**Abstract.** Meiofauna sampling in the proximity of Syd-Hällsö Island (Strömstad, Sweden) revealed a new species of Kinorhyncha from the Skagerrak. The species, *Setaphyes elenae* sp. nov., is distinguished from its congeners by the arrangement of the middorsal cuticular specializations (it has shortened, distally rounded middorsal processes on segments 1 and 9 and middorsal elevations throughout segments 2–8), as well as by the presence of paired laterodorsal setae on segments 3, 5, 7 and 9 and ventromedial setae on segments 3, 5 and 7 in both males and females. The finding of a new species from the north-eastern Atlantic Ocean, provides new valuable information for the recently established genus in the Allomalorhagida.

**Keywords.** Meiofauna, taxonomy, Scalidophora, biodiversity, invertebrates.

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**Introduction**

The phylum Kinorhyncha Reinhard, 1887 encompasses a group of meiobenthic, free-living invertebrates that mainly inhabit the upper centimetres of marine and estuarine sediments, although some species have been found living in hard substrata or associated with macroalgae and marine phanerogams (Higgins 1988). Kinorhynchs have been described from shallow to deep-sea waters across the worldwide oceans,
but certain areas have received much more attention to the detriment of others due to the specialists sampling strategies (Sánchez et al. 2012; Neuhaus 2013; Sørensen et al. 2013; Cepeda et al. 2019).

Kinorhynch biodiversity of the North Sea, together with other north-eastern Atlantic areas nearby, has been studied earlier (Neuhaus, 2013). Currently, 20 species are known in this area: *Campyloderes vanhoeffeni* Zelinka, 1913, *Centroderes spinosus* (Reinhard, 1881), *Condyloperes multispinosus* (McIntyre, 1962), *Echinoderes dujardinii* Claparède, 1863, *E. elongatus* (Nyholm, 1947), *E. higginsi* Huys & Coomans, 1989, *E. levanderi* Karling, 1955, *E. peterseni* Higgins & Kristensen, 1988, *E. setiger* (Greeff, 1869), *E. subfuscus* Zelinka, 1928, *E. worthingi* Southern, 1914, *Paracentrophyes quadridentatus* (Zelinka, 1928), *Pycnophyes calmani* (Zelinka in Southern, 1914), *P. communis* Zelinka, 1908, *P. zelinkaei* Southern, 1914, *Semnoderes armiger* Zelinka, 1928, *Setaphyes dentatus* (Reinhard, 1881), *S. flaveolatus* (Zelinka, 1928), *S. kielensis* (Zelinka, 1928) and *Zelinkaderes submersus* (Gerlach, 1969) (Zelinka 1928; Nyholm 1947; McIntyre et al. 1962; Gerlach 1969; Huys & Coomans 1989; Sørensen et al. 2009; Neuhaus et al. 2013; Altenburger 2016).

In the context of global change and decreasing studies of taxonomy, especially those of small-sized taxa, there is a need to improve taxonomic information about meiofaunal organisms, even in geographic areas that are supposed to be relatively well-studied (Mora et al. 2011; Sørensen & Grzelak 2018). A recent sampling done in the proximities of Syd-Hällsö Island (Strömstad, Sweden) revealed a new species of Kinorhyncha from Skagerrak. Additionally, the species *Echinoderes* cf. *eximus* and *Pycnophyes ancalagon* Sørensen & Grzelak, 2018, the latter recently described from the Svalbard Archipelago (Arctic Ocean), are firstly reported for the boreal area.

**Material and methods**

**Study site**

Sampling was undertaken at a single locality near Syd-Hällsö Island (Strömstad, Sweden) during summer 2017, at the eastern-most limit of the North Sea: 58°56.846′ N, 11°4.896′ E (Fig. 1B). This area encompasses the Skagerrak Strait that connects the North Sea with the Kattegat sea region (Fig. 1A).

Skagerrak, with a surface area of about 32 000 km², is the deepest area of the North Sea basin, with a maximum depth of 700 m (Weering et al. 1993; Rosenberg et al. 1996). Skagerrak is dominated by a deep-reaching flow of water from the central and northern North Sea, with a salinity of about 35 psu and to a lesser extent by a weaker inflow from the southern North Sea with a salinity of about 31–34 psu influenced by river inputs (Rosenberg et al. 1996). Additionally, the shallowest waters of Skagerrak are also subject of upwelling events that cause low-saline currents along the Swedish and Norwegian coastlines (Rodhe 1996).

