Diversity of Alkaliphilic Halobacteria: Proposals for Transfer of *Natronobacterium vacuolatum*, *Natronobacterium magadii*, and *Natronobacterium pharaonis* to *Halorubrum*, *Natrialba*, and *Natronomonas* gen. nov., Respectively, as *Halorubrum vacuolatum* comb. nov., *Natrialba magadii* comb. nov., and *Natronomonas pharaonis* comb. nov., Respectively

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The 16S rRNA genes of three species of the genus *Natronobacterium* (*Natronobacterium gregoryi*, *Natronobacterium pharaonis*, and *Natronobacterium vacuolatum*) were sequenced and compared to that of the previously sequenced species *Natronobacterium magadii*. The sequences revealed that *Natronobacterium pharaonis* was phylogenetically distinct from the other members of the genus and also from other recognized genera of the family Halobacteriaceae. However, *Natronobacterium vacuolatum* and *Natronobacterium magadii* were found to be most closely related to the genera *Halorubrum* and *Natrialba*, respectively. An unidentified haloalkaliphile, strain SSL1, was also closely related to *Natronobacterium magadii* and *Natrialba asiatica*. On the basis of phylogenetic tree reconstructions, signature bases specific for individual genera, and sequences of spacer regions between 16 and 23S rRNA genes, we propose the following changes: *Natronobacterium pharaonis* to be transferred to *Natronomonas* gen. nov. as *Natronomonas pharaonis* gen. nov., comb. nov.; *Natronobacterium vacuolatum* to be transferred to the genus *Halorubrum* as *Halorubrum vacuolatum* comb. nov.; and *Natronobacterium magadii* to be transferred to the genus *Natrialba* as *Natrialba magadii*.

The extremely halophilic, aerobic *Archaea* are classified within the family Halobacteriaceae, which currently contains nine valid genera: *Halobacterium*, *Halococcus*, *Haloarcula*, *Haloferax*, *Halorubrum*, *Halobaculum*, *Natrialba*, *Natronobacterium*, and *Natronococcus* (3, 5, 7, 13, 16, 17).

The alkaliphilic members of the Halobacteriaceae form a distinct physiological group as they require not only high NaCl concentrations but also high pH (between 8.5 and 11) and low Mg²⁺ concentrations (less than 10 mM) for growth. They have been isolated from a variety of alkaline, hypersaline lakes and soils (3, 15, 18, 20–22). Microscopically, the initial isolates consisted of rods and cocci and were accordingly separated into two genera, *Natronobacterium* and *Natronococcus*, by Tindall et al. (21). *Natronobacterium* currently contains four recognized species, *Natronobacterium gregoryi*, *Natronobacterium magadii*, *Natronobacterium pharaonis*, and *Natronobacterium vacuolatum*, with *Natronobacterium gregoryi* as the type species. The present classification of these isolates is based largely on chemotaxonomic criteria, particularly phenotypic properties and lipid composition. Glycolipid analysis, which has been of great use in the taxonomy of the neutrophilic halobacteria, has had little impact in the classification of the natronobacteria, as these lack major amounts of glycolipids in their membranes (3, 15). The complete sequence of the 16S rRNA gene of only one species, *Natronobacterium magadii*, has been reported, and

comparison to other halobacteria has clearly shown that it represents a distinct genus (7, 12). However, the sequences of the other three species of *Natronobacterium* (*Natronobacterium gregoryi*, *Natronobacterium pharaonis*, and *Natronobacterium vacuolatum*) have not been published, so phylogenetic support for their inclusion in this genus is lacking. DNA-DNA hybridization of the four species indicated that they have very little sequence homology (31 to 38%) (15), and while this is good evidence for their classification as separate species, it does not preclude their representing different genera.

The aim of this study was to determine the phylogenetic relationships among the natronobacteria and in particular to ascertain whether the four members of the genus *Natronobacterium* were indeed close relatives. For this purpose, 16S rRNA sequences of the three unsequenced species of *Natronobacterium* and of one unclassified haloalkaliphilic strain, SSL1, were determined, as well as sequences from four other strains of halobacteria. These sequences were incorporated into phylogenetic tree reconstructions of the family Halobacteriaceae.

