Alkalilimnicola ehrlichii sp. nov., a novel, arsenite-oxidizing haloalkaliphilic gammaproteobacterium capable of chemoautotrophic or heterotrophic growth with nitrate or oxygen as the electron acceptor

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A facultative chemoautotrophic bacterium, strain MLHE-1T, was isolated from Mono Lake, an alkaline hypersaline soda lake in California, USA. Cells of strain MLHE-1T were Gram-negative, short motile rods that grew with inorganic electron donors (arsenite, hydrogen, sulfide or thiosulfate) coupled with the reduction of nitrate to nitrite. No aerobic growth was attained with arsenite or sulfide, but hydrogen sustained both aerobic and anaerobic growth. No growth occurred when nitrite or nitrous oxide was substituted for nitrate. Heterotrophic growth was observed under aerobic and anaerobic (nitrate) conditions. Cells of strain MLHE-1T could oxidize but not grow on CO, while CH4 neither supported growth nor was it oxidized. When grown chemoautotrophically, strain MLHE-1T assimilated inorganic carbon via the Calvin–Benson–Bassham reductive pentose phosphate pathway, with the activity of ribulose 1,5-bisphosphate carboxylase (RuBisCO) functioning optimally at 0.1 M NaCl and at pH 7.3. Strain MLHE-1T grew over broad ranges of pH (7.3–10.0; optimum, 9.3), salinity (15–190 g l⁻¹; optimum 30 g l⁻¹) and temperature (13–40°C; optimum, 30°C). Phylogenetic analysis of 16S rRNA gene sequences placed strain MLHE-1T in the class Gammaproteobacteria (family Ectothiorhodospiraceae) and most closely related to Alkalispirillum mobile (98.5%) and Alkalilimnicola halodurans (98.6%), although none of these three haloalkaliphilic micro-organisms were capable of photoautotrophic growth and only strain MLHE-1T was able to oxidize As(III). On the basis of physiological characteristics and DNA–DNA hybridization data, it is suggested that strain MLHE-1T represents a novel species within the genus Alkalilimnicola for which the name Alkalilimnicola ehrlichii is proposed. The type strain is MLHE-1T (=DSM 17681T = ATCC BAA-1101T). Aspects of the annotated full genome of Alkalilimnicola ehrlichii are discussed in the light of its physiology.

INTRODUCTION

Arsenic is a toxic metalloid of environmental concern. Despite its toxicity, certain microbes have evolved that can utilize arsenic for energy generation. Arsenic generally occurs in aquatic environments as either arsenate [HAsO₄²⁻ or As(V)] or arsenite [H₃AsO₃⁻ or As(III)], the latter oxyanion being more toxic than the former. The first As(III)-oxidizing micro-organism was isolated by Green (1918) and since then several prokaryotes with this aerobic

Abbreviations: AODC, acridine orange direct counts; PHB, polyhydroxybutyrate; RuBisCo, ribulose 1,5-bisphosphate carboxylase.

Further details of the methods used for PCR amplification and gene fragment analysis, graphs showing the growth of strain MLHE-1T under varying temperature, salinity and pH conditions, an additional phylogenetic tree, a plot showing RuBisCo activity in cell extracts of MLHE-1T and tables detailing lithotrophic growth of the novel strain on a variety of inorganic electron donors and on a diversity of organic and one-carbon electron donors are available as supplementary material in IJSEM Online.
metabolic capability have been described. Phylogenetically diverse, they include members of the *Alphaproteobacteria* and the *Betaproteobacteria* (Oremland & Stolz, 2003). As(III) oxidation has been noted in both heterotrophic and chemoaotrophic bacteria that have been isolated from a number of As-rich environments including cattle-dipping fluids, soils, sewage, mine drainage and hot springs (Ehrlich, 2002; Santini et al., 2000; Salmassi et al., 2002; Stolz et al., 2006).