Skagerrak forms a natural topographic sediment trap, receiving inputs from the entire north-western European drainage systems and the North Sea shoreline (Weering et al. 1993). Thus, Skagerrak has been evidenced as a major depository of fine-grained sediments in the North Sea, with 59% of particles with an average size less than 63 μm (Weering et al. 1987). Nevertheless, areas with coarse gravel, sandy coves and sandbars are also present in the area (Curini-Galletti et al. 2012).

**Sampling and specimen preparation**

Sampling was done during the summer of 2017 by dredging. The collected sample was taken at a depth of 55–65 m and mainly consisted of very fine, soft mud. Sediment was kept in large plastic boxes and stored at a constant temperature of 15°C. Meiofaunal organisms were then extracted by MgCl₂-decantation and live material was studied and sorted using stereo microscopes.
Kinorhynchs were preserved in 100% ethanol. For light microscopy (LM), unmounted specimens were dehydrated through a graded series of glycerine. After being kept in 100% glycerine for 24 h, kinorhynchs were mounted on glass slides with Fluoromount G® sealed with Depex®. Mounted specimens were studied with an Olympus® BX51-P microscope with differential interference contrast (DIC) optics equipped with an Olympus® DP-70 camera. Identification to genus level was done using the dichotomous keys provided by Sørensen & Pardos (2008) for cyclorhagids and the genus diagnoses provided by Sánchez et al. (2016) for allomalorhagids. For scanning electron microscopy (SEM), some unmounted kinorhynchs were sonically cleaned during 10–15 s and led to chemical point drying using a hexamethyldisilazane-ethanol series. Finally, specimens were coated with gold and mounted on aluminium stubs to be studied with a JSM® 6335-F JEOL SEM at the ICTS Centro Nacional de Microscopía Electrónica (UCM, Spain). Type material was deposited at the Natural History Museum of Denmark (NHMD). Line drawings and image plates composition were done using Adobe® Photoshop and Illustrator CC-2014 software.

Morphometric statistical analyses
Differences in selected morphometric measures (total trunk length, standard width and lateral terminal spines’ length) of *Setaphyes elenae* sp. nov. and its congeners were tested. For this, we selected several specimens of *S. dentatus* (n = 18) and *S. flaveolatus* (n = 14), which are the most morphologically similar, and are also distributed through the north-eastern Atlantic Ocean. Specimens of *S. dentatus* and *S. flaveolatus* are from several Atlantic and Mediterranean locations surrounding the Iberian Peninsula, stored at the Meiofauna Collection of the UCM.

Normality and homoscedasticity of the variables were tested using the Saphiro-Wilk’s test (together with visual methods of density and Q–Q plots) and the Bartlett’s test, respectively. Tukey multiple comparison test and pairwise comparisons between group levels with corrections for multiple testing were used to determine which means amongst the set of means differ from the rest. A one-way analysis of variance (ANOVA) was used to test differences, but when heteroscedasticity was detected, a Welch’s

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**Fig. 1.** A. Map showing the sampling area of Syd-Hällsö Island (Strömstad, Sweden). B. Detail. Red point shows the specific sampling point.
ANOVA was performed instead. All the statistical analyses were run in R ver. 1.1.453 using the ‘stats’ basic package and the ‘car’ package ver. 3.0.5 (Fox et al. 2019).

**Results**

Five species co-occur in the collected sample near Syd-Hällsö Island (Swedish North Sea): *Centroderes spinosus*, *Echinoderes cf. eximus*, *Pycnophyes ancalagon*, *Semnoderes armiger* and *Setaphyes elenae* sp. nov.

Class Allomalorhagida Sørensen *et al.*, 2015

Family Pycnophyidae Zelinka, 1896

Genus *Setaphyes* Sánchez *et al.*, 2016

*Setaphyes elenae* sp. nov. [urn:lsid:zoobank.org:act:6C4FE50F-E39A-451E-B805-60B709B9B9B4](urn:lsid:zoobank.org:act:6C4FE50F-E39A-451E-B805-60B709B9B9B4)

Figs 2–4, Tables 1–3

**Diagnosis**

*Setaphyes* with shortened, distally rounded middorsal processes on segments 1 and 9, and middorsal elevations on segments 2–9, superficially covered by tufts of elongated, thick hairs whose tips sometimes surpass the posterior margin of segment. Unpaired setae in paradorsal position on segments 1–9. Laterodorsal setae on segments 3, 5, 7 and 9; paralateral setae absent. Lateroventral setae on segments 2–10. Ventromedial setae on segments 3, 5 and 7. Paired, small, dot-shaped intracuticular structures (maybe outlets of glandular cells) present in several positions throughout the trunk, with a specific arrangement that differs from males to females. Males with paired, sexually dimorphic ventromedial tubes on segment 2, and females with paired, sexually dimorphic ventrolateral setae on segment 2. Lateral terminal spines present, relatively short, slender. Segment 11 retractable into segment 10.

