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A New Species of the γ-Proteobacterium *Francisella*, *F. adeliensis* Sp. Nov., Endocytobiont in an Antarctic Marine Ciliate and Potential Evolutionary Forerunner of Pathogenic Species

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Abstract

The study of the draft genome of an Antarctic marine ciliate, *Euplotes petzi*, revealed foreign sequences of bacterial origin belonging to the γ-proteobacterium *Francisella* that includes pathogenic and environmental species. TEM and FISH analyses confirmed the presence of a *Francisella* endocytobiont in *E. petzi*. This endocytobiont was isolated and found to be a new species, named *F. adeliensis* sp. nov.. *F. adeliensis* grows well at wide ranges of temperature, salinity, and carbon dioxide concentrations implying that it may colonize new organisms living in deeply diversified habitats. The *F. adeliensis* genome includes the *igl* and *pdp* gene sets (*pdpC* and *pdpE* excepted) of the *Francisella* pathogenicity island needed for intracellular growth. Consistently with an *F. adeliensis* ancient symbiotic lifestyle, it also contains a single insertion-sequence element. Instead, it lacks genes for the biosynthesis of essential amino acids such as cysteine, lysine, methionine, and tyrosine. In a genome-based phylogenetic tree, *F. adeliensis* forms a new early branching clade, basal to the evolution of pathogenic species. The correlations of this clade with the other clades raise doubts about a genuine free-living nature of the environmental *Francisella* species isolated from natural and man-made environments, and suggest to look at *F. adeliensis* as a pioneer in the *Francisella* colonization of eukaryotic organisms.

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Introduction

Like their multicellular descendants, also single-celled eukaryotes host a huge variety of bacteria. Ciliates in particular are a preferential and stable home to bacteria, which may be carried either attached as epibionts to the cell body surface, as is the case of the association between a group (designated as "epixenosomes") of Verrucomicrobia and *Euplotidium itoi* [1], or enclosed as endocytobionts inside the cell body. Being a principal component of the diet of ciliates, which are mostly phagotrophic and filter-feeding, bacteria can easily escape digestion and adopt a new intracellular lifestyle [2]. Roughly 250 ciliate species, among the nearly 10,000 that are in total known, have been detected to be hosts of endocytobiont bacteria. Large-size species of *Paramecium*, *Euplotes*, and *Spirostomum* may be home also of mixed populations of unrelated species of bacteria [3, 4].

The knowledge of the biology and life cycle of endocytobiont bacteria in ciliates is essentially limited to

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species of *Holospora* and *Caedibacter*, which are colonizers of the nuclear apparatus of freshwater species of *Paramecium* [5, 6]. These symbionts have been successfully isolated from host-cell homogenates, but any attempt of cultivation outside their hosts has failed as in the case of any other bacterial symbiont of aerobic ciliates [7].

A substantial contribution to improve this knowledge may now be provided by the isolation and cultivation of *Francisella* bacteria living as endocytobionts in marine species of *Euplotes*, a genus which is quite rich also in freshwater species extensively studied for their symbiotic associations with polymorphic populations of *Polynucleobacter* [8]. *F. endociliophora*, earlier described as a novel subspecies of *F. noatunensis* [9], is the first *Francisella* that has been isolated and genome-sequenced from a marine species of *Euplotes*, *E. raikovi*, dwelling in temperate waters [10]. Here, we report the isolation and genome sequencing of another new species of *Francisella*, *F. adeliensis* sp. nov., living as endocytobiont in a bipolar (Antarctic and Arctic) species of *Euplotes*, *E. petzi*.

The genus *Francisella* comprises species classified as facultative intracellular γ-proteobacteria potentially noxious to their hosts [11, 12]. *F. tularensis*, with its three subspecies, is a specialized intracellular pathogen of both invertebrate and vertebrate hosts, human beings included [13, 14]. *F. noatunensis*, with its two subspecies adapted to different hosts' temperatures, is the etiological agent of the fish disease known as francisellosis [15, 16]. The endosymbiotic *F. persica* (ex *Wolbachia persica*) [17], together with the generalists *F. philomiragia* and *F. novicida*, may harm human beings with a compromised immune system [18–20].

The position that *F. adeliensis* takes in the genome-based phylogenetic tree provides new insights on *Francisella* diversity and helps to decipher the emergence of symbiosis and the evolution of pathogenicity in this genus.

Materials and Methods

Host *E. petzi* Cells

The *E. petzi* cells were isolated from a sample of seawater and sandy bottom collected by means of a sediment trap from Adelie Cove in Antarctica, at a depth of 27 m, a temperature of − 1.2 °C and a salinity of 34‰. Cultures were maintained in the laboratory in cold rooms, at 4 °C, under a cycle of 12 h of very low light and 12 h of dark, as previously described [21]. The green alga *Dunaliella tertiolecta* was used as food source.