*Thermus aquaticus* and *Thermus thermophilus* were found to rapidly oxidize As(III) to As(V), but were unable to grow with As(III) as the sole energy source (Ghihring et al., 2001). Similarly, *Hydrogenophaga* sp. strain NT-14 oxidized As(III) but, although it did not grow as a chemoautotroph, it nonetheless gained a small amount of energy from the reaction (vanden Hoven & Santini, 2004). Strain NT-26, a member of the *Rhizobium* clade of the *Alphaproteobacteria*, grew either as a chemoautotroph by oxidizing As(III) or as a heterotroph (Santini et al., 2000). The arsenic oxidases of these aerobic bacteria constitute a separate group within the broad family of molybdenum-containing enzymes and are composed of two heterologous subunits: the larger (~88 kDa) Mo-containing reactive centre protein and the smaller protein (~14 kDa) that contains a high potential Rieske functional group (Silver & Phung, 2005). In heterotrophic As(III) oxidizers, the enzyme is designated Aox, while in autotrophic As(III) oxidizers, it is designated Aro. Nonetheless, the Aro/Aox enzymes for both chemoaotrophs and heterotrophs that oxidize As(III) are, with some variations in amino acid sequences and in the multiplicity of their corresponding large and small subcomponents, essentially of the same genre (vanden Hoven & Santini, 2004; Santini & vanden Hoven, 2004).

In contrast to the above examples of aerobic As(III) oxidation, little is known about anaerobic arsenite oxidation. We first observed this phenomenon in experiments with *Rhodium* strain MLHE-1T which grows under anaerobic conditions using As(III) as the electron donor, nitrate as the electron acceptor and CO₂ (as bicarbonate) as the carbon source. Strain MLHE-1T is proposed as a non-photosynthetic, novel species of the genus *Ectothiorhodospira* of the class *Gammaproteobacteria*.

**METHODS**

**Culture conditions and strain characterization.** Strain MLHE-1T was isolated from Mono Lake water and maintained in an anaerobic liquid basal salts medium as described previously (Oremland et al., 2002) with 10 mM As(III) as the electron donor and 10 mM nitrate as the electron acceptor. Cells were also grown under heterotrophic conditions with 10 mM acetate as the electron donor and either oxygen (air head space) or 10 mM nitrate as the electron acceptor. Solid media were prepared by mixing 4 % (w/v) agar and double strength of the above basal salts media at 55–60 °C. Single colonies on solid media were grown aerobically with acetate as the electron donor. The standard growth conditions for the characterization of strain MLHE-1T were 28 °C, pH 9.8 and 6 % NaCl. Further experimental details on the ability of this organism to grow over wide ranges of pH, salinity and temperature, as well as its substrate affinities are presented as supplementary material in IJSEM Online.

**Electron microscopy.** For transmission electron microscopy, cells of MLHE-1T were grown either heterotrophically on plates or chemoaotrophically in liquid medium. Cells were fixed with 2.5 % glutaraldehyde in phosphate-buffered saline (plates) or by the addition of glutaraldehyde directly to the medium (liquid cultures) at a final concentration of 2.5 %. Negative staining of whole cells was performed with 1 % uranyl acetate on Formvar-coated grids. Thin sections of Spurr’s embedded samples were prepared following the procedure described by Switzer Blum et al. (1998). Samples were observed with a transmission electron microscope (100CX; JEOL) at 60 kV.

**Arsenic metabolism and amplification of araA.** All of the known araA gene sequences were aligned using CLUSTAL_X and conserved regions were identified. Primers were designed that would amplify a region of ~530 bp. These primers contained the *PstI* recognition sequence (underlined) and were: AraOdegF, 5′-GGCTGCA{-}CAGTGCYGTTGAGMAYGGTTA{-}3′ (binds at nucleotide position 85–108; numbering according to the *araA* gene sequence of strain NT-26) and AraOdegR, 5′-GGCTGCA{-}YTCDGARTTGT{-}AGC{-}YGBCG{-}3′ (binds at nucleotide position 622–599 on the *araA* gene sequence of NT-26). PCR conditions involved incubation at 90 °C, 3 min (1st cycle only); 92 °C, 1 min; 45–50 °C, 1.5 min; 72 °C, 1 min; 72 °C, 5 min (final cycle only) for 40 cycles. PCRs contained 100 ng DNA, 100 ng each primer, 10 % (v/v) DMSO and GoTag Green Master Mix (Promega). Chromosomal DNA of strain MLHE-1T was isolated using the Wizard Genomic DNA Purification Kit (Promega) and the method described by Humphreys et al. (1975) was used to detect the presence of a plasmid. No plasmid was detected.

