	3	3	3	3	3	3	2	3	3	3	3	3	3	3	29	9.0
<i>Henricia</i> sp. 1 (Akkeshi)	3	4	4	3	3	3	3	3	3	4	3	3	3	3	3	3	3	4	3	40	9.0
<i>Henricia nipponica</i> (Muroan)	4	3	3	3	3	3	4	3	4	3	2	3	3	3	3	2	3	3	3	14	4.5
<i>Henricia nipponica</i> (Oshoro)	3	3	3	3	3	3	2	3	3	2	3	3	4	3	3	3	2	3	3	20	6.0
<i>Henricia reniossa</i>	5	5	6	5	5	6	5	6	5	5	5	6	7	6	5	6	6	6	6	64	8.0
<i>Henricia</i> sp. 2 (Abashiri 1)	3	4	5	4	4	5	4	4	4	4	3	4	4	5	4	4	5	5	4	26	7.0
<i>Henricia hayashii</i>	4	4	5	4	4	4	4	4	4	4	4	4	5	5	4	4	5	4	5	57	9.0
<i>Henricia</i> sp. 3 (Abashiri 2)	6	6	6	6	7	7	6	5	6	6	6	7	7	6	6	6	7	6	6	35	7.0
<i>Henricia kinkasana</i>	5	5	6	5	5	4	4	5	5	5	5	5	5	4	4	4	5	4	4	31	6.5
<i>Henricia</i> sp. 4 (Fukushima)	4	4	5	5	6	5	5	6	5	5	5	4	5	5	4	3	4	5	4	45	8.5
<i>Henricia</i> sp. 5 (Okinawa)	4	3	3	4	4	3	4	4	4	4	5	4	4	3	4	4	4	4	5	45	7.5
<i>Henricia</i> sp. 6 (Miyako)	3	3	4	4	4	4	3	4	4	4	4	3	3	4	4	4	3	4	4	60	8.5
<i>Henricia</i> sp. 7 (Nagannu)	4	4	4	4	4	5	4	4	5	4	4	3	4	4	5	4	4	4	3	40	7.5
<i>Henricia regularis</i>	3	4	4	4	3	4	4	4	4	5	4	4	4	5	3	4	4	3	4	27	6.5
<i>Henricia</i> sp. 8 (Okinoerabu)	2	3	3	2	2	2	3	2	2	2	2	3	2	2	2	2	2	2	2	77	10.0
<i>Henricia reticulata</i>	2	2	2	2	2	2	2	2	2	2	1	2	2	2	1	2	2	2	2	83	17.0
<i>Henricia ohshimai</i> (Misaki)	2	2	2	2	2	2	2	2	2	2	3	2	2	2	2	3	2	2	2	56	13.0
<i>Henricia ohshimai</i> (Oshoro)	2	2	2	2	2	3	2	2	2	2	2	2	2	2	2	2	2	2	2	50	11.0

Species	DS_{height}									
<i>Henricia tumida</i>	287	278	260	301	287	249	300	302	283	—
<i>Henricia</i> sp. 1 (Akkeshi)	315	324	246	326	251	287	284	279	381	268
<i>Henricia nipponica</i> (Muroan)	199	216	235	243	211	222	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	282	276	321	290	270	278	274	280	281	263
<i>Henricia reniossa</i>	291	272	324	298	293	300	277	280	315	282
<i>Henricia</i> sp. 2 (Abashiri 1)	372	386	301	—	—	—	—	—	—	—
<i>Henricia hayashii</i>	300	241	244	234	265	305	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	268	319	301	334	301	259	407	295	268	318
<i>Henricia kinkasana</i>	291	315	300	324	321	307	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	446	419	438	424	447	—	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	318	326	324	349	323	309	342	304	318	—
<i>Henricia</i> sp. 6 (Miyako)	335	326	319	330	—	—	—	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	345	353	340	—	—	—	—	—	—	—
<i>Henricia regularis</i>	310	276	302	315	299	318	311	311	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	463	292	298	349	315	255	449	—	—	—
<i>Henricia reticulata</i>	618	510	—	—	—	—	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	433	468	542	521	484	501	526	494	494	498
<i>Henricia ohshimai</i> (Oshoro)	346	342	356	367	375	373	366	356	348	381