Fluorescent In Situ Hybridization (FISH)

E. petzi cells were collected from severely starved cultures, transferred onto glass slides, fixed with 4% formaldehyde in phosphate-buffered saline (PBS) for 10 min at room temperature, and permeabilized by ethanol gradient (50%, 80%, and 100% of ethanol in water, for 10 min each). The fluorescein-labeled probe EUB338 (5′-GCTGCCTCCCGTAGGAT-3′) for eubacteria and the Cy3-labeled probe Bwall1448 (5′-CAACCATTCGCCGGGCCT-3′) for *Francisella* were synthesized by Integrated DNA Technologies (Coralville, IA, USA). Hybridization was performed following the method described by Hugenholtz et al. [22]. Briefly, 2 μl of each probe solution (50 ng/μl) in 20 μl hybridization buffer (0.9 M NaCl, 20 mM Tris-HCl pH 7.0, 15% formamide, 0.1% SDS) were added directly to the cells on slides. Hybridization was performed in a humid chamber at 46 °C for 3 h. Slides were then washed 20 min with washing buffer (318 mM NaCl, 20 mM Tris-HCl pH 7.0, 0.1% SDS) at 48 °C and air dried. Slides were embedded with anti-fading mounting medium and then inspected with a Nikon confocal microscope (Nikon, Amsterdam, The Netherlands).

Light and Transmission Electron Microscopy (TEM)

In light microscopy, bacteria were observed in vivo, as well as after staining with the fluorescent dye Syto-9 (used as recommended by the manufacturer Molecular Probes, Eugene, OR, USA), on an Axiophot microscope equipped with an Axio Cam MRC color video camera and an AxioVision computer-assisted image analysis system (Zeiss, Jena, Germany).

For TEM analysis, bacteria were fixed with 2.5% glutaraldehyde and 6% sucrose in 0.1 M cacodylate buffer, pH 7.2, for 2 h at 4 °C. After three washings at 4 °C in the same buffer, samples were post-fixed with 1% osmium tetroxide in 0.1 M cacodylate buffer, pH 7.2, for 1 h at 4 °C, washed in the same buffer, and dehydrated in a gradient ethanol series. Samples were then infiltrated with mixtures of LRWhite resin/ethanol in different percentages, embedded in pure LRWhite resin, and left to polymerize for 2 days at 50 °C. Resin blocks were cut with a Reichert Ultracut ultramicrotome using a diamond knife. Ultrathin sections (60–80 nm) were collected on copper grids, stained with uranyl acetate and lead citrate, and observed with a JEOL 1200EXII electron microscope (JEOL, Peabody, MA, USA). Micrographs were captured using a SIS VELETA CCD camera (Olympus, Muenster, Germany) equipped with iTEM software.

Isolation, Identification, and Culturing

The isolation of *F. adeliensis* was carried out from *E. petzi* cell lysates following the procedure previously used by Sjödin et al. to isolate *F. endociliophora* [10], taking care to incubate plates at 4 °C. Briefly, *E. petzi* cell samples were bead beaten and acid treated according to Humrighouse et al. [23], before being diluted in PBS and spread on Cysteine Heart Agar

Blood (CHAB) culture plates, supplemented with 10^5 U/l penicillin and 40 mg/l vancomycin as described [24]. The culture plates were incubated at 4 °C for 1 to 2 weeks and monitored for bacterial growth. Colonies were then isolated and maintained in CHAB plates at 4 °C. To identify *F. adeliensis* from other contaminating bacteria, isolated colonies were picked, resuspended in 20 μl of water and immediately lysed by boiling for 3 min. Five microliters of each lysed cell suspension were used as template in PCR, run using two sets of primers: fw1 (5′-GCGTTTACCACGGAGTGATT -3′) and rv1 (5′-TGGAGCCTAGCGGGATC-3′); fw2 (5′-AGTCAGGGAGGAAGTTTATTTGGTT-3′) and rv2 (5′-CACCTTCCTCCGCCTTGT-3′). Positive clones were maintained in CHAB plates for subsequent analysis.

For dot-plate analysis, a *Francisella* colony was suspended in 1 ml PBS buffer and 5 microliters of serial dilutions were spotted on CHAB plates and incubated at 4, 10, 20, 30, and 37 °C. Plates were checked for bacterial growth every 3 days. For growth assays in liquid medium, an overnight culture was used to inoculate aliquots of 50 ml of T medium [25] to reach an OD_{600} ranging from 0.01 to 0.05. Flasks were then incubated at different temperatures, salinity, and CO_2 concentrations. Bacterial growth was monitored by measuring the OD_{600} every day during the first week, then every 3 days. The number of generation/day was calculated using the Origin 8 software.

The presence of the enzymes catalase and oxidase was tested on agar plates using a 3% H_2O_2 solution and an oxidase-strip (OXOID, Monza, Italy), respectively. Motility was determined with the hanging drop technique.

Genome Sequencing and Assembly

Isolated DNA was sequenced using the Nextera XT library protocol on an Illumina MiSeq instrument in addition to a Pacific Biosciences RSII system (10-kb library, 2-h movie length), generating a total of 57,926 PacBio reads with an average read length of 11,653 bp, using single-molecular real-time (SMRT) cells. The initial draft of the genome was generated by assembling PacBio reads using the SMRT analysis system version 2.3.0. Polishing of the draft genome was performed using Illumina reads [26].

Phylogenetic Analysis

The phylogenetic analysis was inferred using the neighbor-joining method [27]. The evolutionary distances among *Francisella* genomes were computed using the number of differences method [28] and are in the units of the number of base differences per sequence. The analysis involved 139 nucleotide sequences, with a total of 213,734 positions in the final dataset. Positions containing gaps and missing data were eliminated. Evolutionary analyses were conducted in MEGA7

[29]. *Fangia hongkongensis* was included as outgroup to generate the genome-based phylogenetic tree.

