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Three novel *Rubripirellula* species isolated from plastic particles submerged in the Baltic Sea and the estuary of the river Warnow in northern Germany

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Abstract Planctomycetes are a unique and important phylum containing mostly aquatic bacteria, which are often associated with phototrophic surfaces. A complex lifestyle, their potential for the production of bioactive small molecules, their unusual cell biology and a large number of giant and hypothetical genes in their genomes make these microorganisms a fascinating topic for further research. Here, we characterise three novel planctomycetal strains isolated from polystyrene and polyethylene particles that were submerged in the German part of the Baltic Sea and the estuary of the river Warnow. All three strains showed typical planctomycetal traits such as division by polar budding and formation of rosettes. The isolated strains were mesophilic and neutrophilic chemoheterotrophs and reached generation times of 10–25 h during laboratory-scale cultivation. Taxonomically, the three strains belong to the genus *Rubripirellula*. Based on our analyses all three strains represent novel species, for which we propose the

names *Rubripirellula amarantea* sp. nov., *Rubripirellula tenax* sp. nov. and *Rubripirellula reticaptiva* sp. nov. The here characterised strains Pla22^T (DSM 102267^T = LMG 29691^T), Poly51^T (DSM 103356^T = VKM B-3438^T) and Poly59^T (DSM 103767^T = LMG 29696^T) are the respective type strains of these novel species. We also emend the description of the genus *Rubripirellula*.

Keywords Marine bacteria · Planctomycetes · Baltic Sea · Biofilm · Brackish water · *Rubripirellula amarantea* · *Rubripirellula reticaptiva* · *Rubripirellula tenax*

Introduction

Planctomycetes are ubiquitous bacteria often associated with nutrient-rich aquatic surfaces. By occupying these ecological niches, Planctomycetes largely contribute to the vital activity of global carbon and nitrogen cycles (Wiegand et al. 2018). Phylogenetically, Planctomycetes form the PVC superphylum alongside with the phyla of Verrucomicrobia, Lentisphaerae, Kiritimatiellaeota, Candidatus Omnitrophica and Chlamydiae (Spring et al. 2016; Wagner and Horn 2006). Based on exceptional traits, such as, presence of compartment-like structures (Lindsay et al. 1997) and lack of peptidoglycan (König et al. 1984), Planctomycetes were thought to be beyond the

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bacterial cell plan (Devos et al. 2013; Fuerst and Sagulenko 2011). It was even speculated that they might represent the missing link between bacteria and eukaryotes (Devos and Reynaud 2010). In the recent years, however, presence of peptidoglycan was confirmed (Jeske et al. 2015; van Teeseling et al. 2015) and compartment-like structures were re-interpreted as invaginations of the cytoplasmic membrane (Acehan et al. 2013; Boedeker et al. 2017; Lage et al. 2013; Santarella-Mellwig et al. 2013). Planctomycetes were thus classified as Gram-negative bacteria (Devos 2014), but still remain exceptional and enigmatic in comparison to well-characterised bacteria of other phyla. Most Planctomycetes divide unusually by budding, while some also perform binary fission or even a combination of both modes of division (Wiegand et al. 2019). The observed lack of canonical divisome proteins including the otherwise universal FtsZ (Jogler et al. 2012; Pilhofer et al. 2008) might also be part of the explanation why Planctomycetes are resistant to many antibiotics (Cayrou et al. 2010; Godinho et al. 2019).

As a result of the absence of nutrients in oligotrophic seawater, several Planctomycetes attach to nutrient-rich aquatic surfaces (Bengtsson et al. 2012; Bondoso et al. 2014, 2015, 2017; Lage and Bondoso 2014; Vollmers et al. 2017). Attachment typically goes along with a lifestyle switch between planktonic swimmer cells and sessile stalked mother cells (Jogler et al. 2011). Once attached to the surface, they start to degrade polymeric compounds for biomass production, e.g. utilising complex sugars released by algae. This strategy was demonstrated for the model polysaccharide dextran (Jeske et al. 2013; Lachnit et al. 2013). Unique sugar-binding pili attached to crateriform structures and an extremely enlarged periplasm are believed to be involved in the uptake of such high molecular weight sugars (Boedeker et al. 2017), which requires further attention in the next years.