MATERIALS AND METHODS

Bacterial strains and culture conditions. The following strains of the genus *Natronobacterium* were used in this study: *Natronobacterium gregoryi* NCIMB 2189^T, *Natronobacterium pharaonis* JCM 8858^T (derived from DSM 2160^T via IFO 14720^T), *Natronobacterium vacuolatum* JCM 9060^T (deposited by W. D. Grant), and strain SSL1 (microscopically long rods, originally isolated from an alkaline saline brine from Sambhar Salt Lake, India, and deposited with the American Type Culture Collection as *Natronobacterium* sp. strain ATCC 43988) (22, 23). These strains were grown at 37°C in haloalkaliphile medium as previously described (20, 23).

Halobacterium trapanicum NCIMB 767 was obtained from the National Collection of Industrial and Marine Bacteria in 1994 and deposited as JCM 8979. Three species of the genus *Halococcus* (*Halococcus saccharolyticus* ATCC 49257,

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Halococcus salifodinae DSM 8989, and *Halococcus turkmenicus* VKM B-1734) were also obtained from culture collections and cultivated as described previously (7).

Sequencing of 16S rRNA genes and thin-layer chromatographic analysis of polar lipids. Genomic DNAs from the four species of *Natronobacterium* were extracted as described before (7). *Halobacterium trapanicum* JCM 8979 and species of *Halococcus* which do not lyse with *N*-lauroylsarcosine were ground with quartz sand, and DNA was recovered by ethanol precipitation. 16S rRNA-encoding genes were amplified by PCR, cloned into plasmid pUC119, and sequenced (7, 8). Final sequences of these genes were unambiguous, as regions which were difficult to read on one strand were easily read by sequencing the complementary strand. Phylogenetic tree reconstructions based on the 16S rRNA sequences were performed as described previously (7).

Membrane lipids were extracted, and thin-layer chromatography was performed as described before (7).

Sequencing of spacer regions. Genomic DNAs from five species of the genus *Halorubrum* (*Halorubrum saccharovorum*, *Halorubrum sodomense*, *Halorubrum lacusprofundi*, *Halorubrum coriense*, and *Halorubrum distributum*) were extracted (7) and kept at -30°C . Spacer regions between the 16S rRNA and 23S rRNA genes from the three species of the genus *Natronobacterium* and the five species of the genus *Halorubrum* were amplified by PCR with a set of specific oligonucleotide primers. The sense primer, 5'-TTAAGTCGTAACAAGGTAGCCGTAG, was based on the sequence from nucleotides 2332 to 2356 of the *Halobacterium halobium* rRNA operon (accession no. X03407), and the antisense primer, 3'-CCTGCACGGTTCGACGCTATTCG, was designed based from nucleotides 2959 to 2981 of the same sequence. Both primers are situated in highly conserved regions. PCR products, which were expected to overlap by 51 bp with the 3' termini of the 16S rRNA genes and by 71 bp with the 5' termini of the 23S rRNA genes, were cloned and sequenced as described above. Sequences of spacer regions from the following halobacteria were taken or extracted from operon sequences deposited with databases: *Halobacterium cutirubrum* K02971, *Halobacterium halobium* X03407, *Halococcus morrhuae* X72588, *Haloferax volcanii* M19341, and *Natronobacterium magadii* X72495.

Nucleotide sequence accession numbers. The 16S rRNA gene sequences for *Natronobacterium gregoryi*, *Natronobacterium pharaonis*, *Natronobacterium vacuolatum*, *Natronobacterium* sp. strain SSL1, and *Halobacterium trapanicum* JCM 8979 were deposited in the DDBJ under accession numbers D87970, D87971, D87972, D88256, and D63786, respectively.

The sequences of the spacer regions for *Halorubrum sodomense*, *Halorubrum saccharovorum*, *Halorubrum lacusprofundi*, *Halorubrum distributum*, *Halorubrum coriense*, *Natronobacterium pharaonis*, *Natronobacterium gregoryi*, and *Natronobacterium vacuolatum* were deposited in the DDBJ under accession numbers AB003404, AB003405, AB003406, AB003407, AB003408, AB003409, AB003410, and AB003411, respectively.