Results

Identification

Total DNA preparations of *E. petzi* subjected to high-throughput sequencing generated 24,800 assembled contigs (Villalobo and Vallesi, unpublished), of which approximately 800 (equivalent to a total of 1.6 Mb) revealed a close similarity to bacterial sequences available from public databases, with the highest matching value of each contig systematically resulting against gene sequences of *Francisella* species.

Among the 800 contigs, one of 5091 bp included the 16S and 23S rRNA gene sequences plus the sequences of the tRNA^{Ile} and tRNA^{Ala} genes (Fig. 1a). Therefore, it revealed to be a typical bacterial rDNA operon. Using the SILVA INcremental Aligner bioinformatics tool [30], the 16S rRNA gene sequence of this operon was classified as belonging to *Francisella* with 94.39% identity and 97 score along 1480 bp. Given that the 3% cutoff rule [31] for a 16S divergence among species was fulfilled, the new 16S rRNA gene sequence was assumed to belong to a new *Francisella* species for which the proposed name is *Francisella adeliensis* nov. sp.. The species name is after that of the Antarctic cove, Adelie, from which *E. petzi*, the *F. adeliensis* host, was collected.

Analyzed in the BLASTN 2.6.1 database [32] for its closest identity, the *F. adeliensis* 16S rRNA gene sequence showed the best alignment (only seven nucleotide variations along 1376 bp) with the 16S gene sequence of an unnamed and uncultured γ-proteobacterium reported to be a chemoautotrophic symbiont on gills of deep-sea clams and mussels collected at a 10-m depth from the fjord of Saanich Inlet, British Columbia [33]. The other two closest counterparts were the 16S sequences of *F. endociliophora* [10] and *F. salina* [24], with 96% of sequence identity along the 1481-bp gene length.

Intracellular Localization

To verify whether *F. adeliensis* resides as endosymbiont inside *E. petzi*, or it coexists as environmental bacteria with *E. petzi* in culture, *E. petzi* cells were starved for 10 days to avoid any possible bacterial contamination from undigested food, and analyzed by fluorescent in situ hybridization (FISH) with two distinct probes: the first ("EUB338," see "Materials and Methods") specific to a 16S rRNA-sequence conserved in most bacterial species, and the second ("Bwall1448") specific to a 23S rRNA region unique to *Francisella* [34]. Both probes generated fluorescent signals within the cytoplasm of *E. petzi* cells (Fig. 1b), and their co-localization provided evidence that *F. adeliensis* was the only guest.

Fig. 1 *F. adeliensis* identification. **a** Schematic representation of the *F. adeliensis* rDNA operon. The relative positions of fluorescent FISH probes and primers used in colony-PCR are indicated. The 33-bp sequence exclusive of *F. adeliensis* rDNA operon is shown. **b** Fluorescent in situ hybridization of *E. petzi* cells: a, signal from fluorescein-labeled probe EUB338 for all eubacteria; b, signal from Cy3-labeled probe Bwall1448 specific for *Francisella*; c, co-localization of signals of the two labeled probes. Scale bar = 20 μm
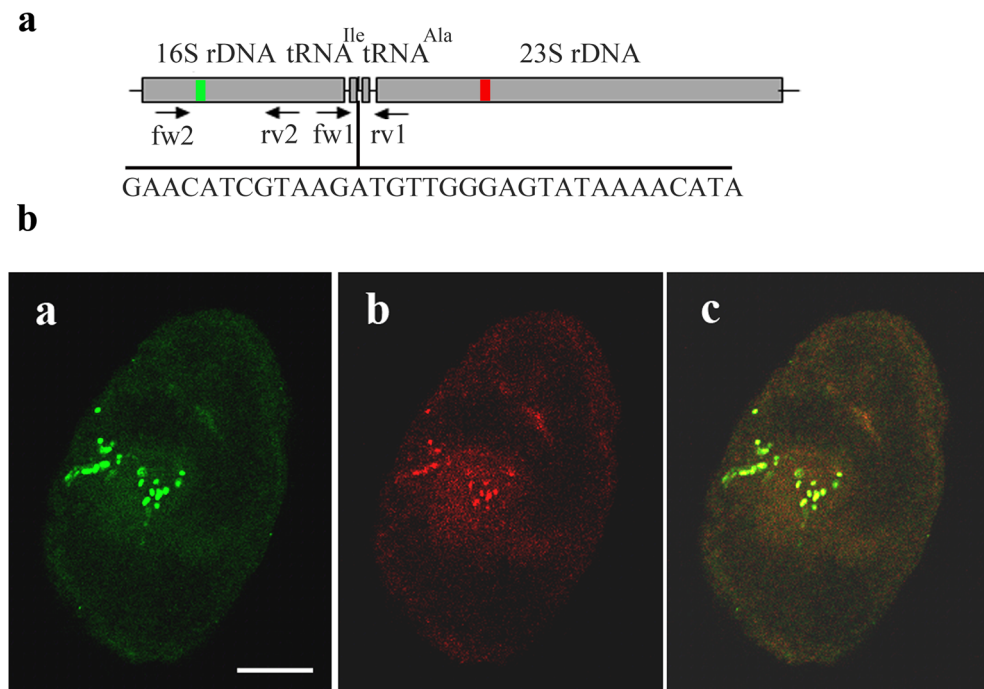
TEM analysis of *E. petzi* cells, suspended without food for not less than 1 week before being used, confirmed the presence of numerous bacteria (Fig. 2a, b). Only occasionally were bacteria observed to be individually dispersed in the cytoplasm, each confined inside a membranous-bound vesicle (Fig. 2e), or apparently free in the cytosol (Fig. 2f, g). Much more often, they appeared clustered together in larger fusogenic membrane-bound structures (Fig. 2c, d). These structures were quite heterogeneous in size and number of enclosed bacteria, and usually located in close proximity of the host's somatic and transcriptionally active nucleus (macronucleus).