Planctomycetes were found to be predominant members in biofilms on the mentioned nutrient-rich aquatic surfaces (Bengtsson and Øvreås 2010; Kohn et al. 2019), which is counter-intuitive when considering their slow growth compared to other natural competitors, e.g. members of the *Roseobacter* clade (Frank et al. 2014; Wiegand et al. 2018). It was proposed that this observation is the result of defense strategies involving secondary metabolite production, which, in presence of a competitor, ultimately lead to

release of antimicrobial compounds. This notion, that Planctomycetes are ‘talented’ producers of such small bioactive molecules, is substantiated by large genomes (Kohn et al. 2016) as well as several predicted gene clusters involved in small molecule production (Graca et al. 2016; Jeske et al. 2016; Wiegand et al. 2019).

Succinctly, Planctomycetes are amongst the most maverick of all bacteria (Wiegand et al. 2018), which motivated us to expand the collection of Planctomycetes by isolating and characterising 79 novel planctomycetal strains presented in an overview article (Wiegand et al. 2019). Here, we validly describe three novel strains, Pla22^T, Poly51^T, and Poly59^T, isolated from plastic particles submerged in the Baltic Sea and the river Warnow in northern Germany. The genus *Rubripirellula*, to which the three here proposed species belong, was described earlier with *Rubripirellula obstinata* as the type species and the only member of the genus so far. The type strain *R. obstinata* LFI^T was isolated from algae (*Laminaria* sp.) at the coast of Porto, Portugal (Bondoso et al. 2015).

Materials and methods

Preparation of cultivation medium

For strain isolation and subsequent cultivations M1H NAG ASW medium [M1 medium with HEPES as buffering agent (M1H) additionally supplemented with *N*-acetyl glucosamine (NAG) as carbon and nitrogen source and artificial seawater (ASW)] was used. This medium is originally based on a recipe described earlier (Staley et al. 1992). This recipe was extensively modified to allow cultivation of a broad range of different planctomycetal strains and was prepared as described before (Kallscheuer et al. 2019).

Isolation of the strains

Strain Pla22^T was sampled on the 4th of September 2014 from polyethylene (PE) pellets (ExxonMobil HDPE HTA 108, diameter 3 mm), which were incubated in 2 m depth for 14 days in the Warnow river north of Rostock, Germany. The sampling site (54.106 N, 12.096 E) is located close to a wastewater treatment plant discharge. The exact setup of sampling was described earlier (Oberbeckmann et al. 2018).

Strain Poly51^T and strain Poly59^T were sampled on the 8th of October 2015 at Heiligendamm, Germany (54.146 N, 11.843 E) from polystyrene (PS) and PE particles, respectively, stored in separate incubators in the water for 14 days in 2 m depth and accessed via Heiligendamm pier. The strains were isolated from the plastic particles as described before (Wiegand et al. 2019). Plastic was chosen as material for sampling as it is cheap and easily available, provides a high surface area, is non-toxic and inert, i.e. it neither reacts with the seawater nor is degraded. Briefly, for the isolation of Pla22^T (September 2014) samples were stored for 8 weeks at 4 °C before isolation of bacterial biofilms by digestion with 2 mg/mL (28 U/mL) β -galactosidase for 30 min at 30 °C and simultaneous vortexing every 5 min followed by 10 min sonication at 30 °C. Separation of plastic from the biofilm was performed by filtration. Biofilms were stored for 11 months at 4 °C prior to plating. For isolation of Poly51^T and Poly59^T (October 2015) incubated plastic particles were washed three times with sterile natural seawater and stored at 4 °C until cultivation (5 days after sampling). Plastic particles were vortexed and 50 μ L of the liquid seawater was used for cultivation on M1H NAG ASW plates containing 8 g/L gellan gum. For selection of Planctomycetes 500 mg/L streptomycin and 200 mg/L ampicillin were added as antibiotic reagents, while 20 mg/L cycloheximide was used to prevent fungal growth. In order to check whether the isolated strains indeed represent Planctomycetes the 16S rRNA gene was amplified by PCR and sequenced as previously described (Rast et al. 2017).