**Etymology**

The species is dedicated to Ms Elena González, sister of the second author.

**Material examined**

**Holotype**

ATLANTIC OCEAN • ♀ adult (mounted in Fluoromount G®); near Syd-Hällsö Island, Skagerrak (Fig. 1B); 58°56.846′ N, 11°4.896′ E; 55–65 m depth; Ulf Jondelius and Fredrik Pleijel leg.; very fine mud; NHMD 655358.

**Paratypes**

ATLANTIC OCEAN • 3 adult ♂♂, 2 adult ♀♀ (all mounted in Fluoromount G®); same collection data as for holotype; NHMD 655359 to 655363.

**Additional non-type material**

ATLANTIC OCEAN • 8 specs (four mounted for LM and four mounted for SEM); same collection data as for holotype; Meiofauna Collection UCM.

**Description**

See Table 1 for measurements and dimensions, Table 2 for summary of cuticular elevation, process, seta, tube, nephridiopore and sensory spot locations, and Table 3 for summary of intracuticular, dot-shaped structure locations.
Head with retractable mouth cone and introvert. The collected specimens were not suitable for head examinations, hence data on number and arrangement of scalids and oral styles are not available.

Neck with four dorsal and two ventral sclerotized placids (Fig. 2A–B, D). Dorsal placids rectangular, with a slightly convex anterior margin; mesial ones broader than lateral ones (Fig. 2B). Ventral placids morphologically similar to dorsal ones but much more elongated, getting thinner towards the lateral sides (Fig. 2A, D).

Trunk with eleven segments (Figs 2A–B, 3A, H, 4A). Segment 1 with one tergal, two episternal and one sub-trapezoidal, midsternal plate; remaining ones with one tergal and two sternal cuticular plates (Figs 2A–D, 3A, H). Tergal cuticular plates slightly bulging middorsally. Sternal plates reach their maximum width at segment 5, but are almost constant in width across the trunk, slightly tapering at the last three trunk segments (Figs 2A–B, 3A, H). Sternal cuticular plates are relatively narrow in the ratio maximum width to total trunk length (MSW-5: TL average ratio = 27.2%), giving the animal a slender appearance (Figs 2A–B, 3A, H, 4A). Middorsal processes on segments 1 and 9, shortened and distally rounded (Figs 2B, 3B, M, 4G, I); middorsal elevations on segments 2–8, pentagonally-shaped, distally rounded, with intracuticular, butterfly-like atria of paradorsal sensory spots (Figs 2B, 3D, F, I, K, 4B, E). Middorsal elevations superficially covered by tufts of elongated, thick cuticular hairs whose tips

| Character | Range ♀ | Mean ♀ | Range ♂ | Mean ♂ | Total range | Total mean | Sd |
|-----------|---------|--------|---------|--------|-------------|------------|----|
| TL (µm)   | 612.67–722.64 | 679.192 | 614.32–647.86 | 625.51 | 612.67–722.64 | 655.33 | 44.06 |
| MSW–5 (µm) | 179.55–188.89 | 183.422 | 168.06–176.25 | 170.72 | 168.06–188.89 | 177.78 | 7.61 |
| MSW–5/TL (%) | 26–29.5 | 27 | 26.2–28.4 | 27.3 | 26–29.5 | 27.2 | 1.22 |
| SSSW (µm) | 141.41–152.44 | 144.178 | 124.68–133.65 | 131.065 | 124.68–152.44 | 138.35 | 8.23 |
| SSSW/TL (%) | 19.7–23 | 21.3 | 19.2–21.9 | 21 | 19.2–23 | 21.1 | 1.14 |
| S1 (µm)   | 87.74–108.41 | 96.584 | 89.16–96.28 | 91.548 | 87.74–108.41 | 94.35 | 6.31 |
| S2 (µm)   | 59.74–82.08 | 69.712 | 62.45–68.07 | 65.068 | 59.74–82.08 | 67.65 | 6.42 |
| S3 (µm)   | 57.56–79.22 | 70.068 | 63.54–68.88 | 66.655 | 57.56–79.22 | 68.55 | 6.04 |
| S4 (µm)   | 70.27–79.51 | 74.618 | 68.79–81.68 | 74.51 | 68.79–81.68 | 74.57 | 4.48 |
| S5 (µm)   | 69.77–83.01 | 77.1 | 69.3–76.08 | 74.438 | 69.3–83.01 | 75.92 | 5.84 |
| S6 (µm)   | 73.96–81.58 | 78.504 | 73.81–83.53 | 77.013 | 73.81–83.53 | 77.84 | 3.51 |
| S7 (µm)   | 74.47–85.94 | 80.752 | 73.33–80.37 | 75.385 | 73.33–85.94 | 78.37 | 4.70 |
| S8 (µm)   | 69.57–86.82 | 80.846 | 77.08–79.98 | 78.138 | 69.57–86.82 | 79.64 | 5.31 |
| S9 (µm)   | 70.24–84.44 | 79.61 | 68.76–78.56 | 74.463 | 68.76–84.44 | 77.32 | 5.43 |
| S10 (µm)  | 64.87–77.97 | 73.66 | 60.93–71.98 | 67.165 | 60.93–77.97 | 70.77 | 5.92 |
| S11 (µm)  | 27.93–39.81 | 32.3 | 27.49–34.23 | 30.615 | 27.49–39.81 | 31.55 | 3.74 |
| LTS (µm)  | 90.64–105.7 | 100.9 | 149.06–168.74 | 158.928 | 90.64–168.74 | 126.69 | 31.29 |
| LTS/TL (%) | 13.9–15.8 | 14.9 | 24–26 | 25.4 | 13.9–26 | 19.6 | 5.60 |