Species	DS_{base}									
<i>Henricia tumida</i>	86	90	94	93	90	101	101	93	85	—
<i>Henricia</i> sp. 1 (Akkeshi)	95	93	90	95	76	101	95	96	89	84
<i>Henricia nipponica</i> (Muroan)	39	35	42	39	41	40	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	60	57	55	54	51	59	57	57	59	52
<i>Henricia reniossa</i>	55	56	63	53	61	78	62	69	66	69
<i>Henricia</i> sp. 2 (Abashiri 1)	63	61	47	—	—	—	—	—	—	—
<i>Henricia hayashii</i>	54	55	53	52	51	55	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	31	33	32	35	41	33	43	37	31	31
<i>Henricia kinkasana</i>	39	39	39	37	37	38	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	50	49	47	50	53	—	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	59	45	52	54	59	52	45	49	42	—
<i>Henricia</i> sp. 6 (Miyako)	62	54	51	49	—	—	—	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	42	56	40	—	—	—	—	—	—	—
<i>Henricia regularis</i>	58	58	60	60	60	51	56	61	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	68	49	46	50	58	37	64	—	—	—
<i>Henricia reticulata</i>	137	128	—	—	—	—	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	101	111	138	124	123	115	124	102	116	128
<i>Henricia ohshimai</i> (Oshoro)	96	109	86	104	101	93	117	87	102	102

(Continued)

Species	DS _{middle}									
<i>Henricia tumida</i>	84	81	90	73	82	93	94	88	79	—
<i>Henricia</i> sp. 1 (Akkeshi)	77	68	80	85	67	79	80	81	75	75
<i>Henricia nipponica</i> (Murooran)	37	34	39	35	39	44	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	59	47	46	48	56	53	42	50	57	49
<i>Henricia reniossa</i>	43	34	50	44	39	50	36	48	48	53
<i>Henricia</i> sp. 2 (Abashiri 1)	43	54	33	—	—	—	—	—	—	—
<i>Henricia hayashii</i>	53	56	46	43	55	52	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	26	29	30	30	26	30	39	33	27	29
<i>Henricia kinkasana</i>	30	32	29	31	26	30	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	36	36	38	34	39	—	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	36	29	36	38	36	27	33	34	29	—
<i>Henricia</i> sp. 6 (Miyako)	45	34	43	37	—	—	—	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	30	38	28	—	—	—	—	—	—	—
<i>Henricia regularis</i>	42	38	42	37	37	36	46	41	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	43	27	23	32	38	29	31	—	—	—
<i>Henricia reticulata</i>	100	93	—	—	—	—	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	87	94	135	114	115	115	115	102	111	122
<i>Henricia ohshimai</i> (Oshoro)	87	89	74	78	73	75	87	76	80	77

Species	DS _{tip}									
<i>Henricia tumida</i>	94	90	106	83	91	83	95	102	91	—
<i>Henricia</i> sp. 1 (Akkeshi)	76	75	78	83	56	78	68	63	80	74
<i>Henricia nipponica</i> (Murooran)	42	34	49	40	38	51	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	68	54	49	63	63	58	50	60	76	58
<i>Henricia reniossa</i>	52	36	59	54	49	58	40	62	51	72
<i>Henricia</i> sp. 2 (Abashiri 1)	39	50	32	—	—	—	—	—	—	—
<i>Henricia hayashii</i>	88	70	65	70	77	74	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	28	34	35	30	31	40	45	35	31	32
<i>Henricia kinkasana</i>	33	37	23	34	30	35	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	35	31	36	39	35	—	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	28	34	44	41	45	28	41	38	32	—
<i>Henricia</i> sp. 6 (Miyako)	40	36	32	29	—	—	—	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	29	36	29	—	—	—	—	—	—	—
<i>Henricia regularis</i>	37	37	37	40	39	26	47	50	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	19	9	15	16	23	20	20	—	—	—
<i>Henricia reticulata</i>	76	71	—	—	—	—	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	78	103	117	90	105	92	94	87	106	95
<i>Henricia ohshimai</i> (Oshoro)	68	64	57	68	53	58	69	57	66	62