Phenotypic Traits

Individual *F. adeliensis* colonies were screened by PCR using two sets of specific primers (Fig. 1a). Primers ("fw1" and "rv1," see "Materials and Methods") of one set were designed to amplify a 360-bp fragment containing a 33-bp sequence lying between the two tRNA coding regions and without counterparts in the rDNA operons of other *Francisella* species. Primers ("fw2" and "rv2") of the second set were designed to amplify a 660-bp fragment of the 16S rRNA coding region shared among other *Francisella* species. Products sequenced from both amplifications showed to fully match the genomic data, confirming the taxonomic identity of the isolated colonies with *F. adeliensis*.

On CHAB plates, *F. adeliensis* colonies look spherical, white and slightly mucoidal, formed by roundish (diameter, 1–1.2 μm) bacteria (Fig. 3a), which are non-motile, catalase-

positive, and oxidase-negative. In solid medium, colonies are visible after 3 days of incubation at temperatures ranging from 20 to 30 °C, and require 6–12 days to grow when incubated at 4 and 10 °C (Fig. 3b). In liquid medium, the highest growth rate was measured at 20 and 30 °C and the lowest at 4 °C (Fig. 3c). The mean numbers of generations/day were counted to be 0.11, 0.29, 0.53, and 0.47 at 4, 10, 20, and 30 °C, respectively, and no growth was observed at 37 °C. Roughly one half of bacteria inoculated on plates at 37 °C died after 16 h of incubation and none survived after 48 h.

In the presence of 5% CO_2, *F. adeliensis* cultures grew with OD_{600} values approximately 60% lower than those measured in ambient atmosphere (0.04%). Instead, no significant variation in the growth rate was observed in cultures left to grow in liquid medium containing salt concentrations ranging from 0 to 35‰, implying that *F. adeliensis* is a strongly euryhaline bacterium (data not shown).

Genomic Features

The *F. adeliensis* genome extends for 2,054,094 bp, a length matching the mean genome size of other *Francisella* species (1.96 ± 0.14 Mbp, Table 1) much more closely than the size of any other bacterial genome (3.82 ± 1.8 Mbp) [35]. It contains 1880 predicted protein coding sequences, 38 tRNA genes, ten rRNA genes (four 5S rRNA, three 16S rRNA, and three 23S rRNA) and one tmRNA gene (Table 1). Its average nucleotide identity (ANI) with the closest *Francisella* genomes is in the range of 77–78.8% (Table 2), which is distant from the 95–96% range usually taken as the minimum threshold value to consider two genome sequences as belonging to the same
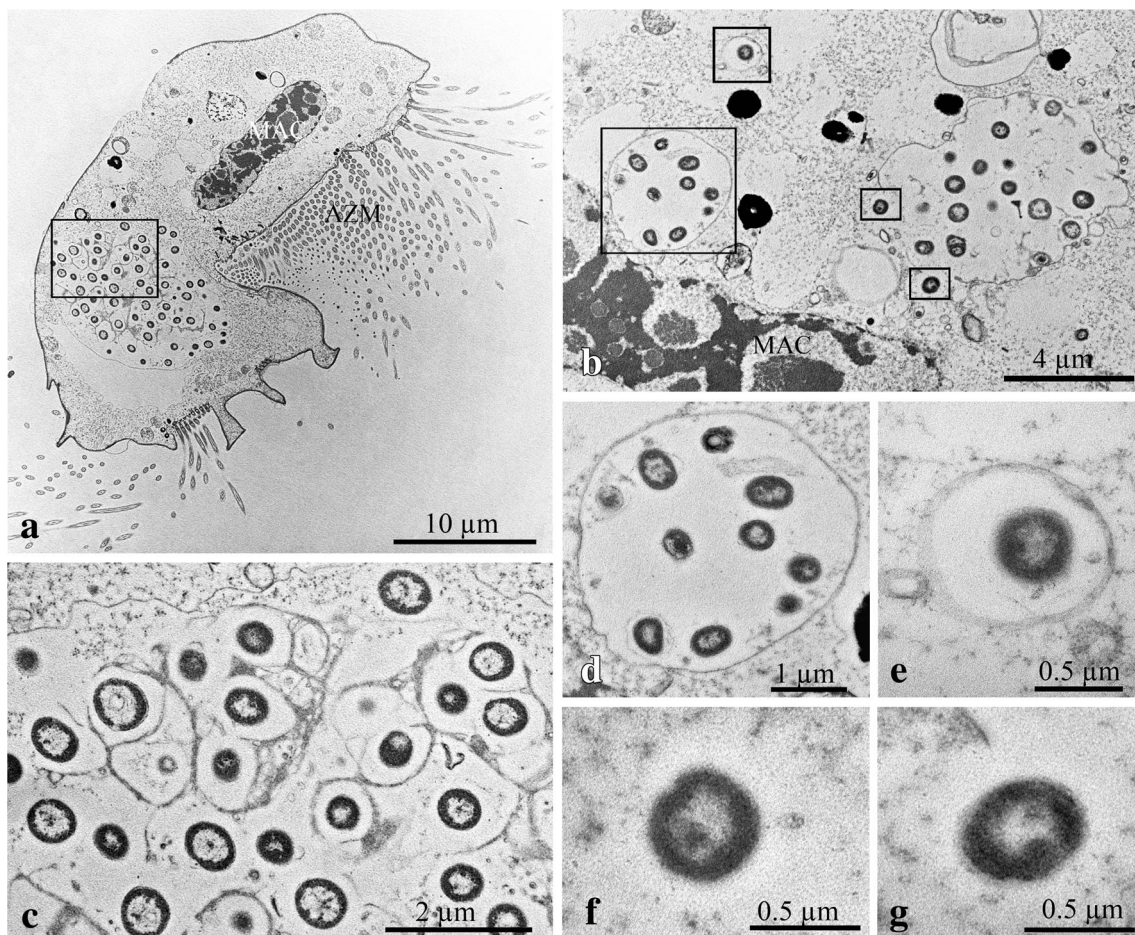
Fig. 2 TEM of *E. petzi* cells containing *F. adeliensis*. **a**, **b** Micrographs showing bacteria individually dispersed in the host cytoplasm, or associated together in groups enclosed in membrane-bound compartments. **c–g** Magnifications of the boxed areas in **a** and **b**. MAC macronucleus, AZM adoral zone ciliary membranelles

