

# Microbial Diversity in Natural Asphalts of the Rancho La Brea Tar Pits<sup>∇</sup>

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**Bacteria commonly inhabit subsurface oil reservoirs, but almost nothing is known yet about microorganisms that live in naturally occurring terrestrial oil seeps and natural asphalts that are comprised of highly recalcitrant petroleum hydrocarbons. Here we report the first survey of microbial diversity in ca. 28,000-year-old samples of natural asphalts from the Rancho La Brea Tar Pits in Los Angeles, CA. Microbiological studies included analyses of 16S rRNA gene sequences and DNA encoding aromatic ring-hydroxylating dioxygenases from two tar pits differing in chemical composition. Our results revealed a wide range of phylogenetic groups within the *Archaea* and *Bacteria* domains, in which individual taxonomic clusters were comprised of sets of closely related species within novel genera and families. Fluorescent staining of asphalt-soil particles using phylogenetic probes for *Archaea*, *Bacteria*, and *Pseudomonas* showed coexistence of mixed microbial communities at high cell densities. Genes encoding dioxygenases included three novel clusters of enzymes. The discovery of life in the tar pits provides an avenue for further studies of the evolution of enzymes and catabolic pathways for bacteria that have been exposed to complex hydrocarbons for millennia. These bacteria also should have application for industrial microbiology and bioremediation.**

Prior studies of subsurface petroleum reservoirs using culture-based methods have revealed diverse microbial communities that are able to live on complex petroleum hydrocarbon mixtures (22, 43). Nonetheless, very little is known yet about the true extent of microbial diversity in natural oil reservoirs and terrestrial petroleum deposits, such as those that occur in oil sands, shales, and natural asphalts. Initial surveys of underground reservoirs using molecular approaches so far suggest that the majority of microorganisms inhabiting these environments are new species that represent a rich pool of novel genetic diversity with potential importance for industrial and petroleum microbiology (43). Compared to underground oil reservoirs, even less is known about terrestrial habitats, where petroleum-degrading soil bacteria have come to inhabit heavy oil seeps, tar sands, and natural asphalts. With the advent of improved DNA extraction and purification methods, such bacteria and their genes may now be accessible for detailed study of their diversity and genes that encode petroleum-degrading enzymes.

The existence of bacteria in petroleum deposits at great depths suggests that many species have evolved specifically for this environment and may be carried to the surface in oil seeps. In soils that are permeated with the asphalt, bacteria may also include indigenous bacteria that survived after asphalt seeped through the soil. Selection of bacterial communities for petroleum substances occurs rapidly after even short-term exposures of soil to petroleum hydrocarbons following oil spills (41, 42). Over time spans encompassing millennia, bacteria that can tolerate this environment would be expected to undergo genetic adaptations that may lead to evolution of new ecotypes and species and enzymes for growth on petroleum hydrocarbons. During adaptation of communities, genes for petroleum hydrocarbon-degrading enzymes that are carried on plasmids

or transposons may be exchanged between species. In turn, new catabolic pathways eventually may be assembled and modified for efficient regulation (27). Other cell adaptations leading to new ecotypes may include modifications of the cell envelope to tolerate solvents (28) and development of community-level interactions that facilitate cooperation within consortia.

Here we describe a survey of microbial diversity in natural asphalts at the Rancho La Brea Tar Pits in California. These natural asphalts are located in Hancock Park in downtown Los Angeles and consist of asphalt-soil mixtures formed by upwelling of heavy oil and asphaltenes in spatially separated seeps that differ in their chemical composition and age. Although the asphalt at Rancho La Brea is commonly called tar, the petroleum hydrocarbons here are correctly referred to as natural asphalt and are comprised of some of the most recalcitrant carbon compounds in nature (9). Our survey examined two excavation sites. The first site, Pit 91, has yielded thousands of plant and animal fossils and is the richest Pleistocene fossil site in the world (10, 16, 45). Carbon dating of fossils from the current depth under excavation in Pit 91 fixes their ages in a range from 10,000 to 38,000 years before the present (16). The second site, Pit 101, was excavated early last century and was closed in the 1920s, after which the pit was covered with a permanent building as part of a museum display. Microbiological studies included analysis of 16S rRNA genes from DNA extracted from the tar pits and a traditional approach employing cultivation of bacteria on agar. In conjunction with microbial diversity, we also surveyed DNA sequences for aromatic ring-hydroxylating dioxygenases that may have application for industrial microbiology and bioremediation of petroleum wastes.