Table 1. Measurements of nine adult specimens of *Setaphyes elenae* sp. nov. (four males and five females) from Skagerrak. Abbreviations: LTS = lateral terminal spines; MSW = maximum sternal width (measured at segment 5); S = segments’ length (followed by number of corresponding segment); Sd = standard deviation; SSSW = standard sternal width (measured at segment 10); TL = total length.
sometimes surpass the posterior margin of the segments (Fig. 4B, E). Intracuticular, minute, dot-shaped, rounded to oval structures (maybe outlets of glandular cells) throughout the cuticle on segments 1–10 (Figs 2A–D, 3B–G, I–N). Location and pairs of these structures per segment differ from males to females (Table 3), and deviations from the bilateral symmetry of their arrangement have been observed in some specimens. Up to three pairs of conspicuous laterodorsal cuticular ridges on segments 2–10 (Figs 2A–B, 3D–G, I–N). Cuticular hairs acicular, elongated, emerging from oval perforation sites, distributed all over the trunk cuticle. Pachycycli and ball-and-socket joints only conspicuous on segments 2–3, reduced on posterior segments (Fig. 2A–D). Apodemes on segments 7–10 (Fig. 2A–C). Primary pectinate fringe finely serrated; secondary pectinate fringe as a double tranverse, hairy-like, wavy row; free flaps covering the anterior part of subsequent segment (Figs 2A–D, 4D). Muscular scars as rounded to oval, hairless areas in laterodorsal and ventromedial positions on segments 1–10 (those of segment 1 in subdorsal and ventrolateral positions), quite inconspicuous (Fig. 2A–D).

Segment 1 with shortened, distally rounded middorsal process still extending beyond the posterior margin of segment (Figs 2B, 3B, 4G). Anterolateral margins of the tergal plate as horn-shaped, short, wide, distally curved and pointed extensions (Figs 2A–B, D, 3A–C, H, 4A). Anterior margin of segment with a reticule-like ornamentation dorsally (Figs 2B, 3B, 4C). Setae absent. Two pairs of sensory spots

Table 2. Summary of nature and arrangement of cuticular elevations, processes, spines, tubes, setae, sensory spots and nephridiopores in Setaphyes elenae sp. nov. Abbreviations: ce = cuticular elevation; cp = cuticular process; LD = laterodorsal; lts = lateral terminal spine; LV = lateroventral; MD = middorsal; ne = nephridiopore; PD = paradorsal; SD = subdorsal; se = seta; ss = sensory spot; tu = tube; VL = ventrolateral; VM = ventromedial. * indicates unpaired structures.

| Segment | MD | PD | SD | LD | LV | VL | VM |
|---------|----|----|----|----|----|----|----|
| 1       | cp* | ss | ss | ss | ss | ss | ss |
| 2       | ce* | se*, ss | ss | ss | se | se (♀) | ss, tu (♂) |
| 3       | ce* | se*, ss | ss | ss | se | se | se |
| 4       | ce* | se*, ss | ss | ss | se | se | ss |
| 5       | ce* | se*, ss | ss | ss | se | se | se |
| 6       | ce* | se*, ss | ss | ss | se | se | ss |
| 7       | ce* | se*, ss | ss | ss | se | se | se |
| 8       | ce* | se*, ss | ss | ss | se | se | ss |
| 9       | cp* | se*, ss | ss | ss, ss | se | se, ne | ss |
| 10      | ss, ss | ss | ss, ss | se | ss |
| 11      |    |     | ls |     |    |    |    |