RESULTS

Nucleotide sequences of 16S rRNA-encoding genes. The 16S rRNA genes from four alkaliphilic halobacteria (*Natronobacterium gregoryi*, *Natronobacterium pharaonis*, *Natronobacterium vacuolatum*, and strain SSL1) were sequenced and compared to the previously determined sequences of *Natronobacterium magadii*, two species of the genus *Natronococcus*, and other closely related species. The lengths of the genes from the five *Natronobacterium* strains were 1,465 to 1,468 bp. The longest 16S rRNA gene from the *Halobacteriaceae* reported so far is that of *Halococcus morrhuae* (1,475 bp), and the shortest one is that of *Natronobacterium pharaonis* (1,465 bp).

Devereux et al. (1) and Fry et al. (2) proposed that a 16S rRNA similarity value less than 93 to 95% indicated that two organisms belonged to different genera. Surprisingly, when the similarity values of the four recognized members of the genus *Natronobacterium* were examined, all were below 92% (the highest similarity, 91.9%, was observed between *Natronobacterium gregoryi* and *Natronobacterium magadii*), indicating that the four species may be sufficiently different to warrant classification in different genera. *Natronobacterium pharaonis* and *Natronobacterium vacuolatum* showed particularly weak similarities to each other (88.3%), to *Natronobacterium gregoryi* (88.5 and 88.1%, respectively), and to *Natronobacterium magadii* (88.8 and 87.1%, respectively). *Natronobacterium magadii* was most similar to the strain SSL1 (95.9%), and similarity to *Natrialba asiatica* 172P1 was 93.3%. The similarity of strain SSL1 to *Natrialba asiatica* was even greater (96.6%).

The 16S rRNA gene from *Halobacterium trapanicum* JCM

8979 (a culture obtained from the National Collection of Industrial and Marine Bacteria in 1994; see Materials and Methods) was also sequenced and found to be almost indistinguishable from those of two *Halococcus morrhuae* strains (98.6 to 98.9% sequence similarity).

Phylogenetic tree reconstruction. The sequences of almost all of the 16S rRNA genes determined so far by us (25 of 37 sequences, including those determined in this work) are unambiguous. Although the 5'- and 3'-terminal sequences of the genes are those of the PCR primers, 20 bp each, these regions are highly conserved as revealed by several sequences determined from genes obtained by the conventional cloning methods. Replacement of C by T in one position in the 3'-terminal antisense primer region has been reported only for the two 16S rRNA genes (i.e., A and B) of *Haloarcula marismortui*. Several strains also show replacement of the 3'-terminal T by A. The sequences determined in the present study were aligned with those previously reported (by us and by other groups), and used to reconstruct phylogenetic trees. Positions with any deletions or uncertain alignment were removed, and the remaining 1,371 positions were used to construct the trees. A variety of algorithms were utilized (maximum likelihood, maximum parsimony, and distance matrix [7]), which gave very similar topologies. A representative example is shown in Fig. 1. As anticipated from the similarity values, the four recognized *Natronobacterium* species and strain SSL1 did not form a monophyletic group but were distributed throughout the tree. *Natronobacterium gregoryi*, the type species, branched just before the *natronococci*, whereas *Natronobacterium magadii* and strain SSL1 branched before *Natrialba*. The exact branching order of *Natronobacterium magadii* and SSL1 was not clear in parsimony or distance matrix trees, but the bootstrap values indicate a clear separation of *Natronobacterium magadii* and *Natronobacterium gregoryi*. *Natronobacterium vacuolatum* was most closely related to the genus *Halorubrum*.

Natronobacterium pharaonis turned out to be only distantly related to the other genera. Even the sequence most similar to that of *Natronobacterium pharaonis*, i.e., *Haloarcula marismortui* A, showed a low similarity value of 89.9%. The lack of bootstrap values at the branch point indicates that parsimony and distance methods could not confidently identify the deepest branching order in the tree (bootstrap values, 54 to 55%). However, the maximum likelihood method (the tree shown) gave strongly supported branch lengths in this region. The evidence from sequence similarity and tree reconstructions appears to present a clear case for this isolate representing a new genus.