Species	DS _{base} /DS _{tip}									
<i>Henricia tumida</i>	0.915	1.000	0.887	1.120	0.989	1.217	1.063	0.912	0.934	—
<i>Henricia</i> sp. 1 (Akkeshi)	1.250	1.240	1.154	1.145	1.357	1.295	1.397	1.524	1.113	1.135
<i>Henricia nipponica</i> (Murooran)	0.929	1.029	0.857	0.975	1.079	0.784	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	0.882	1.056	1.122	0.857	0.810	1.017	1.140	0.950	0.776	0.897
<i>Henricia reniossa</i>	1.058	1.556	1.068	0.981	1.245	1.345	1.550	1.113	1.294	0.958
<i>Henricia</i> sp. 2 (Abashiri 1)	1.615	1.220	1.469	—	—	—	—	—	—	—
<i>Henricia hayashii</i>	0.614	0.786	0.815	0.743	0.662	0.743	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	1.107	0.971	0.914	1.167	1.323	0.825	0.956	1.057	1.000	0.969
<i>Henricia kinkasana</i>	1.182	1.054	1.696	1.088	1.233	1.086	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	1.429	1.581	1.306	1.282	1.514	—	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	2.107	1.324	1.182	1.317	1.311	1.857	1.098	1.289	1.313	—
<i>Henricia</i> sp. 6 (Miyako)	1.550	1.500	1.594	1.690	—	—	—	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	1.448	1.556	1.379	—	—	—	—	—	—	—
<i>Henricia regularis</i>	1.568	1.568	1.622	1.500	1.538	1.962	1.191	1.220	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	3.579	5.444	3.067	3.125	2.522	1.850	3.200	—	—	—
<i>Henricia reticulata</i>	1.803	1.803	—	—	—	—	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	1.295	1.078	1.179	1.378	1.171	1.250	1.319	1.172	1.094	1.347
<i>Henricia ohshimai</i> (Oshoro)	1.412	1.703	1.509	1.529	1.906	1.603	1.696	1.526	1.545	1.645

(Continued)

Species	$DS_{\text{middle}}/DS_{\text{tip}}$									
<i>Henricia tumida</i>	0.894	0.900	0.849	0.880	0.901	1.120	0.989	0.863	0.868	—
<i>Henricia</i> sp. 1 (Akkeshi)	1.013	0.907	1.026	1.024	1.196	1.013	1.176	1.286	0.938	1.014
<i>Henricia nipponica</i> (Murooran)	0.881	1.000	0.796	0.875	1.026	0.863	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	0.868	0.870	0.939	0.762	0.889	0.914	0.840	0.833	0.750	0.845
<i>Henricia reniessa</i>	0.827	0.944	0.847	0.815	0.796	0.862	0.900	0.774	0.941	0.736
<i>Henricia</i> sp. 2 (Abashiri 1)	1.103	1.080	1.031	—	—	—	—	—	—	—
<i>Henricia hayashii</i>	0.602	0.800	0.708	0.614	0.714	0.703	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	0.929	0.853	0.857	1.000	0.839	0.750	0.867	0.943	0.871	0.906
<i>Henricia kinkasana</i>	0.909	0.865	1.261	0.912	0.867	0.857	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	1.029	1.161	1.056	0.872	1.114	—	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	1.286	0.853	0.818	0.927	0.800	0.964	0.805	0.895	0.906	—
<i>Henricia</i> sp. 6 (Miyako)	1.125	0.944	1.344	1.276	—	—	—	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	1.034	1.056	0.966	—	—	—	—	—	—	—
<i>Henricia regularis</i>	1.135	1.027	1.135	0.925	0.949	1.385	0.979	0.820	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	2.263	3.000	1.533	2.000	1.652	1.450	1.550	—	—	—
<i>Henricia reticulata</i>	1.316	1.310	—	—	—	—	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	1.115	0.913	1.154	1.267	1.095	1.250	1.223	1.172	1.047	1.284
<i>Henricia ohshimai</i> (Oshoro)	1.279	1.391	1.298	1.147	1.377	1.293	1.261	1.333	1.212	1.242