**Phylogenetic analysis.** Sequences of the 16S rRNA genes of the closest relatives of strain MLHE-1T based on a BLAST search were obtained from GenBank. Sequence alignments were performed using CLUSTAL_X and a phylogenetic tree was constructed with maximum-parsimony using PAUP* 4.0b (Swofford, 2002).

The G+C content of the DNA (mol%) was determined by total genome analysis courtesy of the DOE Joint Genome Institute (http://genome.jgi-psf.org/draft_microbes/alkhe/alkhe.home.html).
Optical DNA–DNA hybridization assays of strain MLHE-1T against *Alkalilimnicola halodurans* DSM 13718T and *Alkalipirillum mobile* DSM 12769T were conducted in duplicate at the Deutsche Sammlung von Mikroorganismen und Zellkulturen (DSMZ) using 10% formamide in SSC buffer at 69°C (Huß et al., 1983).

**Metabolism of CH₄.** A 1 ml sample of cells of strain MLHE-1T was inoculated into 25 ml ‘Balch’-type culture tubes containing 10 ml basal salts medium sealed under a gas phase of 5% CH₄ in air. Tubes were incubated at 28°C with constant reciprocal shaking (150 r.p.m.). Growth was also attempted using 25 ml conical flasks containing 75 ml basal salts medium inoculated with 10 ml live cell suspensions sealed under a head space of 50% CH₄ and 50% air. Flasks were incubated at 30°C with rotary shaking (~150 r.p.m.). After 3 weeks of incubation, the head space was aseptically changed to 1000 p.p.m. CH₄ in air and further monitored for CH₄ consumption. Growth in both experiments was measured by acridine orange direct counts (AODC; Hobbie et al., 1977). A final washed cell suspension experiment was conducted using cells grown with acetate under air. Cells were centrifuged and washed three times with basal salts medium and then resuspended to a final cell density of 8.6 × 10⁶ cells ml⁻¹. A 10 ml sample of the suspension was dispensed into 37 ml serum bottles and sealed under an air atmosphere with 1000 or 100 p.p.m. CH₄. The suspensions were incubated at 30°C with rotary shaking (150 r.p.m.). CH₄ consumption in the above experiments was tracked by flame ionization gas chromatography (Oremland et al., 2005).

**Metabolism of CO.** *Alkalilimnicola halodurans*, *Alkalipirillum mobile* and strain MLHE-1T were grown under an air head space to stationary phase at 30°C in stoppered 160 ml serum bottles containing 10 ml basal salts medium with 0.05% yeast extract and 25 mM pyruvate in an artificial Mono Lake medium (King, 2003a). CO was added to the bottle head spaces (air, 1.3% O₂ in N₂ or 100% N₂) at starting concentrations ranging from 5 p.p.m. to 30% v/v. Nitrate (10 mM) was included as the electron acceptor for cells incubated under N₂ and a nitrate-free condition served as a live control. Head space subsamples were obtained at intervals using a needle and syringe; CO concentrations were determined by GC as previously described (King, 2003b). Details of the PCR amplification and analysis of *coxL* gene fragments are given in the supplementary material available in IJSEM Online.

**Determination of RuBisCO activity.** Details of the methods used for determination of RuBisCO activity in cell extracts are given in the supplementary material available in IJSEM Online.