species [36]. Consistently with an intracellular lifestyle, the average 32.6% G+C content of the *F. adeliensis* genome closely reflects the $32.38 \pm 0.24\%$ content of the other endosymbiotic *Francisella*, and is significantly lower than the average G+C content ($49.1 \pm 12.4\%$) shown by free-living bacteria [35].

Based on a search for transposable elements and phages carried out with PHASTER and ISFinder softwares [37, 38], the *F. adeliensis* genome contains prophage sequences like other *Francisella*. However, it includes only one IS*Ftu4* insertion sequence element (E-value $1e^{-15}$).

Analysis of the *F. adeliensis* genome for the presence of the ten *igl* (intracellular growth locus) and five *pdp* (pathogenicity determinant proteins) genes, which composethe so-called *Francisella* pathogenicity island responsible for the virulence of *F. tularensiss* [39, 40], indicated that all *igl* genes are present, while the *pdp* gene set lacks the *pdpC* and *pdpE* genes.

The observation that *F. adeliensis* requires complex media to grow in culture suggested a loss of genes responsible for the synthesis of essential amino acids. This hypothesis was verified by screening the *F. adeliensis* genome for the presence of genes responsible for the synthesis of arginine, cysteine, histidine, lysine, methionine and tyrosine for which the pathogenic *F. tularensis* is known to be auxotrophic [41]. Only the histidine and arginine biosynthesis appeared to be genetically supported: the histidine biosynthesis by the complete set of relevant genes, and the arginine biosynthesis by the *argJ* gene that likely replaces the lack of *argA*, *argD*, and *argE* genes [42]. Instead, the biosynthesis of the other four amino acids appeared genetically not supported. The *F. adeliensis* genome lacks the genes *dapD*, *dapC*, and *dapE* encoding enzymes responsible for the lysine biosynthesis [43], as well as the gene encoding cystathionine γ-synthase responsible for the methionine and cysteine biosynthesis [44]. With regard to the tyrosine biosynthesis, the genome contains the complete gene set for the shikimate pathway, but it lacks the gene encoding prephenate dehydrogenase which converts prephenic acid to 4-hydroxyphenyl-pyruvic acid [45]. In conclusion, *F. adeliensis* shows to be prototrophic for arginine and histidine, and auxotrophic for cysteine, lysine, methionine, and tyrosine.
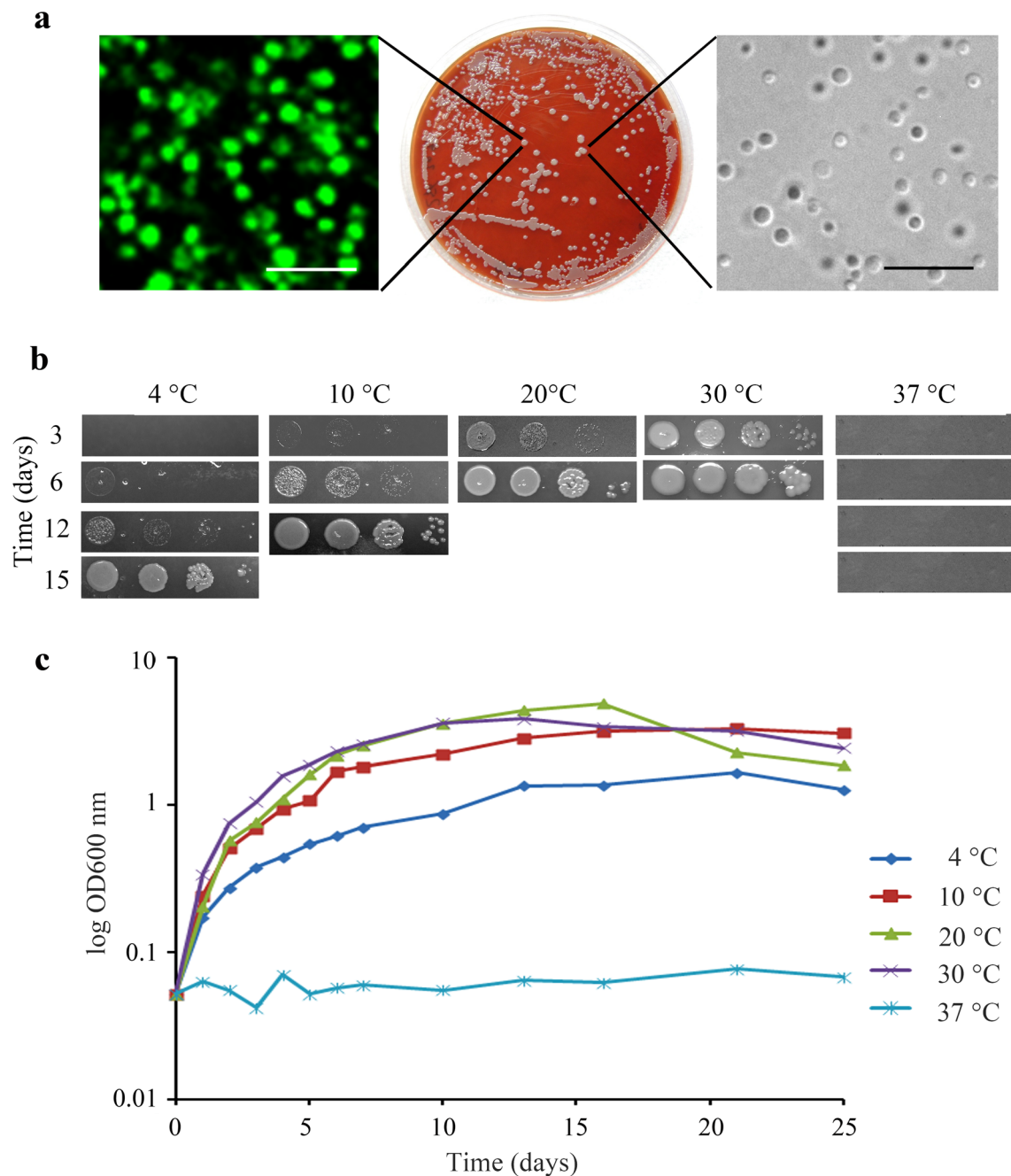
Fig. 3 *F. adeliensis* colonies and growth. **a** Colonies grown on a CHAB plate and bacteria as they appear after staining with Syto-9 fluorescent dye (left picture), and in vivo in differential interference contrast microscopy (right picture). Scale bar, 10 μm. **b** Dot-plate analysis. Serial dilutions of a *F. adeliensis* suspension were spotted on plates and plates incubated at the indicated temperatures for the indicated times (days). **c** Growth curves in liquid medium incubated at the indicated temperatures. Data from a representative experiment are shown; experiments were repeated three times with equivalent results