Species	AdS_{height}				
<i>Henricia tumida</i>	807	714	759	848	—
<i>Henricia</i> sp. 1 (Akkeshi)	977	989	932	1028	1105
<i>Henricia nipponica</i> (Murooran)	473	471	533	538	—
<i>Henricia nipponica</i> (Oshoro)	679	638	665	652	649
<i>Henricia reniessa</i>	1272	1308	1251	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	807	758	758	—	—
<i>Henricia hayashii</i>	1298	1304	1308	1260	1161
<i>Henricia</i> sp. 3 (Abashiri 2)	921	936	941	833	—
<i>Henricia kinkasana</i>	664	754	740	683	704
<i>Henricia</i> sp. 4 (Fukushima)	1033	1113	1120	—	—
<i>Henricia</i> sp. 5 (Okinawa)	1018	1033	965	—	—
<i>Henricia</i> sp. 6 (Miyako)	859	980	866	—	—
<i>Henricia</i> sp. 7 (Nagannu)	1054	1065	—	—	—
<i>Henricia regularis</i>	978	943	951	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	1283	1245	1282	1258	—
<i>Henricia reticulata</i>	2091	2007	2043	—	—
<i>Henricia ohshimai</i> (Misaki)	1911	1959	1956	—	—
<i>Henricia ohshimai</i> (Oshoro)	1548	1536	1547	1482	1562

Species	AdS_{base}				
<i>Henricia tumida</i>	223	178	214	210	—
<i>Henricia</i> sp. 1 (Akkeshi)	181	178	178	203	215
<i>Henricia nipponica</i> (Murooran)	89	101	121	101	—
<i>Henricia nipponica</i> (Oshoro)	139	141	136	125	141
<i>Henricia reniessa</i>	181	149	144	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	102	95	95	—	—
<i>Henricia hayashii</i>	210	176	216	171	197
<i>Henricia</i> sp. 3 (Abashiri 2)	123	123	116	110	—
<i>Henricia kinkasana</i>	101	115	105	120	110
<i>Henricia</i> sp. 4 (Fukushima)	142	127	130	—	—
<i>Henricia</i> sp. 5 (Okinawa)	115	106	95	—	—
<i>Henricia</i> sp. 6 (Miyako)	136	150	129	—	—
<i>Henricia</i> sp. 7 (Nagannu)	113	116	—	—	—
<i>Henricia regularis</i>	147	115	126	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	206	219	216	208	—
<i>Henricia reticulata</i>	355	327	370	—	—
<i>Henricia ohshimai</i> (Misaki)	354	341	355	—	—
<i>Henricia ohshimai</i> (Oshoro)	300	296	316	281	316

(Continued)