**RESULTS AND DISCUSSION**

**Cell morphology and ultrastructure**

Strain MLHE-1T was isolated in a defined basal salts medium from Mono Lake anoxic bottom water (Oremland et al., 2002). Strain MLHE-1T is a rod-shaped bacterium with a length of approximately 1.5–2.5 μm and a diameter of 0.3–0.5 μm and has a negative reaction to Gram staining. Cells from liquid culture exhibit active motility when viewed by phase-contrast microscopy. TEM observations of negatively stained cells revealed a single polar flagellum along with several pili (Fig. 1a). Thin sections of cells grown on acetate revealed the presence of polyhydroxybutyrate (PHB) storage granules in the cytoplasm (Fig. 1b), while cells grown autotrophically (arsenite) lacked PHB granules, but instead contained intracellular inclusions (Fig. 1c).

**Growth and physiological characterization**

Colonies on aerobic agar plates with acetate as the electron donor appeared after 14 days of incubation at 28°C and were circular, convex, smooth and approximately 1 mm in diameter. Cells did not show any pigmentation when grown either on plates or in liquid culture and either in the dark or exposed to light. Growth occurred over a salinity range of 15–190 g l⁻¹ (0–175 g l⁻¹ of added NaCl), with an optimum at 30 g l⁻¹ (15 g l⁻¹ added NaCl). Strain MLHE-1T is alkaliophilic and exhibited an optimum growth rate at pH 9.3, but was unable to grow above pH 10.4. Optimum growth occurred at 30°C (see Supplementary Fig. S1 in IJSEM Online).

**Electron donors and acceptors**

Strain MLHE-1T has been previously shown to be capable of anaerobic, chemoheterotrophic growth with either As(III), H₂ or sulfide serving as the electron donor and with NO₃⁻ as the electron acceptor (Oremland et al., 2002). Strain MLHE-1T

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**Fig. 1.** Electron micrographs of cells of strain MLHE-1T. (a) Negatively stained cell grown heterotrophically on plates. F, flagellum. (b) Thin section of a cell grown heterotrophically on plates showing PHA globules (P). (c) Thin section of a cell grown autotrophically. Arrows point to intracellular inclusions. Bars, 0.25 μm.
also grew autotrophically on thiosulfate with nitrate as the electron acceptor (see Supplementary Table S1 in IJSEM Online). Under experimental conditions, strain MLHE-1T did not grow with oxygen as the electron acceptor with either As(III), sulfide or thiosulfate serving as electron donors, but enhanced aerobic growth on hydrogen was noted. Anaerobic growth was also noted on the As(III)-containing minerals orpiment and realgar, which were soluble in the alkaline basal salts medium, but no growth was noted with insoluble arsenopyrite. No growth was noted on a number of other potential inorganic electron donors, including Fe(II), NO\textsubscript{2}, Mn(II) and Sb(III).

Heterotrophic growth occurred on several organic acids, but not on any of the sugars or amino acids tested (see Supplementary Table S2 in IJSEM Online). Aerobic growth was observed on the following complex substrates: yeast extract, Casamino acids and casitone, but when nitrate served as the electron acceptor, growth occurred only on yeast extract. Growth did not occur with the following electron donors when As(V) (10 mM) was provided as the electron acceptor: lactate (10 mM), pyruvate (10 mM), succinate (10 mM), acetate (10 mM), glucose (10 mM), hydrogen (100 % in head space), sulfide (4 mM), yeast extract (0.5 %), Casamino acids (0.3 %) or casitone (0.3 %) (not shown). Finally, growth was not observed when N\textsubscript{2}O replaced nitrate and with acetate, As(III) or sulfide serving as the electron donors (not shown).

**Amplification of aroA**

Attempts to amplify a portion of the aroA gene of strain MLHE-1T were unsuccessful and no amplicons were obtained. Indeed, a BLAST search of the genome for homologues of the aroA gene did not turn up any positive responses, despite the confirmed ability of this organism to grow as a lithoautotroph using As(III) with nitrate (but not oxygen) as its electron acceptor (Supplementary Table S1; Oremland et al., 2004). Interestingly, two homologues of dissimilatory arsenate reductase (arrA) were identified as well as genes for arsenic resistance (arsC). Nevertheless, strain MLHE-1T was not able to grow using As(V) as an electron acceptor (Supplementary Table S1), nor did it produce As(III) when grown aerobically as a heterotroph in the presence of 5 mM As(V).