Signature bases of 16S rRNA genes. By using the aligned sequences, a number of signature sequences (24) which were specific for each genus were revealed. These are as follows: *Halobacterium* (*Halobacterium cutirubrum*, *Halobacterium halobium*, and strain Y12), 77C, 106G, 128T, 213C, 222T, 407A, and 1383T; *Haloarcula* (*Haloarcula marismortui*, *Haloarcula sinaiensis*, *Haloarcula vallismortis*, and *Haloarcula aidiensis*), 235T, 255A, 665A, 670G, 675C, 883C, 937T, 968C, 972T, 1173A, 1200T, 1202T, 1243- (- means a gap), 1250A, and 1458A; *Haloferax* (the four recognized species), 1109A, 1218T, 1220T, and 1227A; *Halococcus* (two strains of *Halococcus morrhuae*, *Halococcus saccharolyticus*, *Halococcus salifodinae*, and *Halobacterium trapanicum* JCM 8979, but not *Halococcus turkmenicus*), 25T, 46C, 90C, 200C, 211C, 215A, 495A, 499A, 1246C, 1271T, and 1280A; *Natrialba* (*Natrialba asiatica*), 385G, 502G, and 1108A; *Halobaculum* (*Halobaculum gomorense*), 106T, 213A, 246T, 521T, 603A, 690T, 702C, 706A, 777C, and 943T; *Natronobacterium gregoryi* and two *Natronococcus* spp. (*Natronococcus occultus* and *Natronococcus amylolyticus*),

FIG. 1. Phylogenetic tree reconstruction of the Halobacteriaceae with complete 16S rRNA sequences. The representative tree shown here was derived by using maximum likelihood (fDNAml). Branches with poorly supported lengths have been collapsed. Bootstrap values (100 replicates) shown at the nodes were obtained by parsimony (above) and distance matrix (below) methods and added to the maximum likelihood tree. Only bootstrap values above 70% are shown. The scale bar represents 0.02 expected changes per site. *Methanospirillum hungatei* (accession no. M60880) was used as the outgroup.

1312C, 1313G, 1324C, and 1325G; *Halorubrum* (six recognized species and *Natronobacterium vacuolatum*; bases which are confined to the six recognized species are underlined), 114T, 179A, 189C, 206A, 611G, 683C, 783C, 794G, 795G, 818C, 1070C, 1087G, 1101G, 1105G, 1115C, 1121G, 1127C, 1208A, and 1274C; and *Natronobacterium pharaonis*, 55G, 64-, 65-, 71-, 72-, 73-, 85T, 109G, 381A, and 477C. The sequence numbers refer to the refined alignment (1 to 1483) as reported previously (8).

Sequences of spacer regions. Lengths (in base pairs) of the total spacer regions ranged from 528 (*Halobacterium halobium*) to 377 (*Natronobacterium pharaonis*). The values are as follows, together with the lengths before and after tRNA^{Ala} genes: *Halobacterium cutirubrum*, 509, 105, and 332; *Halobacterium halobium*, 528, 105, and 351; *Halococcus morrhuae*, 433, 148, and 213; *Haloferax volcanii*, 391, 97, and 222; *Natronobac-*

terium magadii, 405, 130, and 204; *Natronobacterium gregoryi*, 456, 154, and 230; *Natronobacterium pharaonis*, 377, 95, and 210; *Natronobacterium vacuolatum*, 480, 167, and 241; *Halorubrum saccharovororum*, 514, 161, and 281; *Halorubrum sodomense*, 517, 164, and 281; *Halorubrum lacusprofundi*, 509, 158, and 279; *Halorubrum coriense*, 512, 163, and 277; and *Halorubrum distributum*, 517, 164, and 281. Another species of the genus *Halorubrum*, *Halorubrum trapanicum* NRC 34021, was not available from the National Research Council since this strain died out at the National Research Council (11a).

Alignment of the 13 sequences was difficult and required the insertion of many gaps. However, it was clear that *Natronobacterium vacuolatum* showed the greatest similarity to members of the genus *Halorubrum*. The five recognized species of *Halorubrum* were easily aligned with *Natronobacterium vacuolatum*, and the alignment showed long stretches of conserved

sequence. If sequences from members of the other genera were introduced, the alignment was severely disrupted. This was also reflected by the tRNA^{Ala} genes (72 bp), which were largely conserved among the five *Halorubrum* spp. and *Natronobacterium vacuolatum*, except for a one-base difference in *Halorubrum lacusprofundi* (³²C→T). Alignment with the other seven tRNA^{Ala} genes revealed differences in nine bases. Three of these (bases 5, 17, and 68) are conserved among the five recognized species of the *Halorubrum*-*Natronobacterium vacuolatum* group but differ from the other sequences, at bases 5 and 68, which form a hydrogen bond, T and A in the *Halorubrum*-*Natronobacterium vacuolatum* group but C and G in the rest; and at base 17, which is in the D arm of the cloverleaf secondary structure, C in the *Halorubrum*-*Natronobacterium vacuolatum* group but T in the rest.