## MATERIALS AND METHODS

**Characterization of chemical properties in Pits 91 and 101.** Ten-gram samples of the asphalt-permeated soil were physically broken into small aggregates and placed in beakers containing 10 ml deionized water or 10 mM CaCl<sub>2</sub> buffer. The

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TABLE 1. Chemical properties of asphalt-permeated soil samples from Pit 91 and Pit 101 of Rancho La Brea Tar Pits<sup>a</sup>

Sample site	pH of suspension		EC ( $\mu\text{S}/\text{cm}$ )	% C	% N	% S	C/N ratio	Concn ( $\mu\text{g}/\text{ml}$ ) of:												
	1:1 ( $\text{CaCl}_2$ )	1:5 ( $\text{H}_2\text{O}$ )						Na	Ca	Mg	K	Al	Cd	Cr	Fe	Zn	Cu	Mn	Pb	Ni
Pit 91	6.3	5.36	46	22.04	0.3	0.97	74.16	13	2,902	265	86.1	356	2.2	2.65	484	33.9	23.4	13	ND	89.9
Pit 101	8.44	7.59	4,610	7.04	0.18	0.53	46.74	178	3,696	4,245	1,011	11,395	1.52	19.9	14,133	126	37.1	178	58.1	39.4

<sup>a</sup> EC, electrical conductivity; ND, not detected.

suspensions were mixed using a magnetic stir bar for 1 h, after which pH and salinity were determined. Salinity was determined in water solutions using a conductivity meter. Other samples for metal analyses were acid digested in nitric perchloric acid using digestion bombs and a microwave oven (USEPA SW-846, method 3051). Heavy metals and cations were analyzed by inductively coupled plasma mass spectrometry using an ELAN500 mass spectrometer (Perkin-Elmer-Sciex Instruments, Concord, Ontario, Canada). Carbon, nitrogen, and sulfur were analyzed using a Flash EA 1112 NC analyzer (Thermo Electron Corporation, Milan, Italy).

**Sampling and DNA extraction.** Previously unexposed samples were removed from approximately 10 cm under the surfaces of Pit 91 and Pit 101 of the Rancho La Brea Tar Pits in Los Angeles in October 2004. A total of five samples were taken along a 3-m transect from the center of each pit. Samples were removed from the pits with sterile, autoclaved spatulas and were transferred into sterile 50-ml plastic tubes with screw caps and transported to the laboratory for processing. The samples from each pit were pooled prior to extraction. One of the challenges in conducting this survey was the difficulty in extracting high-quality DNA for use in cloning and sequencing. DNA was extracted from the asphalt-soil mixtures by first freezing approximately 5-g aliquots of the asphalt in liquid nitrogen. The frozen samples were transferred to a sterile ceramic mortar and were then ground under liquid nitrogen to a fine powder. Approximately 0.5-g subsamples were processed to extract DNA by bead beating using the BIO 101 Fastprep DNA extraction kit for soil following the manufacturer's protocols. Extracted DNA was concentrated using a Savant Speed Vac system (GMI Inc., Ramsey, MN), and subsamples were combined to obtain a high concentration of DNA. The DNA was purified using a QIAquick gel extraction kit (QIAGEN, Chatsworth, CA) according to the manufacturer's instructions. The purified DNA was concentrated again for use in construction of clone libraries.

**Phylogenetic analysis.** 16S rRNA genes were amplified by PCR and purified with a QIAquick PCR purification kit (QIAGEN). The purified PCR products from five runs were combined and used to construct clone libraries using the pGEM-T Easy vector (Promega) with selected primer sets. The *Bacteria* were detected using bacterium-specific primers, 27F and 1492R (21). The *Archaea* were detected using the domain-specific PCR primers, Ar4F and Ar958R (17). *Pseudomonas* spp. were identified using the *Pseudomonas*-selective PCR primers, Ps289F and Ps1258R (47). Dioxygenases were detected using the PCR primer set adoF to adoR for aromatic ring-hydroxylating dioxygenases (36, 37). The sequencing primers were T7 and SP6. After obtaining raw sequences using Chromas 2 (Technelysium Pty. Ltd., Tewantin, Queensland, Australia), putative chimeric sequences were identified using Bellerophon (12) and identified chimeric sequences (43 of 278 bacterial sequences) were excluded. The 16S rRNA sequences were aligned using the NAST aligner, and the aligned sequences were compared to the Lane mask (21) using the Greengenes web site (4, 5). Evolutionary distances were calculated with the Kimura 2-parameter method, and a phylogenetic tree was constructed by the neighbor-joining method (31) with MEGA3 for Windows (19). Bootstrap analyses of the neighbor-joining data were conducted based on 1,000 samples to assess the stability of the phylogenetic relationships.

**Statistical analyses.** The computer program DOTUR (32) was used to calculate species richness estimates and diversity indices. A second program, LIBSHUFF (34), was used to compare the similarities of bacterial clone libraries in Pits 91 and 101. The distance matrices for both programs were obtained using an algorithm located at the Greengenes website (4, 5).