Light microscopy

Phase contrast (Phaco) analyses were performed employing a Nikon Eclipse Ti inverted microscope with a Nikon DS-Ri2 camera (blue LED). Specimens were immobilised in MatTek glass bottom dishes (35 mm, No. 1.5) employing a 1% agarose cushion (Will et al. 2018). Images were analysed using the Nikon NIS-Elements software (version 4.3). To determine the cell size, at least 100 representative cells were counted manually (Annotations and Measurements, NIS-Elements) or by using the NIS-Elements semi-automated Object Count tool (smooth: 4 $\times$, clean: 4 $\times$, fill holes: on, separate: 4 $\times$).

Electron microscopy

For field emission scanning electron microscopy bacteria were fixed in 1% (v/v) formaldehyde in HEPES buffer (3 mM HEPES, 0.3 mM CaCl₂, 0.3 mM MgCl₂, 2.7 mM sucrose, pH 6.9) for 1 h on ice and washed once employing the same buffer (Rast et al. 2017). Cover slips with a diameter of 12 mm were coated with a poly-L-lysine solution (Sigma-Aldrich) for 10 min, washed in distilled water and air-dried. 50 μ L of the fixed bacteria solution was placed on a cover slip and allowed to settle for 10 min. Cover slips were then fixed in 1% glutaraldehyde in TE buffer (20 mM TRIS, 1 mM EDTA, pH 6.9) for 5 min at room temperature and subsequently washed twice with TE buffer before dehydrating in a graded series of acetone (10, 30, 50, 70, 90, 100%) on ice for 10 min at each concentration. Samples from the 100% acetone step were brought to room temperature before placing them in fresh 100% acetone. Samples were then subjected to critical-point drying with liquid CO₂ (CPD 300, Leica). Dried samples were covered with a gold/palladium (80/20) film by sputter coating (SCD 500, Bal-Tec) before examination in a field emission scanning electron microscope (Zeiss Merlin) using the Everhart–Thornley HESE2 detector and the inlens SE detector in a 25:75 ratio at an acceleration voltage of 5 kV.

Genome information

Genome information of the three isolated strains is available under accession numbers SJPI00000000 (Pla22^T), SJPW00000000 (Poly51^T) and SJPX00000000 (Poly59^T). The corresponding 16S rRNA gene sequences can be found under accession numbers MK554581 (Pla22^T), MK554552 (Poly51^T) and MK554553 (Poly59^T).

Physiological analysis

For determination of the pH optimum 100 mM HEPES was used for cultivations at pH 7.0, 7.5 and 8.0. For cultivation at pH 5.0 and 6.0 HEPES was replaced by 100 mM 2-(*N*-morpholino)ethanesulfonic acid (MES), whereas 100 mM *N*-cyclohexyl-2-aminoethane-sulfonic acid (CHES) served as a buffering agent at pH 9.0 and 10.0. Cultivations for determination of the pH optimum were performed at 28 °C. For

determination of the temperature optimum the strains were cultivated at temperatures ranging from 10 to 40 °C in M1H NAG ASW medium at pH 7.5.