Species	AdS _{middle}				
<i>Henricia tumida</i>	202	166	191	178	—
<i>Henricia</i> sp. 1 (Akkeshi)	135	139	188	203	196
<i>Henricia nipponica</i> (Murooran)	83	86	112	92	—
<i>Henricia nipponica</i> (Oshoro)	129	128	118	125	124
<i>Henricia reniossa</i>	153	149	135	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	89	82	75	—	—
<i>Henricia hayashii</i>	203	180	183	206	169
<i>Henricia</i> sp. 3 (Abashiri 2)	106	106	103	90	—
<i>Henricia kinkasana</i>	78	89	71	73	78
<i>Henricia</i> sp. 4 (Fukushima)	110	117	120	—	—
<i>Henricia</i> sp. 5 (Okinawa)	102	111	98	—	—
<i>Henricia</i> sp. 6 (Miyako)	100	121	116	—	—
<i>Henricia</i> sp. 7 (Nagannu)	88	100	—	—	—
<i>Henricia regularis</i>	119	111	114	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	163	178	203	165	—
<i>Henricia reticulata</i>	276	295	318	—	—
<i>Henricia ohshimai</i> (Misaki)	363	377	350	—	—
<i>Henricia ohshimai</i> (Oshoro)	271	234	230	251	302

Species	AdS _{tip}				
<i>Henricia tumida</i>	252	198	246	230	—
<i>Henricia</i> sp. 1 (Akkeshi)	154	156	244	271	277
<i>Henricia nipponica</i> (Murooran)	91	102	111	105	—
<i>Henricia nipponica</i> (Oshoro)	156	163	148	139	143
<i>Henricia reniossa</i>	163	170	159	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	95	99	84	—	—
<i>Henricia hayashii</i>	261	226	262	234	244
<i>Henricia</i> sp. 3 (Abashiri 2)	161	150	148	137	—
<i>Henricia kinkasana</i>	93	104	102	105	91
<i>Henricia</i> sp. 4 (Fukushima)	120	116	119	—	—
<i>Henricia</i> sp. 5 (Okinawa)	95	104	107	—	—
<i>Henricia</i> sp. 6 (Miyako)	110	150	127	—	—
<i>Henricia</i> sp. 7 (Nagannu)	102	100	—	—	—
<i>Henricia regularis</i>	126	107	126	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	130	143	177	136	—
<i>Henricia reticulata</i>	290	291	327	—	—
<i>Henricia ohshimai</i> (Misaki)	395	377	350	—	—
<i>Henricia ohshimai</i> (Oshoro)	264	224	236	238	285

Species	AdS _{base} /AdS _{tip}				
<i>Henricia tumida</i>	0.885	0.899	0.870	0.913	—
<i>Henricia</i> sp. 1 (Akkeshi)	1.175	1.141	0.730	0.749	0.776
<i>Henricia nipponica</i> (Murooran)	0.978	0.990	1.090	0.962	—
<i>Henricia nipponica</i> (Oshoro)	0.891	0.865	0.919	0.899	0.986
<i>Henricia reniossa</i>	1.110	0.876	0.906	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	1.074	0.960	1.131	—	—
<i>Henricia hayashii</i>	0.805	0.779	0.824	0.731	0.807
<i>Henricia</i> sp. 3 (Abashiri 2)	0.764	0.820	0.784	0.803	—
<i>Henricia kinkasana</i>	1.086	1.106	1.029	1.143	1.209
<i>Henricia</i> sp. 4 (Fukushima)	1.183	1.095	1.092	—	—
<i>Henricia</i> sp. 5 (Okinawa)	1.211	1.019	0.888	—	—
<i>Henricia</i> sp. 6 (Miyako)	1.236	1.000	1.016	—	—
<i>Henricia</i> sp. 7 (Nagannu)	1.108	1.160	—	—	—
<i>Henricia regularis</i>	1.167	1.075	1.000	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	1.585	1.531	1.220	1.529	—
<i>Henricia reticulata</i>	1.224	1.124	1.131	—	—
<i>Henricia ohshimai</i> (Misaki)	0.896	0.905	1.014	—	—
<i>Henricia ohshimai</i> (Oshoro)	1.136	1.321	1.339	1.181	1.109

(Continued)