**Phylogenetic analysis**

The 16S rRNA gene sequence for strain MLHE-1T has been published previously (Oremland et al., 2002). However, four novel strains belonging to the genus have been isolated and characterized recently (Sorokin et al., 2006). These novel strains, Alkalilimnicola sp. AHN1, Alkalilimnicola sp. Z7008, Alkalilimnicola sp. AGDZ and Alkalilimnicola sp. ALPS2, share 98.8, 98.3, 98.2 and 97.3 % gene sequence similarity with strain MLHE-1T and, along with Alkalilimnicola halodurans (98.6 % gene sequence similarity) and Alkalilimnicola halodurans (98.5 % similarity), form a tight clade (Fig. 2).

The DNA G+C content of strain MLHE-1T is 67.5 mol% and is similar to that of Alkalilimnicola halodurans (66.2 mol%) and Alkalilimnicola halodurans (65.6 mol%) (Table 1). Since the G+C contents of DNA from these strains were so similar, hybridizations of DNA from strain MLHE-1T to DNA from Alkalilimnicola halodurans and Alkalilimnicola halodurans were performed. The DNA–DNA relatedness values (duplicates) of strain MLHE-1T with Alkalilimnicola...
Halodurans (45.7%; 42.6 %) and Alkalispirillum mobile (14.7%; 28.8 %) were lower than the recommended value of 70 % DNA–DNA similarity for strains belonging to the same species (Wayne et al., 1987). Because of the much closer similarity of strain MLHE-1T to members of the genus Alkalilimnicola, it is proposed that strain MLHE-1T represents a novel species in this genus.

**Arsenite oxidation in closely related species**

Two species closely related to strain MLHE-1T were tested for their ability to grow by oxidizing As(III), hydrogen and sulfide with nitrate as the electron acceptor (Table 1). Neither Alkalispirillum mobile nor Alkalilimnicola halodurans were capable of oxidizing As(III) or hydrogen with nitrate; however, both strains were able to oxidize sulfide with nitrate and growth was confirmed for both by AODC. Growth did not occur with arsenite when oxygen was provided as the electron acceptor (data not shown).

**Metabolism of CH4**

No growth or consumption of the 1 % CH4 head space was observed during incubation of shaken culture tubes (data not shown) or of the conical flasks (50 % CH4 in air) (see Supplementary Table S1). When the original flask head space was replaced with a 1 % CH4 in air mixture to allow monitoring of gas consumption, we did observe a near-complete loss of CH4 over an interval of about 3 weeks that continued upon the addition of more CH4 to the head space. Nonetheless, we were unable to attribute this observation to methane oxidation (as opposed to leakage) because incubated subaliquots of the aqueous phase neither consumed any added 14CH4 nor produced 14CO2 from this radiotracer (data not shown). When the flask experiment was repeated, no loss of head space CH4 was observed over prolonged incubation under identical conditions. In addition, no CH4 consumption was noted by washed cell suspensions. We conclude that strain MLHE-1T is not a

**Table 1.** Comparison of characteristics of strain MLHE-1T and those of its two closest taxonomic relatives

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<thead>
<tr>
<th>Characteristic</th>
<th>1</th>
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<th>3</th>
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<tr>
<td><strong>Morphology</strong></td>
<td>Rods</td>
<td>Oval rods</td>
<td>Vibrio</td>
</tr>
<tr>
<td>Salinity range (NaCl g l⁻¹)</td>
<td>0–175</td>
<td>0–280</td>
<td>0–250</td>
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<tr>
<td>Salinity optimum (NaCl g l⁻¹)</td>
<td>15</td>
<td>30–80</td>
<td>20</td>
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<tr>
<td>pH optimum</td>
<td>9.3</td>
<td>9.5</td>
<td>9–10</td>
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<tr>
<td>Temperature optimum (°C)</td>
<td>30</td>
<td>20–50</td>
<td>35–38</td>
</tr>
<tr>
<td>DNA G+C content (mol%)</td>
<td>67.5</td>
<td>65.6</td>
<td>66.2</td>
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<tr>
<td><strong>Mobility</strong></td>
<td>Single, polar flagellum</td>
<td>Subpolar flagella</td>
<td>Single, polar flagellum</td>
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<td>Substrates for growth (electron acceptor 10 mM nitrate)</td>
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<tr>
<td>10 mM As(III)</td>
<td>+</td>
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<td>–</td>
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<tr>
<td>100 % H₂</td>
<td>+</td>
<td>–</td>
<td>–</td>
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<tr>
<td>4 mM sulfide</td>
<td>+</td>
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methanotroph, nor does it have the ability to oxidize (but not grow on) CH₄, as it apparently does for CO (see below).