Phylogenetic Relationships

To assess the *F. adeliensis* interspecific relationships, the *F. adeliensis* genome was compared with the other *Francisella* genomes available from NCBI using 139 gene sequences for a total of 213,734 nucleotide positions. As shown in Fig. 4, *F. adeliensis* forms its own clade with a high statistical support. Together with the clade formed by

F. frigiditurris, a species recently isolated from the water of a cooling tower [46], it precedes the split of four other major clades in which all the other *Francisella* species are subdivided in full accord with the recently proposed genome-based *Francisella* phylogeny [46, 47]. One of the four clades is specific to species, such as *F. tularensis* and *F. novicida*, that are pathogenic to terrestrial hosts, and *F. persica* (formerly *Wolbachia persica*) isolated from ticks

Table 1 Genome comparison of ten different *Francisella* species. The accession numbers of the examined genomes are indicated

Species	Origin	Genome size (bp)	Pred. proteins	tRNAs	rRNA (16S+23S)	% G+C
F. adeliensis CP021781	*E. petzi*	2,054,094	1880	38	10 3 + 3	32.6
F. endociliophora NZ_CP009574	*E. raikovi*	2,015,987	1891	38	10 3 + 3	32.4
Allofrancisella guangzhouensis NZ_CP010427.1	Cooling tower water	1,658,482	1423	38	10 3 + 3	32.0
F. noatunensis subsp. orientalis FNO24 NZ_CP011922.1	Nile tilapia	1,862,322	1449	39	10 3 + 3	32.3
F. philomiragia GA01-2794 NZ_CP009440.1	Human	2,148,038	1999	40	10 3 + 3	32.4
F. hispaniensis 3523 NC_017449	Human	1,945,310	1798	38	10 3 + 3	32.3
F. tularensis subsp. holarctica LVS NC_007880	Vaccine strain	1,895,994	1766	38	10 3 + 3	32.2
F. tularensis subsp. tularensis WY96 NZ_CP012037.1	Human	2,005,074	1871	38	10 3 + 3	32.4
F. tularensis subsp. mediasiatica FSC147 NC_010677	Gerbil	1,893,886	1659	38	10 3 + 3	32.3
F. salina TX077308 NC_015696	Seawater	2,035,931	1884	39	10 3 + 3	32.9

[12, 17]. The second one includes species such as *F. noatunensis* that are pathogenic to fish, as well as *F. salina* isolated from a seawater sample [46]. The third one includes species such as *F. endociliophora* and *F. halioticida* isolated from marine hosts, together with *F. uliginis* isolated from a seawater sample [9, 12, 46]. And the fourth one is specific to *Francisella* species that have been isolated from waters of cooling systems, and are usually described as "environmental" species and regarded as belonging to the genus *Allofrancisella* [48, 49].