Species	AdS _{middle} /AdS _{tip}				
<i>Henricia tumida</i>	0.802	0.838	0.776	0.774	—
<i>Henricia</i> sp. 1 (Akkeshi)	0.877	0.891	0.770	0.749	0.708
<i>Henricia nipponica</i> (Murooran)	0.912	0.843	1.009	0.876	—
<i>Henricia nipponica</i> (Oshoro)	0.827	0.785	0.797	0.899	0.867
<i>Henricia reniessa</i>	0.939	0.876	0.849	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	0.937	0.828	0.893	—	—
<i>Henricia hayashii</i>	0.778	0.796	0.698	0.880	0.693
<i>Henricia</i> sp. 3 (Abashiri 2)	0.658	0.707	0.696	0.657	—
<i>Henricia kinkasana</i>	0.839	0.856	0.696	0.695	0.857
<i>Henricia</i> sp. 4 (Fukushima)	0.917	1.009	1.008	—	—
<i>Henricia</i> sp. 5 (Okinawa)	1.074	1.067	0.916	—	—
<i>Henricia</i> sp. 6 (Miyako)	0.909	0.807	0.913	—	—
<i>Henricia</i> sp. 7 (Nagannu)	0.863	1.000	—	—	—
<i>Henricia regularis</i>	0.944	1.037	0.905	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	1.254	1.245	1.147	1.213	—
<i>Henricia reticulata</i>	0.952	1.014	0.972	—	—
<i>Henricia ohshimai</i> (Misaki)	0.919	1.000	1.000	—	—
<i>Henricia ohshimai</i> (Oshoro)	1.027	1.045	0.975	1.055	1.060

Species	AdP _{height}							
<i>Henricia tumida</i>	989	976	995	954	1050	982	—	—
<i>Henricia</i> sp. 1 (Akkeshi)	977	959	967	965	—	—	—	—
<i>Henricia nipponica</i> (Murooran)	558	547	543	—	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	708	678	740	690	—	—	—	—
<i>Henricia reniessa</i>	1289	1326	1211	1277	1215	—	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	825	778	813	773	—	—	—	—
<i>Henricia hayashii</i>	1206	1128	—	—	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	918	826	934	—	—	—	—	—
<i>Henricia kinkasana</i>	792	791	803	791	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	888	914	973	1021	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	925	893	924	1012	—	—	—	—
<i>Henricia</i> sp. 6 (Miyako)	1108	1055	1124	1150	1134	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	994	934	946	—	—	—	—	—
<i>Henricia regularis</i>	777	787	804	726	777	—	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	1627	1666	1562	1601	1564	1495	1552	1601
<i>Henricia reticulata</i>	1988	1617	1800	1728	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	1622	1610	1316	1597	1546	—	—	—
<i>Henricia ohshimai</i> (Oshoro)	1395	1267	1315	1290	—	—	—	—

Species	AdP _{width}							
<i>Henricia tumida</i>	994	1007	990	1040	1031	1010	—	—
<i>Henricia</i> sp. 1 (Akkeshi)	1121	1055	1090	1131	—	—	—	—
<i>Henricia nipponica</i> (Murooran)	534	532	512	—	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	690	634	666	658	—	—	—	—
<i>Henricia reniessa</i>	1266	1154	1201	1234	1145	—	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	818	855	815	827	—	—	—	—
<i>Henricia hayashii</i>	1100	1055	—	—	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	964	894	932	—	—	—	—	—
<i>Henricia kinkasana</i>	798	791	765	789	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	815	820	849	856	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	909	930	926	901	—	—	—	—
<i>Henricia</i> sp. 6 (Miyako)	1007	976	993	1006	965	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	906	885	902	—	—	—	—	—
<i>Henricia regularis</i>	757	732	785	717	779	—	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	1870	1896	1666	1879	1945	1914	1769	1853
<i>Henricia reticulata</i>	1792	1916	2000	1923	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	1561	1545	1474	1592	1645	—	—	—
<i>Henricia ohshimai</i> (Oshoro)	1275	1399	1456	1384	—	—	—	—

(Continued)