Fig. 2 (opposite page). Line art illustrations of adult Setaphyes elenae sp. nov. A. ♀, ventral overview. B. ♂, dorsal overview. C. ♀, segments 10–11, ventral view. D. ♀, segments 1–2, ventral view. Abbreviations: bsj = ball-and-socket joint; dpl = dorsal placid; gcoI = type I glandular cell outlet; ldcr = laterodorsal cuticular ridge; ldse = laterodorsal seta; ldss = laterodorsal sensory spot; lts = lateral terminal spine; lvse = lateroventral seta; mde = middorsal elevation; mdp = middorsal process; ms = muscular scar; ps = penil spine; ppf = primary pectinate fringe; psf = secondary pectinate fringe; vlcr = ventrolateral cuticular ridge; vls = ventrolateral placid; vmt = ventromedial tube; vpl = ventral placid. Scale bar = 100 µm.
in ventrolateral position longitudinally arranged, and one pair in paradorsal, subdorsal and laterodorsal positions (Figs 2A–B, D, 3B–C, 4G). Sensory spots on this and subsequent segments as oval areas with several rows of cuticular micropapillae surrounding a single pore (Fig. 4B, G, J).

Segment 2 with middorsal elevation not projecting beyond the posterior margin of segment (Figs 2B, 3D, 4E). Unpaired seta in paradorsal position, on this and following segments indifferently located to the right or to the left of the middorsal cuticular specialization, not following any particular pattern, near the anterior margin of the segment (Figs 2B, 3D, 4E); paired setae in lateroventral position (Figs 2A–B, D, 3E); females furthermore with paired, sexually dimorphic setae in ventrolateral position (Figs 2A, 3E). Sexually dimorphic male tubes in ventromedial position (Fig. 2D); detailed morphology of these tubes not determined. Paired sensory spots in paradorsal, laterodorsal and ventrolateral position, the latter near the ventrolateral-ventromedial limit, not longitudinally aligned with the following ventromedial sensory spots (Figs 2A–B, D, 3D–E).

| Segment | PD | SD | LD | PL | VL | VM | PV |
|---------|----|----|----|----|----|----|----|
| 1       | 3× (♀), 2× (♂) | 2× (♀), 1× (♂) | 1× (♀), 2× (♂) | 3× |
| 2       | 1× | 3× (♀), 1× (♂) | 2× (♀), 3× (♂) | 1× |
| 3       | 1× | 3× (♀), 2× (♂) | 2× | 1× | 1× | 2× (♀), 3× (♂) | 1× |
| 4       | 1× | 3× (♀), 1× (♂) | 2× | 1× | 3× |
| 5       | 1× | 3× (♀), 2× (♂) | 2× | 1× | 3× | 3× (♀), 3× (♂) | 1× |
| 6       | 1× | 2× | 1× | 1× | 2× (♀), 3× (♂) | 1× |
| 7       | 1× | 2× | 1× | 2× (♀), 3× (♂) | 1× |
| 8       | 1× | 3× (♀), 2× (♂) | 1× | 1× | 2× (♀), 3× (♂) | 1× |
| 9       | 1× | 2× | 1× | 3× |
| 10      | 1× | 3× (♀), 1× (♂) | 3× (♀), 1× (♂) | 3× (♀), 1× (♂) | 1× (♀) |
| 11      |    |    |    |    |    |    |    |

Segment 2 with middorsal elevation not projecting beyond the posterior margin of segment (Figs 2B, 3D, 4E). Unpaired seta in paradorsal position, on this and following segments indifferently located to the right or to the left of the middorsal cuticular specialization, not following any particular pattern, near the anterior margin of the segment (Figs 2B, 3D, 4E); paired setae in lateroventral position (Figs 2A–B, D, 3E); females furthermore with paired, sexually dimorphic setae in ventrolateral position (Figs 2A, 3E). Sexually dimorphic male tubes in ventromedial position (Fig. 2D); detailed morphology of these tubes not determined. Paired sensory spots in paradorsal, laterodorsal and ventrolateral position, the latter near the ventrolateral-ventromedial limit, not longitudinally aligned with the following ventromedial sensory spots (Figs 2A–B, D, 3D–E).

**Table 3.** Summary of nature and arrangement of intracuticular structures (maybe outlets of glandular cells) in *Setaphyes elenae* sp. nov., including sexually dimorphic differences. Abbreviations: LD = laterodorsal; PD = paradorsal; PL = paralateral; PV = paraventral; SD = subdorsal; VL = ventrolateral; VM = ventromedial.