**FISH.** Fluorescent in situ hybridization (FISH) was carried out as described previously (49) with minor modifications. Oligonucleotides were 5' end labeled with fluorescent dyes and included Eub338 labeled with Cy3, ARCH915 labeled with Cy5, and Pae997 with Bodipy FL. Details on these probes are available at probeBase (24). Cells were photographed with a Leica TCS/SP2 UV confocal microscope.

**Cultivation.** Culturable bacteria were isolated by serial dilutions of water suspensions of asphalt-soil mixtures on agar plates containing DSMZ medium

371 amended with 20% NaCl and 10% tryptic soy agar and M9 minimum medium. The plates were incubated at 28°C for 2 to 3 weeks, after which individual isolates were transferred and processed for sequencing of 16S rRNA gene sequences. Isolates were placed in glycerol medium and transferred to a -80°C freezer for long-term preservation. Bacterial isolates were tested on agar medium with 1% asphalt as a sole carbon source.

**Nucleotide sequence accession numbers.** All sequences were deposited in GenBank under accession numbers DQ001614 to DQ001623, DQ001626 to DQ001638, DQ001641, DQ001642, DQ001644, DQ001646, DQ001647, and EF157073 to EF157279 (for *Bacteria*), DQ192039 to DQ192061 (for isolates), DQ062817 to DQ062856 (for dioxygenase), AY860440 to AY860443 and AY939988 to AY940011 (for *Archaea*), and AY940013, AY940019 to AY940022, AY940024 to AY940026, AY940028, and AY940029, AY940032, and AY860446 to AY860448 (for *Pseudomonas stutzeri*).

## RESULTS

Detailed analyses of the two tar pits revealed differences in both the chemical composition and the microbial community composition of the asphalt samples. The asphalt-permeated soil from Pit 91 contained a greater concentration of petroleum hydrocarbons, was slightly acidic, and had a relatively low salinity and metal content (Table 1). In contrast, water suspensions of asphalt-permeated soil from Pit 101 were alkaline (pH 8.4) and contained a high concentration of salts and metals. The salinity of 1:1 asphalt water suspensions of Pit 101 was 4,610  $\mu\text{S cm}^{-1}$ , which was 100 times higher than that of Pit 91. Materials from both of the tar pits are impermeable to surface water from rainfall, and bacteria in this matrix are subject to water deficits and high salinity. Water contained in the asphalt is present in stratified water pockets and pore spaces that occur throughout the pit and are likely the main sites for microbial growth. The occurrence of microbial activity in the tar pits is visually evident from the continual evolution of methane bubbles at various locations in the pits where there is still viscous liquefied asphalt.

Here a total of 235 bacterial clones were sequenced to identify the predominant phylogenetic groups (see Fig. 2). The most striking difference in the microbial community composition of the two pits was the presence of halophilic *Archaea* in the highly saline Pit 101, which were not detected in Pit 91 (Fig. 1). Among these were 27 clones representing clusters of closely related species from two unclassified genera that were most similar to *Natronococcus* and *Natronobacterium* (Fig. 1). Of the 29 *Archaea* sequences that were obtained, only 2 were outside of the clusters representing the new genera.

In addition to *Archaea*, there were also differences in the distribution of bacterial phyla between the two pits. The predominant bacteria in both pits were *Gammaproteobacteria* (purple sulfur bacteria). Pit 91 contained 3 unclassified families (60 clones) in the order *Chromatiales*, none of which was found in Pit 101 (Fig. 2). Other families of the *Gammaproteobacteria*