Species	AdP _{top}							
<i>Henricia tumida</i>	772	823	801	802	849	802	—	—
<i>Henricia</i> sp. 1 (Akkeshi)	905	820	857	911	—	—	—	—
<i>Henricia nipponica</i> (Muroan)	428	401	441	—	—	—	—	—
<i>Henricia nipponica</i> (Oshoro)	490	493	527	534	—	—	—	—
<i>Henricia reniessa</i>	922	941	988	1032	981	—	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	632	629	452	611	—	—	—	—
<i>Henricia hayashii</i>	954	869	—	—	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	720	677	733	—	—	—	—	—
<i>Henricia kinkasana</i>	601	781	520	622	—	—	—	—
<i>Henricia</i> sp. 4 (Fukushima)	604	628	747	741	—	—	—	—
<i>Henricia</i> sp. 5 (Okinawa)	679	677	620	666	—	—	—	—
<i>Henricia</i> sp. 6 (Miyako)	657	756	838	827	736	—	—	—
<i>Henricia</i> sp. 7 (Nagannu)	732	592	589	—	—	—	—	—
<i>Henricia regularis</i>	608	583	572	616	581	—	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	1545	1549	1458	1545	1612	1613	1493	1618
<i>Henricia reticulata</i>	1543	1545	1420	1682	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	1303	1249	1271	1124	1165	—	—	—
<i>Henricia ohshimai</i> (Oshoro)	1087	1068	909	1064	—	—	—	—

Species	AmP _{width}						
<i>Henricia tumida</i>	549	582	620	—	—	—	—
<i>Henricia</i> sp. 1 (Akkeshi)	661	635	631	737	—	—	—
<i>Henricia nipponica</i> (Muroan)	354	412	378	364	350	317	—
<i>Henricia nipponica</i> (Oshoro)	401	459	456	—	—	—	—
<i>Henricia reniessa</i>	755	762	774	815	775	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	487	440	434	—	—	—	—
<i>Henricia hayashii</i>	632	737	677	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	581	560	597	—	—	—	—
<i>Henricia kinkasana</i>	476	440	471	409	342	—	—
<i>Henricia</i> sp. 4 (Fukushima)	522	533	546	538	529	—	—
<i>Henricia</i> sp. 5 (Okinawa)	492	502	487	494	—	—	—
<i>Henricia</i> sp. 6 (Miyako)	519	573	525	582	564	—	—
<i>Henricia</i> sp. 7 (Nagannu)	458	441	472	518	—	—	—
<i>Henricia regularis</i>	465	459	495	454	—	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	878	727	761	664	—	—	—
<i>Henricia reticulata</i>	1015	1109	1154	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	971	958	891	—	—	—	—
<i>Henricia ohshimai</i> (Oshoro)	918	1000	862	—	—	—	—

Species	AmP _{ridge}						
<i>Henricia tumida</i>	838	861	883	—	—	—	—
<i>Henricia</i> sp. 1 (Akkeshi)	887	876	992	870	—	—	—
<i>Henricia nipponica</i> (Muroan)	467	527	531	441	528	458	—
<i>Henricia nipponica</i> (Oshoro)	623	600	709	—	—	—	—
<i>Henricia reniessa</i>	1342	1397	1323	1333	1349	—	—
<i>Henricia</i> sp. 2 (Abashiri 1)	869	777	919	—	—	—	—
<i>Henricia hayashii</i>	1358	1396	1275	—	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	1096	1090	1020	—	—	—	—
<i>Henricia kinkasana</i>	1030	953	1071	997	1001	—	—
<i>Henricia</i> sp. 4 (Fukushima)	1111	1180	1084	1149	1106	—	—
<i>Henricia</i> sp. 5 (Okinawa)	988	953	982	1033	—	—	—
<i>Henricia</i> sp. 6 (Miyako)	1169	1169	1078	1178	1110	—	—
<i>Henricia</i> sp. 7 (Nagannu)	880	876	870	936	—	—	—
<i>Henricia regularis</i>	844	837	797	761	—	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	1525	1661	1545	1410	—	—	—
<i>Henricia reticulata</i>	1649	1681	1683	—	—	—	—
<i>Henricia ohshimai</i> (Misaki)	1703	1656	1773	—	—	—	—
<i>Henricia ohshimai</i> (Oshoro)	1251	1420	1439	—	—	—	—