Fig. 3 (opposite page). Light micrographs showing trunk overviews and details of cuticular trunk characters of ♀, holotype (NHMD 655358) (A–L, N) and ♂, paratype (NHMD 655361) (M) of *Setaphyes elenae* sp. nov. A. Dorsal overview. B. Dorsal view on right half of segment 1. C. Ventral view on left half of segment 1. D. Dorsal view on right half of segment 2. E. Ventral view on left half of segment 2. F. Dorsal view on right half of segment 3. G. Ventral view on left half of segment 3. H. Ventral overview. I. Dorsal view on right half of segment 4. J. Ventral view on left half of segment 4. K. Dorsal view on right half of segment 8. L. Ventral view on left half of segment 8. M. Dorsal view on right half of segment 9. N. Ventral view on left half of segment 9. Abbreviations: ldse = laterodorsal seta; lts = lateral terminal spine; lvse = lateroventral seta; mde = middorsal elevation; mdp = middorsal process; pdse = paradorsal seta; vlse = ventrolateral seta; vmse = ventromedial seta. Numbers after abreviations indicate correspong segment; sensory spots are marked as dashed circles, and glandular cell outlets as continuous circles. Scale bars: A, H = 100 µm; B–G, I–N = 20 µm.
Fig. 4. Scanning electron micrographs showing general overview and details of the cuticular trunk morphology of a non-type specimen of *Setaphyes elenae* sp. nov. **A.** Dorsal overview. **B.** Middorsal elevation of segment 4. **C.** Cuticular ornamentation of anterior margin of segment 1. **D.** Detail of primary and secondary pectinate fringes of segment 5. **E.** Middorsal to paradorsal view of segment 2. **F.** Laterodorsal seta of segment 5. **G.** Middorsal process of segment 1. **H.** Ventral view of segments 4–5. **I.** Dorsal view of segment 10. **J.** Subdorsal sensory spots of segment 8. Abbreviations: mde = middorsal elevation; mdp = middorsal process; pdse = paradorsal seta; ppf = primary pectinate fringe; s = segment; vmse = ventromedial seta. Numbers after abbreviations indicate corresponding segment; sensory spots are marked as dashed circles. Scale bars: A = 100 µm; B, D, F–G, J = 1 µm; C, E, H–I = 10 µm.
Segment 3 with middorsal elevation as on the preceding segment (Figs 2B, 3F). Unpaired seta in paradorsal position (Figs 2B, 3F); paired setae in laterodorsal, lateroventral and ventromedial positions (Figs 2A–B, 3F–G). Paired sensory spots in paradorsal, subdorsal, laterodorsal and ventromedial positions, the latter mesially shifted compared to the previous ones, aligned with those of the following segments (Figs 2A–B, 3F–G).

Segment 4 with middorsal elevation as on the preceding segment (Figs 2B, 3I, 4B). Unpaired seta in paradorsal position (Figs 2B, 3I); paired setae in lateroventral position (Figs 2A–B, 3J). Paired sensory spots in paradorsal, subdorsal, laterodorsal and ventromedial positions (Figs 2A–B, 3I–J).

Segment 5 similar to segment 3 in the arrangement of the cuticular elevation, setae and sensory spots (Figs 2A–B, 4F, H).

Segment 6 similar to segment 4 in the arrangement of the cuticular elevation, setae and sensory spots (Fig. 2A–B).

Segment 7 similar to segments 3 and 5 in the arrangement of the cuticular elevation, setae and sensory spots (Fig. 2A–B).

Segment 8 with middorsal elevation as on the preceding segment (Figs 2B, 3K). Unpaired seta in paradorsal position (Figs 2B, 3K); paired setae in lateroventral position (Fig. 2A–B). Two pairs of sensory spots in subdorsal position, and one pair in paradorsal, laterodorsal and ventromedial positions (Figs 2A–B, 3K–L, 4J).

Segment 9 with middorsal process as that of segment 1 (Figs 2B, 3M, 4I). Unpaired seta in paradorsal (Figs 2B and 3M); paired setae in laterodorsal and lateroventral positions (Fig. 2A–B). Two pairs of sensory spots in subdorsal position, and one pair in paradorsal, laterodorsal and ventromedial positions (Figs 2A–B, 3M–N, 4I). Nephridiopores in lateroventral position.

Segment 10 without middorsal cuticular specialization. Paired setae in lateroventral position (Figs 2A–C, 4I). Two pairs of sensory spots in subdorsal position, and one pair in laterodorsal and ventrolateral positions, the latter near the tergosternal junction (Figs 2A–C, 4I).