Phylogenetic analysis

16S rRNA gene phylogeny was computed for strains Pla22^T, Poly51^T and Poly59^T, the type strains of all described planctomycetal species (available in May 2019) and all isolates recently published (Wiegand et al. 2019). The 16S rRNA gene sequences were aligned with SINA (Pruesse et al. 2012). The phylogenetic analysis was done with RAxML (Stamatakis 2014) employing a maximum likelihood (ML) approach with 1000 bootstraps, the nucleotide substitution model GTR, gamma distributed rate variation and estimation of proportion of invariable sites (GTRGAMMAI option). Three 16S rRNA genes of bacterial strains from the PVC superphylum were used as outgroup. Average nucleotide identities (ANI) were calculated using OrthoANI (Lee et al. 2016) and average amino acid identities (AAI) were calculated using the aai.rb script of the enveomics collection (Rodriguez-R and Konstantinidis 2016). Values of percentage of conserved proteins (POCP) were calculated as described before (Qin et al. 2014). The *rpoB* nucleotide sequences were taken from the above-mentioned as well as other publicly available genome annotations and the sequence identities were determined as described (Bondoso et al. 2013). Upon extracting only those parts of the sequence that would have been sequenced with the described primer set the alignment and matrix calculation was done with Clustal Omega (Sievers et al. 2011).

Results and discussion

Phylogenetic analysis

The phylogenetic positions of strains Poly59^T, Poly51^T and Pla22^T were determined by 16S rRNA gene analysis as shown in Fig. 1. The three strains cluster monophyletically with *R. obstinata* LF1^T (Bondoso et al. 2015). An assessment of different phylogenetic markers describing the relationship between the type strain and the novel isolates can be found in Table 1. The 16S rRNA gene similarities of Poly59^T, Poly51^T and Pla22^T to *R. obstinata* LF1^T are

all below the species threshold of 98.7% (Stackebrandt and Ebers 2006), but above the proposed genus threshold of 94.5% (Yarza et al. 2014). The comparison of *rpoB* sequence fragments as described (Bondoso et al. 2013) gave similarity results between 80.9 and 82.9%, which are values below the proposed species threshold, but above the given genus threshold (Bondoso et al. 2013). This implies that none of the novel strains belong to the species *Rubripirellula obstinata*, but that all are members of the genus *Rubripirellula*. The ANI values of under 95% support this result of separate species (Kim et al. 2014). Additionally, the two-way AAI in the range of 60–80% (Luo et al. 2014) and a POCP of > 50% (Qin et al. 2014) also indicate that all strains belong to the same genus. The two closely related strains Poly51^T and Poly59^T have a 16S rRNA gene sequence identity of 98.4%, an *rpoB* sequence identity of 85.8%, and an ANI of 75.4% implying that they also form two separate species.

Morphological and physiological analysis

Cell morphologies and cell sizes of strains Pla22^T, Poly51^T and Poly59^T were analysed by light microscopy (Fig. 2) and scanning electron microscopy (Fig. 3) during exponential growth. The obtained images were compared to the already described strain *R. obstinata* LF1^T. All four strains form mainly loose aggregates and divide by polar budding. Buds have the same shape as the mother cell. Flagella formation was only observed for LF1^T. All four strains contain fimbriae at the budding pole but lack a stalk. A holdfast structure was only observed in strain LF1^T. Only Poly51^T and LF1^T formed a visible capsule. Detailed information on morphology, locomotion and mechanism of cell division is summarised in Table 2.

Cells of strain Pla22^T are round grain rice-shaped with an average size of $1.7 \pm 0.3 \mu\text{m}$ in length and $0.9 \pm 0.2 \mu\text{m}$ in width (Fig. 2a, g). Colonies have an amaranth pink color. The strain is aerobic and grew at pH values ranging from 6.0 to 8.5 with an optimum at pH 7.5 (Fig. 4). Growth was observed at temperatures from 10 to 36 °C with optimal growth at 33 °C. In M1H NAG ASW medium a maximal growth rate of 0.068 h^{-1} was observed, which corresponds to a doubling time of approximately 10 h (Fig. 4).