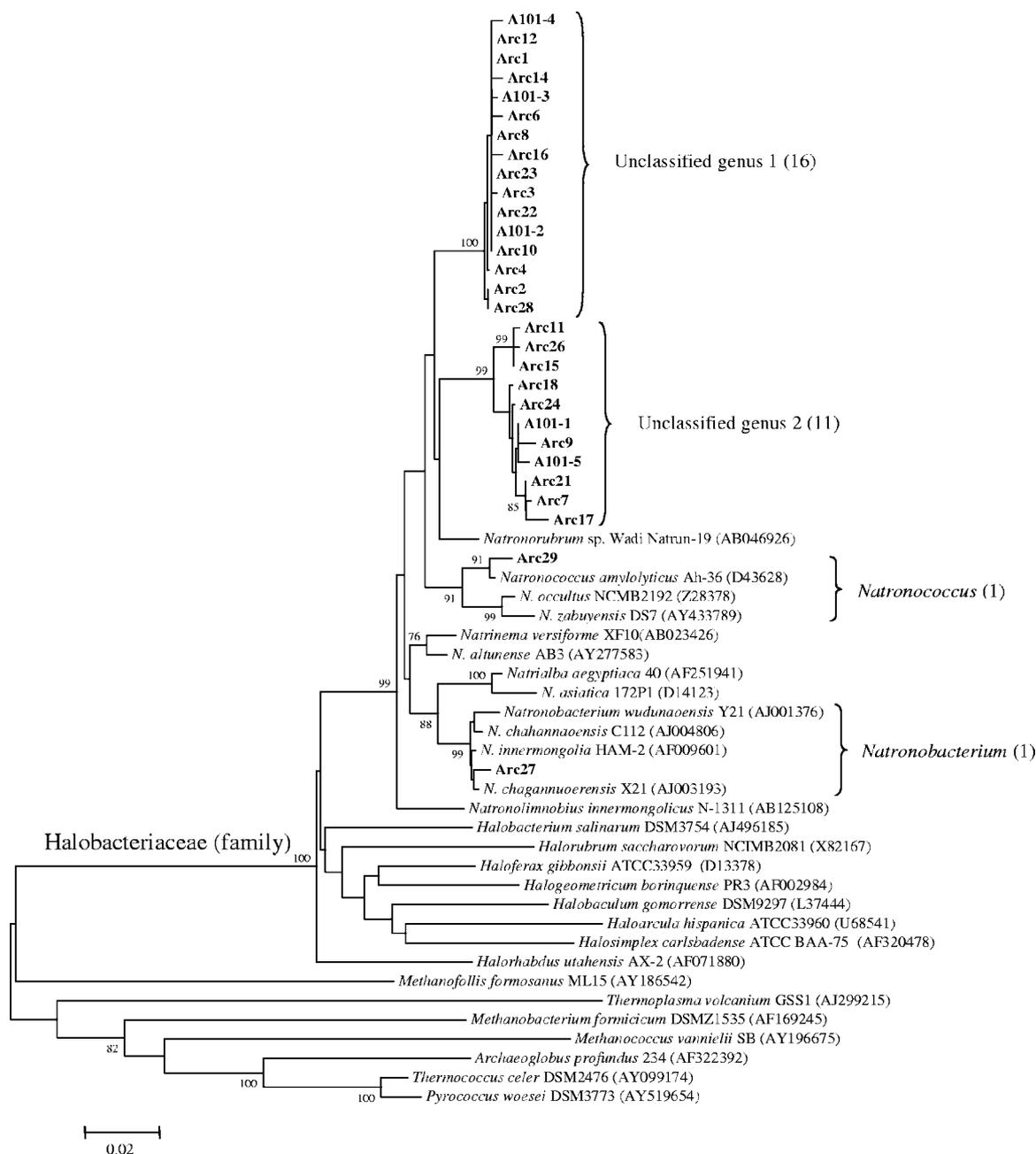


FIG. 1. Phylogeny of halophilic *Archaea* identified in an asphalt soil mixture from Pit 101 of the Rancho La Brea Tar Pits. Genera are indicated by vertical bars on the right of the tree, and values in parentheses are clone numbers. The robustness of the topology was estimated by bootstrap resampling of the neighbor-joining data. Bootstrap values greater than 75 are shown at branch points. Scale bar denotes 0.02 changes per nucleotide.

in Pit 91 included *Xanthomonadaceae* (5 clones) and *Pseudomonadaceae* (5 clones). (Fig. 2A and C).

Pit 101 contained a greater breadth of diversity of *Gamma*-*proteobacteria* and *Alphaproteobacteria* than Pit 91. This included two unclassified families and one new order with two more unclassified families (Fig. 2B). In addition to 16 clones representing the family *Rubrobacteraceae*, other unique taxa in Pit 101 included 5 additional phyla that were represented by 9 clones from the *Planctomycetes*, 6 clones of *Gemmatimonadetes*, 1 clone of BRC1, and 2 clones each of *Nitrospira* and *Verrucomicrobia* (Fig. 2D). Taxa that were common to both

Pit 91 and Pit 101 included species from *Alpha*-, *Beta*-, and *Gammaproteobacteria* and various representatives from the *Acidobacteria*, *Actinobacteria*, *Bacteroidetes*, and *Clostridia* (Fig. 2).

Sequences obtained using the *Pseudomonas*-selective primers included 14 clones of *Pseudomonas stutzeri*, which represented 8 new genomovars of this species (Fig. 3). All but 1 of the 14 clones was obtained from Pit 91, suggesting a differential distribution of *P. stutzeri* in the two tar pits (Fig. 3).