DISCUSSION

In this study, phylogenetic tree reconstructions have clearly shown the genus *Natronobacterium* as being polyphyletic, i.e., it contains a diverse group of organisms that show closer relationships to widely differing groups within the family *Halobacteriaceae*. Each of the four current species probably belongs to a separate genus. Unlike the other halobacterial genera, which possess a variety of membrane glycolipids, the haloalkaliphiles contain only trace amounts or no glycolipids. Though phosphatidylglycerol and phosphatidylglycerophosphate-methyl ester are the main membrane phospholipids of the alkaliphiles as in the other genera, minor amounts of additional phospholipids (PL1 to PL4; only the structure of PL2 has been determined previously [11]) have been detected by thin-layer chromatography among the haloalkaliphiles (14, 15). *Natronobacterium gregoryi* possesses phospholipids PL1 and PL3, *Natronobacterium pharaonis* has PL1, *Natronobacterium magadii* has PL3, and *Natronobacterium vacuolatum* contains PL3 and PL4. On the other hand, *Natronococcus occultus* has PL1 and PL2, while *Natronococcus amylolyticus* does not contain PL2 (9).

Since the phylogenetic data presented in this study indicate that each of the four species of the genus *Natronobacterium* probably belongs to a separate genus, the known differences in minor phospholipids may be considered as additional taxonomic markers.

We believe that the four species of the genus *Natronobacterium* belong to different genera as discussed below.

Natronobacterium gregoryi, which clustered tightly with the two species of the genus *Natronococcus*, should remain the type species of the genus *Natronobacterium*. This is supported by the four signature bases described above.

Natronobacterium pharaonis was only distantly related to the other taxa and would appear to represent a novel genus. Sequence comparisons with other 16S rRNA genes, including that of the recently proposed species *Haloarcula mukohataei* (4), showed similarities of less than 89.9%. Ten signature bases were also observed. We propose to transfer *Natronobacterium pharaonis* to *Natronomonas* gen. nov.

Natronobacterium vacuolatum showed a 16S rRNA sequence similarity of 95.6% to *Halorubrum saccharovororum*, the type species, indicating that they may belong to the same genus. In addition, a close examination of the 37 aligned sequences, used in reconstructing the tree, has revealed many sequence signatures. As described in Results, sequences of the six species of *Halorubrum* possess 19 genus-specific signature bases (i.e., not found in the other 31 sequences). The *Natronobacterium vacuolatum* sequence shares 16 of these 19 bases (the three bases which are confined to the six recognized species of *Halorubrum*

are underlined). On the other hand, there are two positions (781 and 1406) where five of the six species and *Natronobacterium vacuolatum* have the same specific signature bases and two positions (1264 and 1265) where *Halorubrum coriense* (only) has different bases from the other 35 sequences. Furthermore, six bases, 785A, 793T, 1234G, 1242A, 1267T, and 1284A, were confined to the six recognized species of the genus *Halorubrum*, *Natronobacterium vacuolatum*, and *Halobaculum gomorrense*. Altogether, these data seem to support the view that *Natronobacterium vacuolatum* is related to members of the genus *Halorubrum*. No glycolipids have been detected in *Natronobacterium vacuolatum* (15), whereas most of the species of the genus *Halorubrum* are characterized by the presence of glycolipid sulfated diglycosyldiether-3 (S-DGD-3) (7), but an exception has been reported, i.e., sulfated diglycosyldiether-5 (DGD-5) for *Halorubrum trapanicum* NRC 34021 (13). We cannot exclude the possibility that a trace amount of glycolipid is present in *Natronobacterium vacuolatum*.

Given the evidence from 16S rRNA gene sequences and the additional data from spacer region sequences, we believe there is a strong case to support the transfer of *Natronobacterium vacuolatum* to the genus *Halorubrum* and for the strain to be called *Halorubrum vacuolatum* (comb. nov.).