Strain Poly51^T has a similar shape and color as strain Pla22^T, but the cells are slightly smaller and

Fig. 1 16S rRNA gene-based phylogeny. The phylogenetic tree highlights the position of the three here investigated strains in relation to their closest described relatives. 16S rRNA gene phylogeny was computed using the maximum likelihood

method. Bootstrap values after 1000 re-samplings are given at the nodes (in %). The outgroup consists of three 16S rRNA genes from the PVC superphylum

Table 1 Phylogenetic markers

Strain	16S rRNA sequence similarity (%)	<i>rpoB</i> sequence similarity (%)	Average nucleotide identity (ANI) (%)	Average amino acid identity (AAI) (%)	Percentage of conserved proteins (POCP) (%)
Pla22 ^T	96.60	80.90	71.80	66.80	54.70
Poly51 ^T	95.90	82.90	71.60	64.30	52.30
Poly59 ^T	95.30	82.90	71.10	64.10	51.30

Comparison of the different phylogenetic markers 16S rRNA, *rpoB*, ANI, two-way AAI and POCP between the novel *Rubripirellula* strains and *Rubripirellula obstinata* LF1^T

appear chubbier (Poly51^T: $1.4 \pm 0.2 \times 0.9 \pm 0.1 \mu\text{m}$) (Figs. 2c, h, 3c). Strain Poly51^T grows at a pH range of 6.5–9.0 with an optimum at pH 8.0. The

temperature optimum of the strain is between 22 and 24 °C, while cell growth was observed at temperatures ranging from 10 to 28 °C (Fig. 4). The strain did not

Fig. 2 Light microscopy images and cell size plots of the three isolated strains. The mode of cell division (**a**, **c**, **e**) and a general overview of cell morphology (**b**, **d**, **f**) is shown in the pictures.

The scale bar is 1 μm . For determination of the cell size (**g**, **h**, **i**) at least 100 representative cells were counted manually or by using a semi-automated object count tool

grow at temperatures of 30 °C or higher. The highest observed growth rate is 0.028 h^{-1} corresponding to a generation time of 25 h.

Cells of Poly59^T appear round grain rice-shaped to round (Figs. 2e, 3e, f) with an average cell size of $1.5 \pm 0.3 \mu\text{m}$ (length) $\times$ $1.0 \pm 0.2 \mu\text{m}$ (width), are pink-colored and show the same behavior as Poly51^T with regard to pH (growth range 6.5–9.0, optimum 7.5–8.0) (Fig. 4). Comparable to Poly51^T, strain Poly59^T grows best at 24 °C, but in contrast could also grow at 30 and 33 °C. Its highest grow rate was calculated to 0.043 h^{-1} , corresponding to a doubling time of approx. 16 h. The type species *R. obstinata* LF1^T has a temperature optimum at 25 °C (range 10–30 °C) and grows in a pH range of 7.5–10.5 (Bondoso et al. 2015). Thus, the temperature optimum of LF1^T is very similar to Poly51^T and Poly59^T, but different from Pla22^T (33° C). Remarkably, the temperature optimum of Pla22^T being 8 °C higher than that of the three other strains is the most striking difference and is also reflected by a higher growth rate of Pla22^T at the optimal temperature. While Poly51^T, Poly59^T and LF1^T were isolated from the Baltic Sea or the Atlantic Ocean, Pla22^T was isolated from the

estuary of the river Warnow close to a wastewater treatment plant. The water temperature profile in the sampling areas might be considerably different and allows at least to speculate that the higher temperature optimum of Pla22^T might be related to the sampling location. With regard to pH, LF1^T, growing at a range of 7.5–10.5, is slightly more alkaliphilic than the other three strains, which grow at a pH range of 6.5–9.0. None of the strains grew at pH 5.5 or lower and the observed differences between the strains were less pronounced compared to the parameter temperature.