Discussion

The isolation reported here of *F. adeliensis* from an Antarctic strain of *E. petzi* follows the isolation of *F. endociliophora* from *E. raikovi* [10], which is a species distributed in the Caspian and Mediterranean Seas and Eastern Atlantic Ocean [50], and the identification of DNA sequences of a taxonomically undetermined *Francisella* in the genome of *E. focardii* [51], which is a species endemic to Antarctic coastal waters [52]. Altogether, these findings strongly suggest that *Francisella*/*Euplotes* associations are relatively common in

the marine environment, and two additional considerations reinforce this hypothesis. The first consideration is related to the bipolar biogeographic distribution that characterizes the species structure of the *F. adeliensis*'s host, *E. petzi* [21, 53]. Embracing Arctic and White Sea populations in addition to Antarctic and peri-Antarctic ones, this distribution clearly implies that the *F. adelinesis* association with *E. petzi* is likely not restrained to the Antarctic waters where it has been detected. Being extended to the high latitudes of both the hemispheres, it appears to be virtually global and the analysis of other bipolar *Euplotes* species for their symbiotic associations with *F. adeliensis* and/or its close relatives may definitively establish these global dimensions. The second and more significant consideration is related to the psychrophilic and euryhaline behavior shown by *F. adeliensis*. Growing well at temperatures ranging from 4 to 30 °C and promptly adapting to 0–35 ‰ variations in the ambient salinity, *F. adeliensis* appears capable of colonizing other organisms independently of their adaptation to live in marine, brackish, or lacustrine habitats of either cold or temperate areas.

The 16S rRNA gene sequences are the molecules of choice for phylogenetic reconstructions, but their use in devising a *Francisella* phylogenetic tree has frequently been biased by

Table 2 ANI in percent between known *Francisella* genomes

	F. endociliophora NZ_CP009574.1	*A. guangzhouensis* NZ_CP010427.1	*F.noatunensis subsp. orientalis* FNO24 NZ_CP011922.1	*F. philomiragia GA01-2794* NZ_CP009440.1	*F. tularensis subsp. tularensis* WY96 NZ_CP012037.1
F. adeliensis CP021781	78.84	77.07	77.58	77.89	77.69
F. endociliophora NZ_CP009574.1	–	78.4	80.44	81.51	80.22
A. guangzhouensis NZ_CP010427.1	–	–	79.30	78.5	78.99
F. noatunensis subsp. orientalis FNO24 NZ_CP011922.1	–	–	–	95.15	82.09
F. philomiragia GA01-2794 NZ_CP009440.1	–	–	–	–	88.239