The 16S rRNA gene of *Natronobacterium magadii* showed a high sequence similarity to that of strain SSL1 (95.9%) and to those of *Natrialba asiatica* 172P1^T (type strain) and *Natrialba asiatica* B1T (93.3 and 93.7%, respectively). The similarity between SSL1 and *Natrialba asiatica* 172P1 and that between SSL1 and *Natrialba asiatica* B1T were 95.9 and 96.6%, respectively. These figures (pairwise comparisons) seem to suggest that *Natronobacterium magadii*, strain SSL1, and *Natrialba asiatica* may be members of the same genus, *Natrialba*. On the other hand, the lipid analyses do not support the close 16S rRNA sequence similarities. No glycolipid has been detected in *Natronobacterium magadii*, but strain SSL1 has a minor amount of glycolipid DGD-4 (glucose 1→6 glucose-glycerol diether [10, 23]). *Natrialba asiatica* has a glycolipid, S2-DGD (mannose-2,6-disulfate 1→2-glucose-glycerol diether [7]). The structures of these lipids suggest that the enzymes involved in the biosynthesis are different between the two. However, one of the two strains of *Natrialba asiatica*, B1T, is able to grow at alkaline pH (from 5 to 10) (7). Furthermore, *Natronobacterium magadii* and strain SSL1 do share one of the three signature bases of the genus *Natrialba*, 502G, and strain SSL1 shares 1108A. Moreover, 568A is confined to *Natrialba asiatica* B1T, *Natronobacterium magadii*, and strain SSL1. Altogether, these data (high similarities in 16S rRNA genes, signature bases, and growth at alkaline pH) seem to support the view that *Natronobacterium magadii* and strain SSL1 are members of the genus *Natrialba*, and we propose to transfer *Natronobacterium magadii* to the genus *Natrialba* as *Natrialba magadii* and to call strain SSL1 *Natrialba* sp. strain SSL1.

The inferred tree (Fig. 1) also includes three species of the genus *Halococcus* (*Halococcus saccharolyticus*, *Halococcus salifodinae*, and *Halococcus turkmenicus*) and an unvalidated species of the genus *Haloarcula*, *Haloarcula aidinensis* (25). Validation of the latter species is currently in progress (26). Further taxonomic investigation of the three species of *Halococcus* will be described in a separate paper.

Description of *Natronomonas* gen. nov. *Natronomonas* Kamekura, Dyall-Smith, Upasani, Ventosa, and Kates (Na.tro.no.mo'nas. Gr. n. *natron*, derived from Arabic *natrun*, soda [sodium carbonate], Gr. fem. n. *monas*, unit; M.L. fem. n. *Natronomonas*, the soda unit). Gram-negative rods. Colonies are pigmented red because of the presence of C₅₀-carotenoid.

Chemolithotrophic and aerobic. Halophilic, requiring at

least 2 M NaCl. Alkaliphilic, growing between pH 7 and 10 with optimum at 8.5. Moles percent G+C content of DNA is 64.3 (T_m), 61.2 to 62.1 (buoyant density). Polar lipids are glycerol-diether analogs of phosphatidylglycerol, methyl ester of phosphatidylglycerophosphate, phosphatidic acid, and an unknown phospholipid, PL1. Both diphytanyl moieties (C_{20} and C_{20}) and phytanyl-sesterterpanyl moieties (C_{20} and C_{25}) detected.

Type species is *Natronomonas pharaonis*.

Description of *Natronomonas pharaonis* (Soliman and Trüper 1982) comb. nov. (basonym, *Natronobacterium pharaonis*). The first alkaliphilic extreme halophile described. Pleomorphic motile rods, originally isolated from the alkaline brines of eutrophic desert lakes of Wadi Natrun, Egypt. Detailed characteristics are described elsewhere by Soliman and Trüper (18).

Type strain is Gabara, deposited with the Deutsche Sammlung von Mikroorganismen as DSM 2160^T = ATCC 35678^T = NCIMB 2260^T.

Strain SP1, which showed 96% homology with DSM 2160^T in DNA-DNA hybridization, was isolated from a Kenyan soda lake, Lake Magadi, and deposited as NCIMB 2191 (21). NCIMB 2191 contains a small amount of disesterterpanyl moiety (C_{25} and C_{25}), whereas DSM 2160^T does not (19).

Description of *Halorubrum vacuolatum* (Mwatha and Grant 1993) comb. nov. (basonym, *Natronobacterium vacuolatum*). The properties of the type strain M24 (NCIMB 13189) were described previously (15).

Description of *Natrialba magadii* (Tindall et al. 1984) comb. nov. (basonym, *Natronobacterium magadii*). The properties of the type strain MS3 were described previously (21).

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