Genomic characteristics

All three strains show similarities in genome size (6.9–8.0 Mb) and G + C content (53.7–56.2%). The genome size (7.1 Mb) and G + C content (54.3%) of *R. obstinata* LF1^T also fit in the ranges obtained for the novel species. The number of putative open reading frames (ORFs) ranges from 5182 to 6274 (of which 2096–2640 are annotated as hypothetical proteins), which corresponds to 746–785 putative protein-coding genes per Mb. These values give a nearly identical coding density of $89.0 \pm 0.3\%$. Genomic

Fig. 3 Scanning electron microscopic pictures of the three novel strains. The scale bar is 1 μm

characteristics are summarised in Table 2. The observed genome sizes of the three strains are not only comparable to *R. obstinata* LF1^T, but also to other closely related species, e.g. *Rhodopirellula baltica* (7.1 Mb) and *Roseimaritima ulvae* (8.1 Mb). The genome of Pla22^T (6.9 Mb) is around 1 Mb smaller than those of Poly51^T and Poly59^T (both 7.9 Mb). *R.*

obstinata LF1^T contains two copies of the 16S rRNA gene, while the gene is present in single copy in the three other strains. The number of 67 tRNAs in LF1^T is similar to Pla22^T (74 tRNAs) and Poly59^T (71 tRNAs), but much lower compared to Poly51^T (120 tRNAs).

Table 2 Phenotypic and genotypic features of the three characterized *Rubripirellula* species in comparison to *R. obstinata* LF1^T (Bondoso et al. 2015)

Characteristics	Pla22 ^T	Poly51 ^T	Poly59 ^T	<i>R. obstinata</i> LF1 ^T
<i>Phenotypic features</i>				
Size (length × width)	1.7 × 0.9 μm	1.4 × 0.9 μm	1.5 × 1.0 μm	1.5–2.0 × 1.3–1.7 μm
Shape	Round grain rice-shaped	Round grain rice-shaped	Round grain rice-shaped to round	Ovoid to pear-shaped
Colony color	Amaranth pink	Amaranth pink	Pink	Red
Aggregates	Yes	Yes	Yes	Yes
Division	Budding	Budding	Budding	Budding
Flagella	n.o.	n.o.	n.o.	Yes
Crateriform structures	Polar	Polar	Polar	Polar
Fimbriae	Matrix or fiber, at budding pole	Matrix or fiber	Fibercap at budding pole	At budding pole
Capsule	n.o.	Yes	n.o.	yes
Bud shape	Like mother cell	Like mother cell	Like mother cell	Like mother cell
Budding pole	Polar	Polar	Polar	Polar
Stalk	n.o.	n.o.	n.o.	n.o.
Holdfast structure	n.o.	n.o.	n.o.	yes
<i>Genotypic features</i>				
Genome size (bp)	6,945,823	7,988,747	7,852,560	7,094,218
Plasmids (bp)	n.o.	n.o.	n.o.	n/a
G+C (%)	53.7 ± 0.9	56.2 ± 2.1	54.8 ± 1.7	54.3 ± 1.7
Completeness (%)	96.91	98.28	98.28	98.28
Contamination (%)	1.72	1.72	1.72	1.72
Protein-coding genes	5182	6274	6088	5870
Hypothetical proteins	2096	2794	2640	2888
Protein-coding genes/ Mb	746	785	775	827
Coding density (%)	89.3	88.8	89.3	84.4
16S rRNA genes	1	1	1	2
tRNA genes	74	120	71	67

n.o. not observed, n/a not available

Conclusion

Based on our physiological, genomic and phylogenetic analysis the three strains represent each a novel species of the genus *Rubripirellula*, for which we present the here characterised strains as respective type strains.

Emended description of the genus *Rubripirellula*
Bondoso et al. 2016

The description of the genus *Rubripirellula* is as given previously (Bondoso et al. 2015), with the following modification: Colonies are amaranth pink- to red-coloured.