Culturable bacteria from both tar pits were represented by relatively few species. The 10% tryptic soy agar medium



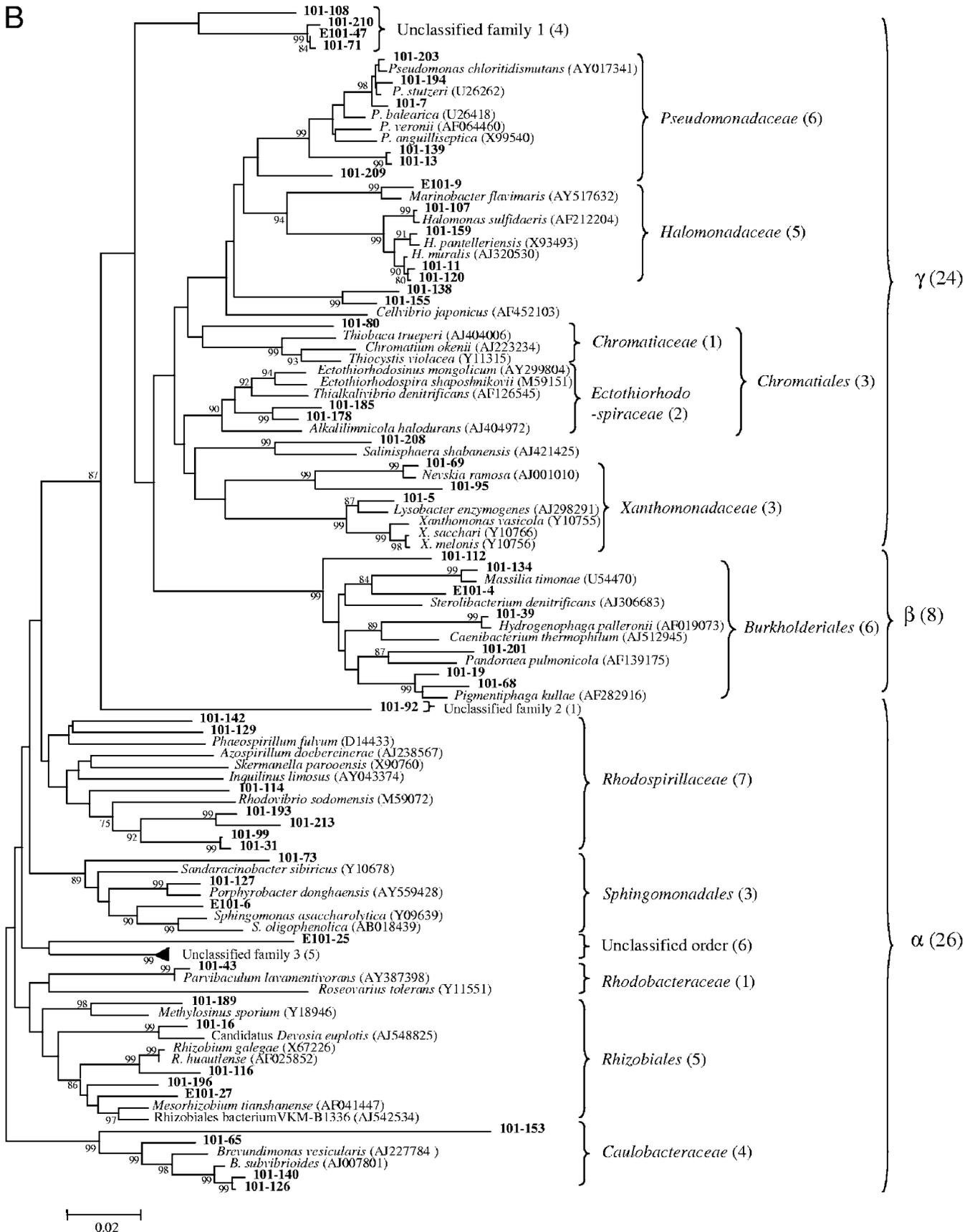
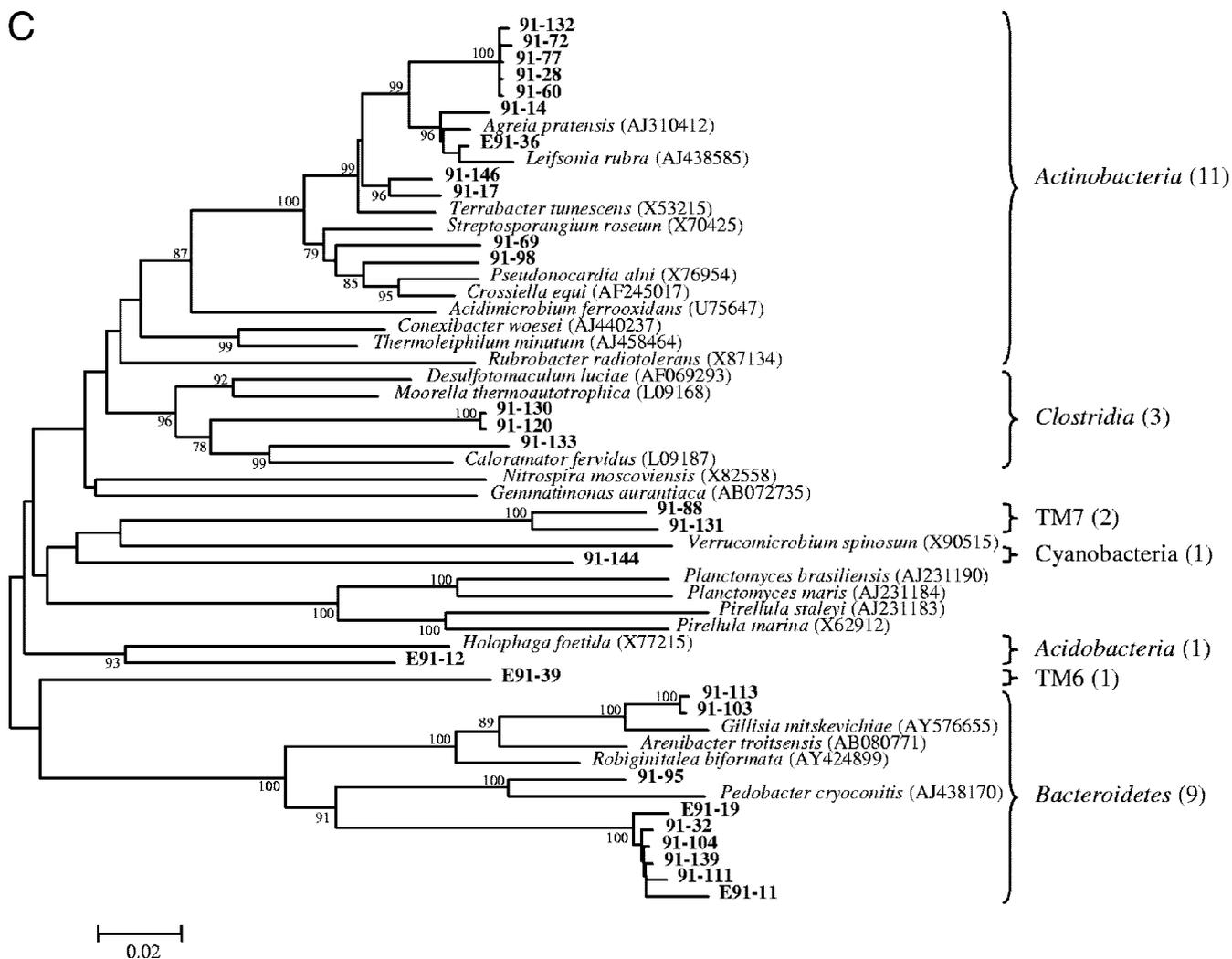