Our tests for the presence of aerobic methanotrophy in strain MLHE-1T were prompted by the identification of a methane monooxygenase C homologue (mnoC), formate dehydrogenase (e.g. fdhABC) and a serine hydroxymethyltransferase (glyA) in the genome of the novel strain. The latter suggested the presence of at least a partial downstream mechanism for C₁ oxidation and carbon fixation, with the former being confirmed by the ability of strain MLHE-1T to grow on formate (see Supplementary Table S2 in IJSEM Online). However, the full complement of mno genes is lacking from the genome of strain MLHE-1T and no methanol dehydrogenase homologues were found. Type I methanotrophs of the genus *Methylobacter* dominate the methane-oxidizing flora of Mono Lake, but molecular signals (pmoA amplicons) were also found for Type II methanotrophs of the *Methylocystis* genus (Lin et al., 2005). It is conceivable that there may have been some conjugative transfer of DNA between *Gammmaproteobacteria* of the *Methylobacter* type and those of the *Ectothiorhodospira* types such as strain MLHE-1T. However, Type I methanotrophs have the Calvin–Benson–Bassham pathway of C assimilation, while the genome of strain MLHE-1T has annotation for the serine pathway, a feature of Type II methanotrophs of the *Alphaproteobacteria* (Anthony, 1982). The serine transhydroxymethyltransferase (glyA) may function in serine biosynthesis from glycine.

**Metabolism of CO**

Strain MLHE-1T readily oxidized CO at a concentration range of between 100 and 1000 p.p.m. during stationary phase culture incubations (Fig. 3). When cultures were incubated with 30 % CO head space concentrations, uptake was initially negligible (7 days) and no growth was observed in the presence of this elevated level of CO. However, during extended incubations (>60 days), strain MLHE-1T reduced CO concentrations by about one-third to 20 %, but CO uptake was not coupled to an increase in cell density. *Alkalispirillum mobile* and *Alkalimnicola halodurans* also oxidized CO at concentrations <1000 p.p.m., with activity by the former greater than the latter. In addition, *Alkalispirillum mobile* reduced 30 % CO head space concentrations to values <5 % over a 60 day interval, but did not grow at the expense of CO.

Cell suspensions of pyruvate-grown strain MLHE-1T were able to consume trace concentrations of CO that mimicked ambient mixing levels of this gas in the lower troposphere. Cells oxidized the initially enclosed 5 p.p.m. CO and lowered the final head space concentrations to <0.1 p.p.m. (>99 % oxidation) within a few days of incubation (data not shown). Activity was noted in cells incubated under air, under microaerophilic conditions (1.3 % O₂) and under anaerobic conditions with nitrate as the electron acceptor. No significant CO oxidation by cells was noted under anaerobic conditions lacking nitrate. Anaerobic oxidation of CO by several diverse nitrate-respiring bacteria has been recently reported (King, 2006).

PCR amplification of strain MLHE-1T genomic DNA with primers specific for form I coxL resulted in a product, the sequence of which was identical to a sequence annotated as a putative large subunit CO dehydrogenase gene in the genome of strain MLHE-1T. This sequence clustered with other form I coxL partial gene sequences from members of the classes *Betaproteobacteria* and *Gammmaproteobacteria* after a neighbour-joining phylogenetic analysis; bootstrap support was high (82 % for 1000 bootstrap replicates; see Supplementary Fig. S2 in IJSEM Online). As in the case of CH₄ illustrated above, we were prompted to examine CO metabolism by strain MLHE-1T as the genome annotation indicated the presence of a CO dehydrogenase operon that included all three structural genes (coxFEDLSM). In this instance, however, we not only found active uptake of CO in this species, as well as in two neighbouring species (Fig. 3), but also successfully generated amplicons of coxL using established primer sets.