Segment 11 without cuticular appendages, partly retractable into segment 10 (Fig. 4A, I). Tergal plate triangular, with a concave and distally pointed posterior margin; sternal plates form a pair of ventral extensions rounded distally (Figs 2A–C, 4I). Males with two sexually dimorphic pairs of stout, thick penile spines (Fig. 2C). Lateral terminal spines’ length differs from males (relatively longer, LTS average ratio = 158.93 μm) to females (relatively shorter, LTS average ratio = 100.9 μm).

**Discussion**

*Setaphyes elenae* sp. nov. clearly agrees with the main diagnostic characters of the genus, including the absence of ventrolateral setae on segment 5, the presence of unpaired paradorsal setae on segments 2–9 and paired lateroventral setae on segments 2–10, ball-and-socket joints only conspicuous at segments 2–3 (reduced on posterior segments), and the presence of scattered, dot-shaped, intracuticular structures (sometimes called as cuticular scars) at both tergal and sternal plates (Sánchez et al. 2016). However, it can easily be distinguished from the remaining species of the genus by the unique arrangement of setae and middorsal cuticular specializations.

Regarding the seta arrangements, *Setaphyes elenae* sp. nov. has paired laterodorsal setae on segments 3, 5, 7 and 9, a pattern similar to that of *S. dentatus* and *S. flaveolatus* (Reinhard 1881; Zelinka 1928; Sánchez et al. 2016), with the exception that these congeners lack the setae on segment 9. The remaining
Fig. 5. Boxplots showing the ranges of different body measurements of *Setaphyes elenae* sp. nov., *S. dentatus* (Reinhard, 1881) and *S. flaveolatus* (Zelinka, 1928). A. Total trunk length. B. Standard sternal width. C. Lateral terminal spines’ length.
species of the genus also possess laterodorsal setae also even segments. In addition, *S. elenae* sp. nov. is characterized by bearing shortened, distally rounded middorsal processes on segments 1 and 9 exclusively, while both *S. dentatus* and *S. flaveolatus* have middorsal processes on segments 1 and 7–9. The pattern of ventromedial setae also differs between *S. dentatus* (on segments 4–5 and 7–9, females furthermore on segment 3), *S. flaveolatus* (on segments 5 and 7–8, females furthermore on segment 3) and *S. elenae* sp. nov. (on segments 3, 5 and 7).

*Setaphyes cimarensis* Sánchez *et al.*, 2018, *S. dentatus*, *S. elenae* sp. nov. and *S. flaveolatus* are also characterized by having patterns of cuticular ornamentation that can be used to discriminate congeners (Sánchez *et al.* 2018). Longitudinal, parallel, fold-like cuticular thickenings are present on segment 10 in *S. dentatus* and *S. cimarensis*, which are absent in *S. elenae* sp. nov. and *S. flaveolatus*. Moreover, *S. flaveolatus* has a continuous, reticule-like ornamentation only present in the middle region of the tergal plate, whereas *S. dentatus* and *S. elenae* sp. nov. possess a similar ornamentation extended throughout the anterior margin of the plate, and *S. cimarensis* is characterized by having small, rounded, isolated depressions near the anterior margin of the plate with a net-like structure (Sánchez *et al.* 2018).

The only statistically significant differences were found in the total trunk length (TL) between *Setaphyes elenae* sp. nov. and *S. flaveolatus* (*p* = 0.00042), so we can conclude that these two morphologically similar species may be also distinguished by the total trunk length (Fig. 5A). Oppositely, *S. dentatus* and *S. elenae* sp. nov. overlap, and statistically significant differences were not found (*p* = 0.51153) (Fig. 5A). *Setaphyes flaveolatus* is the smallest of the tested species, with a total trunk length of 500–600 μm, followed by *S. dentatus* and *S. elenae* sp. nov. with a total trunk length of 600–800 μm (Fig. 5A). The remaining analysed morphometric measurements were not significantly different between *S. elenae* sp. nov. and the aforementioned congeners (Fig. 5B–C).