FIG. 2—Continued.



yielded five distinct isolates of *Pseudomonas* spp., eight isolates of *Bacillus* spp., and five isolates of *Citrobacter* spp. The 20% salt medium yielded three isolates of haloalkalophilic *Bacillus* spp. from Pit 101 (Fig. 4). All of the isolates obtained on the above media were further shown to grow on M9 medium with asphalt as the sole carbon source.

To study the physical distribution of *Archaea* and *Bacteria* in the asphalt from Pit 101, samples from the tar pit were microscopically examined using FISH with DNA probes targeted against *Bacteria*, *Archaea*, and *Pseudomonas* spp. (Fig. 5). Cells from each group were observed to occur in dense clusters of colonies along with randomly dispersed individual cells. The very close association of cells from different phylogenetic groups suggests that there may be consortia that cooperatively solubilize, degrade, and detoxify the complex substances contained in the asphalt. Such consortia also may be conducive for horizontal gene transfer of plasmids and DNA sequences that encode catabolic genes for use of petroleum hydrocarbons.

**Diversity analyses.** Rarefaction curves of Pit 91 and Pit 101 libraries showed significantly different numbers of operational taxonomic units (OTUs) at distance values of 0.03 correspond-

ing to the species level, 0.05 at the genus level, and 0.1 at the family/class level (Fig. 6A). Only 37 OTUs were obtained from the Pit 91 bacterial library, while 80 OTUs were obtained from the Pit 101 bacterial library. Indices of diversity were higher for the Pit 101 library than for the Pit 91 library. The Shannon and Simpson index values were 2.8 and 0.139 for the Pit 91 library, compared with 4.2 and 0.014 for the Pit 101 library. Richness ACE (abundance-based coverage estimator) and Chao1 (13) values were 64 and 58 in Pit 91 and 268 and 242 in Pit 101 (Table 2). There was a significant difference between the rarefaction curves for the Pit 91 and Pit 101 libraries based on 0%, 3%, 5% and 10% differences (Fig. 6A). LIBSHUFF comparisons of each of the libraries indicated that two communities, Pit 91 and Pit 101, were significantly different ( $P < 0.001$ ). At an evolutionary distance of 0.03, coverages of the libraries were 80% and 46% in Pit 91 and Pit 101, respectively (Fig. 6B).