**RuBisCO activity**

Strain MLHE-1T grows as a chemoautotroph as previously demonstrated by its ability to fix ¹⁴CO₂ (added as ¹⁴C-bicarbonate) into cell material and by successful amplification (~800 bp fragment) of the *cbbL* gene of form I RuBisCO (Oremland et al., 2002). However, genome annotation indicates not only the presence of RuBisCO, with both large and small subunits (encoded by the *cbbL* and *cbbS* genes, respectively), but also the possible presence of components of a C₄ or reverse TCA cycle (phosphoenolpyruvate carboxylase; *ppc*) and a serine pathway (see CH₄ oxidation of the former greater than the latter. In addition,
section above). Therefore, we conducted enzyme assays with cell extracts to test for the actual presence of RuBisCO activity. A lysate from cells of strain MLHE-1T grown chemoautotrophically exhibited optimal RuBisCO activity at pH 7.7 and 140 mM (8.2 g L^{-1}) NaCl (see Supplementary Fig. S3 in IJSEM Online). These results were consistent with previous studies of enzymes obtained from halo- and alkaliphilic bacteria (Tabita & McFadden, 1976). At pH 8.5 (140 mM NaCl), lysates buffered with either Bicine-NaOH or CHES-NaOH showed nearly identical activities, indicating that CHES is an acceptable buffer at higher pH values. The lysate from strain MLHE-1T strongly cross-reacted with antisera raised against the Synecococcus 6301 RuBisCO and no cross-reactivity was observed using an antiserum that was raised against the Rhodospirillum rubrum form II enzyme (data not shown).

Maximal enzyme activities occurred at salinities and pH values that were considerably lower than those observed for optimal growth (see Supplementary Fig. S1). We attribute this result to the fact that the internal cytoplasmic pH and salinity of strain MLHE-1T are circumneutral and of low salinity compared with that of its external milieu, a common attribute of extremophiles. In addition, the genome shows the presence of PEP carboxylase as an alternate means of CO₂ fixation, however it is difficult to assess the presence of genes that encode the enzyme signature of the reductive TCA pathway in strain MLHE-1T since these reversible catalysts (pyruvate-ferredoxin oxidoreductase and 2-oxoglutarate-ferredoxin oxidoreductase) may also function in purely heterotrophic metabolism. In no instance has a functional Calvin–Benson–Bassham pathway been shown to be used in concert with the reductive TCA pathway of CO₂ assimilation, making it unlikely that the reductive TCA route is operational here.

CONCLUSION

Strain MLHE-1T has a highly flexible metabolism, capable of growth either as a heterotroph or as a chemoautotroph, with the additional ability to use either nitrate or oxygen as electron acceptors. In Mono Lake, this would allow it to survive as a lithotroph by exploiting the chemical redox gradients occurring within oxic/anoxic boundaries (i.e. chemocline) imposed by either seasonal (monomixis) or prolonged (meromixis) water column stratification (Oremland et al., 2004). In such an environment, it could couple the oxidation of reduced constituents (arsenite, sulfide) diffusing upwards from the anoxic bottom water with downward diffusing oxidants (nitrate, oxygen) from the aerobic epilimnion or the suboxic portion of the chemocline. It could also sustain itself as a conventional heterotroph by feasting upon the sinking remains of decomposing phytoplankton and zooplankton blooms. However, its ecological significance and abundance within Mono Lake has yet to be determined, as a survey of microbial diversity in this system did not detect this strain (Humayoun et al., 2003).