(Continued)

Species	AmP _{neck}					
<i>Henricia tumida</i>	458	496	511	—	—	—
<i>Henricia</i> sp. 1 (Akkeshi)	560	549	556	611	—	—
<i>Henricia nipponica</i> (Muroan)	327	361	340	329	321	290
<i>Henricia nipponica</i> (Oshoro)	366	415	400	—	—	—
<i>Henricia reniossa</i>	684	664	692	707	696	—
<i>Henricia</i> sp. 2 (Abashiri 1)	448	417	404	—	—	—
<i>Henricia hayashii</i>	549	571	565	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	525	504	515	—	—	—
<i>Henricia kinkasana</i>	370	356	371	338	313	—
<i>Henricia</i> sp. 4 (Fukushima)	387	416	403	412	400	—
<i>Henricia</i> sp. 5 (Okinawa)	470	451	474	400	—	—
<i>Henricia</i> sp. 6 (Miyako)	464	473	471	487	491	—
<i>Henricia</i> sp. 7 (Nagannu)	440	441	472	482	—	—
<i>Henricia regularis</i>	438	428	451	401	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	805	689	739	608	—	—
<i>Henricia reticulata</i>	908	1042	1102	—	—	—
<i>Henricia ohshimai</i> (Misaki)	903	850	831	—	—	—
<i>Henricia ohshimai</i> (Oshoro)	841	943	835	—	—	—

Species	AmP _{width} /AmP _{neck} /r					
<i>Henricia tumida</i>	0.1332	0.1304	0.1348	—	—	—
<i>Henricia</i> sp. 1 (Akkeshi)	0.1312	0.1285	0.1261	0.1340	—	—
<i>Henricia nipponica</i> (Muroan)	0.2406	0.2536	0.2471	0.2459	0.2423	0.2429
<i>Henricia nipponica</i> (Oshoro)	0.1826	0.1843	0.1900	—	—	—
<i>Henricia reniossa</i>	0.1380	0.1434	0.1398	0.1441	0.1392	—
<i>Henricia</i> sp. 2 (Abashiri 1)	0.1553	0.1507	0.1535	—	—	—
<i>Henricia hayashii</i>	0.1279	0.1434	0.1331	—	—	—
<i>Henricia</i> sp. 3 (Abashiri 2)	0.1581	0.1587	0.1656	—	—	—
<i>Henricia kinkasana</i>	0.1979	0.1901	0.1953	0.1862	0.1681	—
<i>Henricia</i> sp. 4 (Fukushima)	0.1587	0.1507	0.1594	0.1536	0.1556	—
<i>Henricia</i> sp. 5 (Okinawa)	0.1396	0.1484	0.1370	0.1647	—	—
<i>Henricia</i> sp. 6 (Miyako)	0.1316	0.1425	0.1311	0.1406	0.1351	—
<i>Henricia</i> sp. 7 (Nagannu)	0.1388	0.1333	0.1333	0.1433	—	—
<i>Henricia regularis</i>	0.1633	0.1650	0.1689	0.1742	—	—
<i>Henricia</i> sp. 8 (Okinoerabu)	0.1091	0.1055	0.1030	0.1092	—	—
<i>Henricia reticulata</i>	0.0658	0.0626	0.0616	—	—	—
<i>Henricia ohshimai</i> (Misaki)	0.0827	0.0867	0.0825	—	—	—
<i>Henricia ohshimai</i> (Oshoro)	0.0992	0.0964	0.0938	—	—	—