**Bacterial dioxygenase sequences.** Taxonomic relationships for the dioxygenase sequences that were identified are shown in Fig. 7. The phylogenetic trees for these sequences included reference dioxygenases that were not found in the

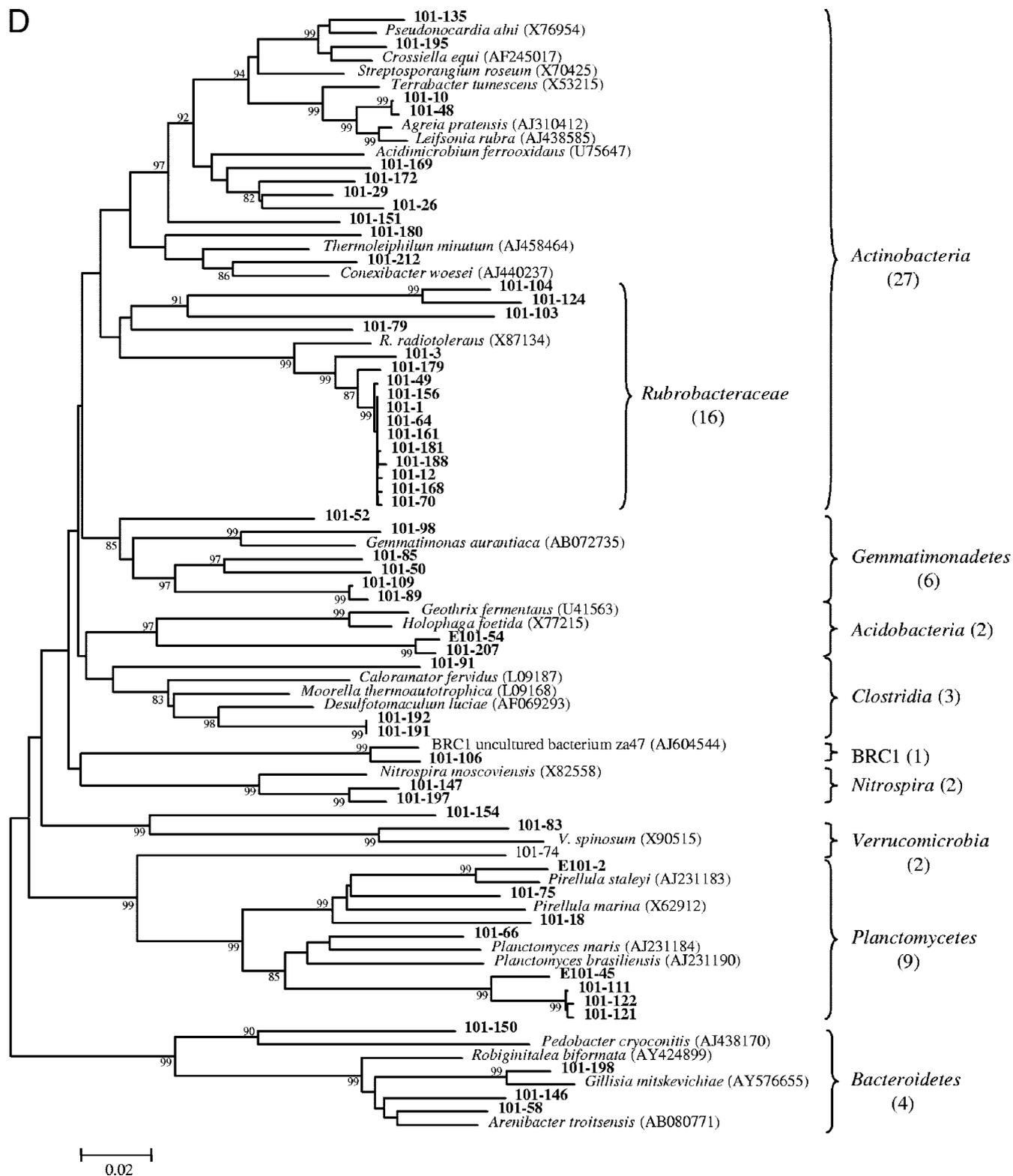


FIG. 2—Continued.

tar pits but that provide an indication of the similarities of the new sequences to those of previously described enzymes. Sequences from Pit 91 were predominantly associated with those of known proteobacterial biphenyl dioxygenases. This

cluster of biphenyl dioxygenases was comprised of 12 sequences from Pit 91 and 2 sequences from Pit 101. Overall similarities to known biphenyl dioxygenases ranged from 79 to 95%. Detailed analysis revealed at least four subclusters