Fig. 4 Evolutionary relationships of *F. adeliensis*. The optimal tree with the sum of branch length = 284,909.68814135 is shown. The tree is drawn to scale, with branch lengths in the same units as those of the evolutionary distances used to infer the phylogenetic tree. The scale bar corresponds to 5000 nucleotide differences. Growth style and environment of each species are indicated by colored dots on the right; the six major branches of the tree are enclosed in colored rectangles. The position of *F. adeliensis* is highlighted in bold
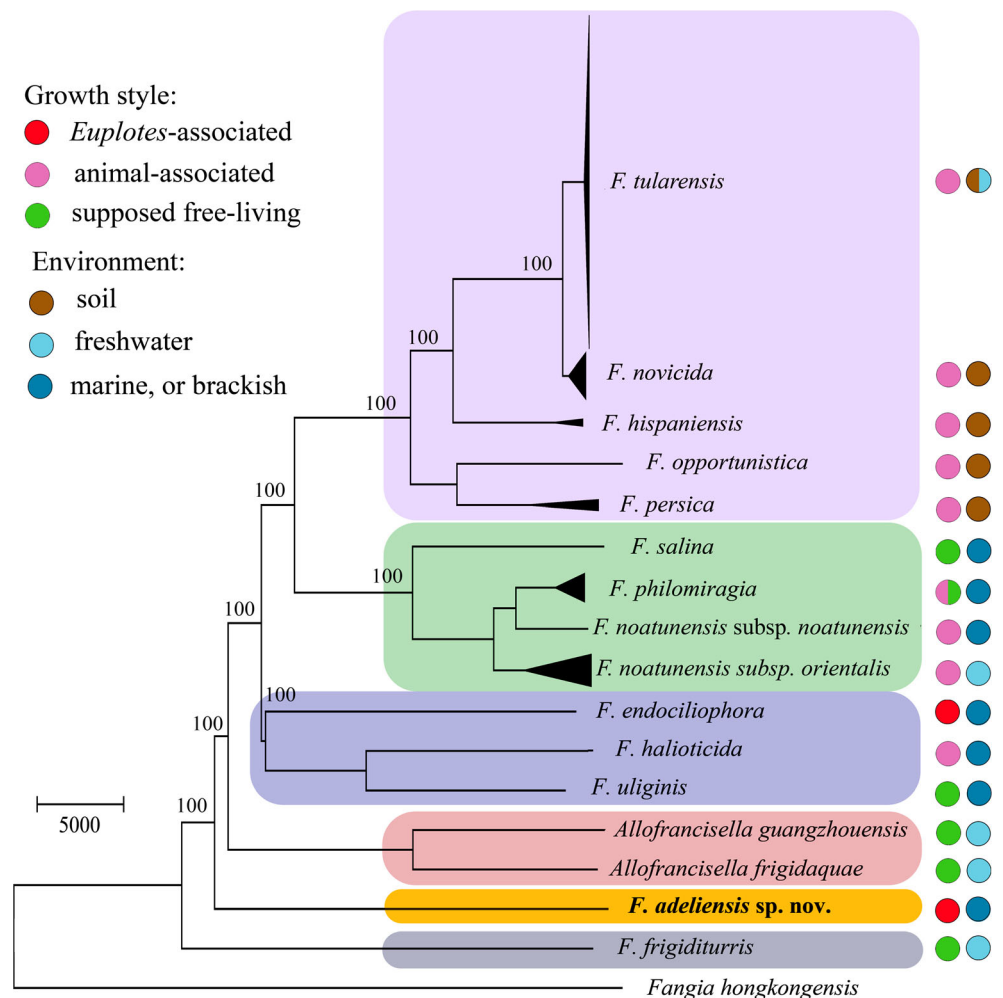
branches supported by low bootstrap values due to the particularly high degree of conservation that these sequences show in *Francisella*. Only the recent availability of genomic data provided more solid grounds to trace the phylogenetic relationships among *Francisella* species, producing phylogenetic trees with more solid statistic support [46, 54]. In the genome-based tree updated with the inclusion of *F. adeliensis* (shown above in Fig. 4), *F. adeliensis* branches surprisingly distant from all the intracellular *Francisella*, including *F. endociliophora* endocytobiont in *E. raikovi*. It correlates much closer to the two earliest branching clades that are uniquely formed by *Francisella* species, namely *F. frigiditurris*, *Allofrancisella frigidaquae*, and *A. guangzhouensis*, isolated from cooling towers. As such, they are collectively regarded as environmental species.

Granted that these species are really free living—considering the strong acidic conditions used for their isolation, it cannot be excluded that they have actually been isolated from some eukaryotic microorganisms living inside the cooling towers—this correlation implies that *F. adeliensis* foreruns the *Francisella* adaptive evolution in replacing a free-living

lifestyle with an intracellular/endosymbiotic style. And the *F. adeliensis* acquisition of the endosymbiotic lifestyle is likely to be quite ancient, considering that a single IS element is present in its genome. In effect, a low number of mobile genetic elements is widely accepted to be a distinctive trait of an ancient stage of intracellular life and an expansion of these elements is regarded to be distinctive of initial stages of host restriction [55, 56]. In addition, the finding that *F. adeliensis* is auxotrophic for cysteine, lysine, methionine, and threonine, and likely depends on the host for nutrient supply, establishes a close physiological analogy with pathogenic strains of *F. tularensis*, whose virulence depends on the activity of the *Francisella* pathogenicity island cluster of genes [40]. In a mouse model of tularaemia, it has been shown that among these genes *F. tularensis* and *F. novicida* particularly need the expression of the *pdpC* gene in order to escape from phagosomes and become free in the cytosol [57]. Neither the *pdpE* gene, which is not directly involved in *F. tularensis* virulence, nor the *pdpC* gene was identified in the *F. adeliensis* genome. However, evidence from TEM analysis indicated that *F. adeliensis* may produce cytosolic stages in addition to

more common fusogenic membrane-bound structures closely recalling the "*Francisella* containing vacuoles" involved in the autophagy-mediated mechanism of *F. tularensis* re-entry into the endocytic compartment [58]. Although these cytosolic stages suggest that *Euplotes* is a potential ecological reservoir for the evolution of pathogenic *Francisella*, the observation that *F. adeliensis* is unable to proliferate at 37 °C would exclude any ability to colonize and be harmful to homothermic, warm-blood organisms.

Description of *Francisella adeliensis* Sp. Nov.

Francisella adeliensis (a.de.lien'sis. L. adj. of Adelie) is named after Adelie Cove, the location in Antarctica where the host, the ciliate *Euplotes petzi*, was collected in 2005 [21]. The type strain is deposited at the Swedish Defence Research Institute (FOI), *Francisella* strain collection # FSC1327. Within its host, *F. adeliensis* resides in the cytoplasm, as determined by TEM and FISH analysis carried out with the *Francisella*-specific probe Bwall1448 [34]. Cells are Gram-negative, roundish in shape, non-motile, catalase-positive, and oxidase-negative. They grow at a wide range of temperature (4–30 °C), salinity (0–35‰), and carbon dioxide concentrations (0.04–5%). The *F. adeliensis* complete genome sequence is available at GenBank, with the accession number CP021781, and supporting sequencing data are deposited in Bioproject PRJNA389235.

Compliance with Ethical Standards

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

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