Fig. 4 Temperature and pH optima of the isolated strains. In the upper panel, the given data points show the average growth rates obtained after cultivation of the three isolated strains in M1H NAG ASW medium in biological triplicates at different

temperatures and a constant pH of 7.5. In the bottom panel, the data points show the average growth rates for cultivation at different pH values and a constant temperature of 28 °C

Description of *Rubripirellula amarantea* sp. nov

Rubripirellula amarantea (a.ma.ran'te.a. N.L. fem. adj. *amarantea* of amaranth; corresponding to the amaranth colour of the cells).

Cells are round grain rice-shaped (length: $1.7 \pm 0.3 \mu\text{m}$, width: $0.9 \pm 0.2 \mu\text{m}$), form aggregates and divide by polar budding. Colonies have amaranth pink color. Cells of the type strain grow at ranges of 10–36 °C (optimum 33 °C) and at pH 6.0–8.5 (optimum 7.5). The genome of the type strain has a size of 6,945,823 bp and a G + C content of $53.7 \pm 0.9\%$. The type strain genome (acc. no. SJPI00000000) and 16S rRNA gene sequence (acc. no. MK554581) are available from GenBank.

The type strain is Pla22^T (DSM 102267^T = LMG 29691^T), isolated from polyethylene particles incubated in brackish water of the Warnow river estuary

close to a wastewater treatment plant near Rostock, Germany.

Description of *Rubripirellula tenax* sp. nov

Rubripirellula tenax (te'nax. L. fem. adj. *tenax* holding fast, clinging; corresponding to the characteristic of the cells to be adhesive).

Cells are round grain-rice shaped (length: $1.4 \pm 0.2 \mu\text{m}$, width: $0.9 \pm 0.1 \mu\text{m}$), form aggregates and divide by polar budding. Pink amaranth coloured colonies are formed. The temperature optimum is 22–24 °C (growth observed from 10 to 28 °C). The type strain failed to grow at 30 °C or higher. The preferred pH is 8.0, but growth is also observed at pH 6.5–9.0. The type strain genome has a size of 7,988,747 and a G + C content of $56.2 \pm 2.1\%$. The type strain genome (acc. no. SJPW00000000) and 16S

rRNA gene sequence (acc. no. MK554552) are available from the GenBank database.

The type strain is Poly51^T (DSM 103356^T = VKM B-3438^T), isolated from polystyrene particles incubated in the Baltic Sea in 2 m depth near Heiligendamm, Germany.

Description of *Rubripirellula reticaptiva* sp. nov

Rubripirellula reticaptiva (re.ti.cap.ti'va. L. n. *rete* a fishing net; L. fem. adj. *captiva* captive; N.L. fem. adj. *reticaptiva* corresponding to the capture of the cells in water).

Cells are round rice grain-shaped to round ($1.5 \pm 0.3 \mu\text{m} \times 1.0 \pm 0.2 \mu\text{m}$), form aggregates and divide by polar budding. Colonies have a red colour. The preferred temperature and pH is 24 °C and 7.5–8.0, respectively, while growth is observed in the range of 10–33 °C and at pH 6.5–9.0. The type strain genome has a size of 7,852,560 and a G + C content of $54.8 \pm 1.7\%$. The type strain genome (acc. no. SJPX00000000) and 16S rRNA gene sequence (acc. no. MK554553) are available from GenBank.

The type strain is Poly59^T (DSM 103767^T = LMG 29696^T), isolated from polyethylene particles incubated in the Baltic Sea in 2 m depth near Heiligendamm, Germany.

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Author contributions NK wrote the manuscript, analyzed the data and prepared the figures; SW and MJ performed the genomic and phylogenetic analysis; AH isolated the strains and performed the initial cultivation and strain deposition; SHP and CB performed the light microscopic analysis; MSMJ contributed to text preparation and revised the manuscript; MR performed the electron microscopic analysis; CJ and MJ took the samples in the Baltic Sea and the river Warnow and CJ supervised the study. All authors read and approved the final version of the manuscript

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

Ethical statement This article does not contain any studies with animals performed by any of the authors.

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