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**References**

Altenburger A. 2016. The neuromuscular system of *Pycnophyes kielensis* (Kinorhyncha: Allomalo-rhagida) investigated by confocal laser scanning microscopy. *EvoDevo* 7: 25. https://doi.org/10.1186/s13227-016-0062-6

Cepeda D., Sánchez N. & Pardos F. 2019. First extensive account of the phylum Kinorhyncha from Haiti and the Dominican Republic (Caribbean Sea), with the description of four new species. *Marine Biodiversity* 49: 2281–2309. https://doi.org/10.1007/s12526-019-00963-x

Curini-Galletti M., Artois T., Delogu V., De Smet W.H., Fontaneto D., Jondelius U., Leasi F., Martínez A., Meyer-Wachsmuth I., Nilsson K.S., Tongiorgi P., Worsaae K. & Todaro, M.A. 2012. Patterns of diversity in soft-bodied meiofauna: dispersal ability and body size matter. *PLoS ONE* 7: e33801. https://doi.org/10.1371/journal.pone.0033801

Fox J., Weisberg S., Price B., Adler D., Bates D., Baud-Bovy G., Bolker B., Ellison S., Firth D., Friendly M., Gorjane G., Graves S., Heiberger R., Krivitsky P., Labossiere R., Maechler M., Monette G., Murdoch D., Nilsson H., Ogle D., Ripley B., Venables W., Walker S., Winsemius D., Zeileis A. & R Core Team. 2019. Package ‘car’. Ver. 3.0-5. https://r-forge.r-project.org/projects/car/
Gerlach S.A. 1969. *Cateria submersa* sp. n., ein cryptorhager Kinorhynch aus dem sublitoralen Mesopsammal der Nordsee. *Veröffentlichungen des Instituts für Meeresforschung in Bremerhaven* 12: 161–168.

Higgins R.P. 1988. Kinorhyncha. In: Higgins R.P. & Thiel H. (eds) *Introduction to the Study of Meiofauna*: 328–331. Smithsonian Institution Press, Washington D.C.

Huys R. & Coomans A. 1989. *Echinoderes higginsi* sp.n. (Kinorhyncha, Cyclorhagida) from the southern North Sea with a key to the genus *Echinoderes* Claparède. *Zoologica Scripta*: 18: 211–221. https://doi.org/10.1111/j.1463-6409.1989.tb00446.x

McIntyre A.D. 1962. The class Kinorhyncha (Echinoderida) in British waters. *Journal of the Marine Biological Association of the United Kingdom* 42: 503–509. https://doi.org/10.1017/S0025315400054217

McIntyre, A.D. 1964. Meiobenthos of sub-littoral muds. *Journal of the Marine Biological Association of the United Kingdom* 44 (3): 665–674. https://doi.org/10.1017/S0025315400027843

Neuhaus B. 2013. Kinorhyncha (= Echinodera). In: Schmidt-Rhaesa A. (ed.) *Handbook of Zoology. Gastrotricha, Cycloneuralia and Gnathifera*. Volume 1: *Nematomorpha, Priapulida, Kinorhyncha, Loricifera*: 1–69. De Gruyter, Göttingen. https://doi.org/10.1515/9783110272536

Sánchez N., Herranz M., Benito J. & Pardos F. 2012. Kinorhyncha from the Iberian Peninsula: new data from the first intensive sampling campaigns. *Zootaxa* 3402: 24–44. https://doi.org/10.11646/zootaxa.3402.1.2

Sørensen M.V. & Pardos F. 2008. Kinorhynch systematics and biology – an introduction to the study of kinorhynchs, inclusive identification keys to the genera. *Meiofauna Marina* 16: 21–73.
Sørensen M.V., Heiner I. & Hansen J.G. 2009. A comparative morphological study of the kinorhynch genera *Antygomonas* and *Semnoderes* (Kinorhyncha, Cyclorhagida). *Helgoland Marine Research* 63: 129–147. [https://doi.org/10.1007/s10152-008-0132-9](https://doi.org/10.1007/s10152-008-0132-9)

Sørensen M.V., Rho H.S., Min W.G., Kim D. & Chang S.Y. 2013. Occurrence of the newly described kinorhynch genus *Meristoderes* (Cyclorhagida: Echinoderidae) in Korea, with the description of four new species. *Helgoland Marine Research* 67 (2): 291–319. [https://doi.org/10.1007/s10152-012-0323-2](https://doi.org/10.1007/s10152-012-0323-2)

Weering T.C.E., Berger G.W. & Kalf J. 1987. Recent sediment accumulation in the Skagerrak, Northeastern North Sea. *Netherlands Journal of Sea Research* 21: 177–189. [https://doi.org/10.1016/0077-7579(87)90011-1](https://doi.org/10.1016/0077-7579(87)90011-1)

Weering T.C.E., Rumohr J. & Liebezeit G. 1993. Holocene sedimentation in the Skagerrak: a review. *Marine Geology* 111: 379–391. [https://doi.org/10.1016/0025-3227(93)90142-I](https://doi.org/10.1016/0025-3227(93)90142-I)

Zelinka C. 1928. *Monographie der Echinodera*. Engelmann, Leipzig.

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