Nonetheless, such a flexible metabolism has also been observed in other arsenite-oxidizing bacteria of terrestrial origin, be they aerobes like strain NT-26 (Santini et al., 2000) or denitrifiers like strains DAO1 and DAO10 (Rhine et al., 2006). Such metabolic flexibility would give such strains a competitive survival edge in aquifers, for example, where nitrate, organic electron donors and arsenic species may all be present but at variable concentrations in both time and space (Kent & Fox, 2004). In the case of strains DAO1 and DAO10, they also exhibited similar autotrophic substrate affinities to strain MLHE-1T in that they could only grow on As(III) with nitrate, but in the case of H₂ were capable of aerobic or anaerobic growth (sulfide was not tested). These two strains conduct a full dissimilatory reduction of nitrate to N₂ while MLHE-1T will only reduce nitrate to nitrite and is incapable of N₂O reduction. Preliminary analysis of the full genome of strain MLHE-1T indicates a functional operon for respiratory nitrate reductase (narLXK₂GHJJ), as well as a nitric oxide reductase (norDQBC, Nor) and nitrous oxide reductase (nosLYDZR, Nos), but critically it lacks a respiratory nitrite reductase (nirK or nirS). Nevertheless, attempts to grow strain MLHE-1T on nitrous oxide were unsuccessful suggesting that Nor and Nos are inoperable (Zumft, 1997).

Finally, with regard to the oxidation of inorganic electron donors, the annotated genome of strain MLHE-1T lists several hydrogenases (mbhL3, hupL; large and small subunits of 'nickel-iron hydrogenase'). A diversity of uptake hydrogenases could explain our observation of H₂-dependent growth under both aerobic and anaerobic conditions (see Supplementary Table S1). However, with regard to sulfide and arsenite oxidation, the genome data are ambiguous. No sox genes were detected, however, homologues of genes encoding a putative sulfide dehydrogenase and sulfite reductase (dsrABFCHMKLJOP) have been identified. Nevertheless, determination of the enzyme(s) involved in these reactions awaits further biochemical analysis. Although we have clearly established that strain MLHE-1T can oxidize arsenite, we have not, however, found evidence for homologues of arsenite oxidase, either the catalytic subunit (AroA) or the Rieske subunit (AroB), and our unsuccessful attempt to amplify an aroA-like sequence would tend to confirm its absence. The presence of two arr homologues is also perplexing in that strain MLHE-1T is unable to grow under anaerobic conditions by using As(V) as an electron acceptor. This suggests that either strain MLHE-1T has a novel enzyme for oxidation of As(III) unrelated to Aro or that it employs one (or both) of its structurally similar Arr reductases (Silver & Phung, 2005) to oxidize As(III) by running in the reverse direction.

Description of Alkalilimnicola ehrlichii sp. nov.

Alkalilimnicola ehrlichii (ehr.li ch.ii. N.L. gen. n. ehrlichii of Ehrlich, named in honour of Professor Emeritus Henry Lutz Ehrlich for his broad scientific, teaching and leadership contributions to the field of geomicrobiology, with
specif... work on the bacterial oxidation of arsenic.

Cells are Gram-negative, motile rods (1.5–2.5 × 0.3–0.5 μm). Colonies on agar plates are circular, convex and smooth with a diameter of approximately 1 mm and are non-pigmented. The optimum pH and temperature are pH 9.3 and 30 °C, respectively. Growth occurs over a salinity range of 15–190 g NaCl l⁻¹, with an optimum of 30 g l⁻¹. Facultatively anaerobic and facultatively chemoautotrophic. Halkoalkaliphilic. Non-photorrophic. Capable of growth with inorganic electron donors (arsenite, hydrogen, sulfide and thiosulfate) and nitrate as the electron acceptor. Heterotrophic growth is observed under both aerobic and anaerobic (nitrate) conditions on a variety of organic acids, but not on sugars (glucose, fructose or galactose) or alcohols (methanol, ethanol). Unable to grow aerobically with other inorganic electron donors, except with hydrogen. Autotrophic growth occurs via the Calvin–Benson–Bassham Cycle. The G+C content of the DNA is 67.5 mol%.

The type strain, MLHE-1T (= DSM 17681T = ATCC BAA-1101T), was isolated from Mono Lake, an alkaline hypersaline soda lake, in California, USA.

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REFERENCES