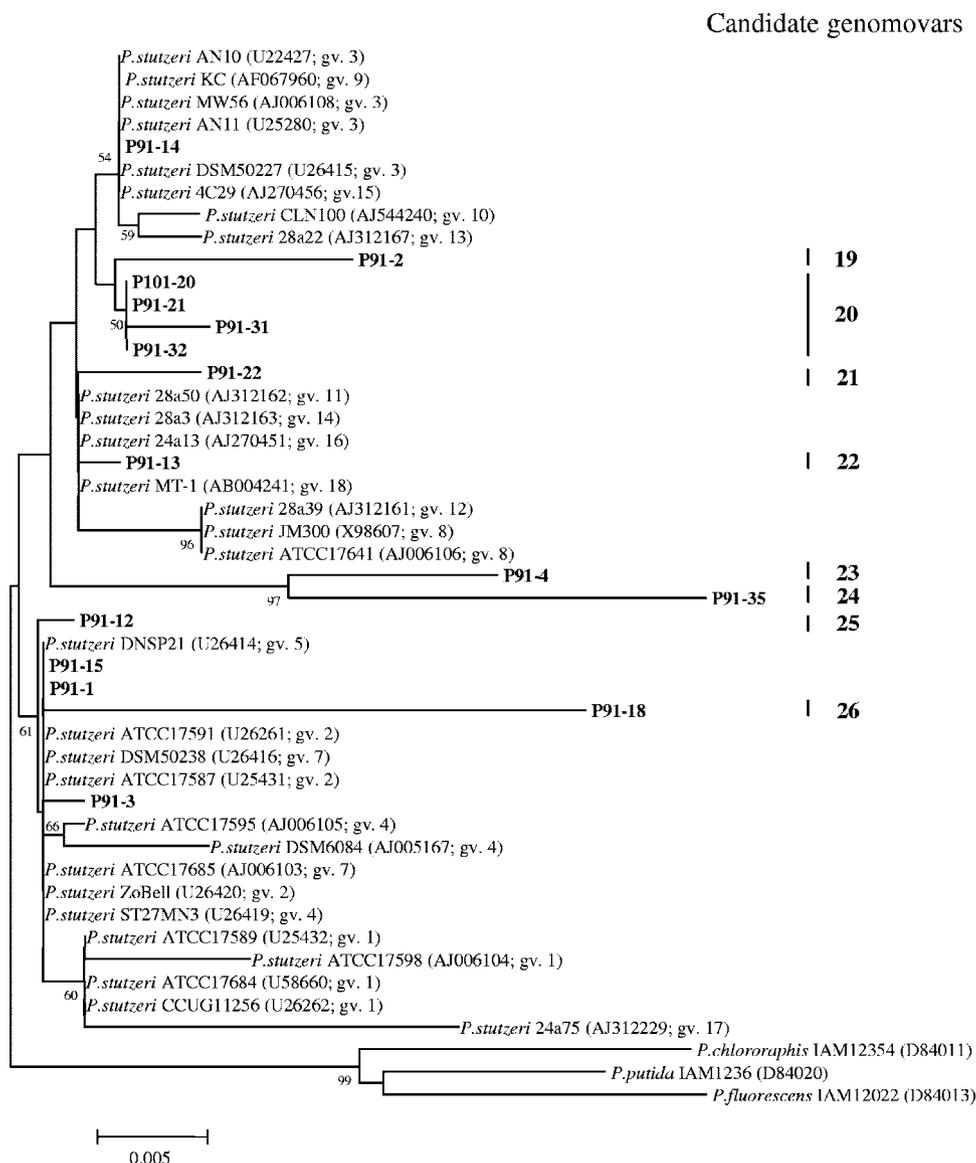


FIG. 3. Phylogeny of *Pseudomonas stutzeri*. 16S rRNA gene sequences identified in asphalt from Pits 91 and 101 of the Rancho La Brea Tar Pits using PCR primers for *Pseudomonas* spp. Candidate genomovars (gv.) are indicated by vertical bars on the right of the tree. The robustness of the topology was estimated by bootstrap resampling of the neighbor-joining data. Bootstrap values greater than 50 are shown at branch points. Scale bar denotes 0.005 changes per nucleotide.

within the biphenyl dioxygenase group. As with the microbial 16S rRNA genes, there were striking differences in the clusters of dioxygenases represented by the two sites. Ring-hydroxylating dioxygenases from Pit 101 were predominantly comprised of a new group of 18 closely related sequences that represent a novel, phylogenetically deep group of enzymes. This cluster was most closely associated with sequences encoding benzene and toluene dioxygenases but was sufficiently distant that it is not possible to infer whether these dioxygenases utilize these substances or instead transform other substrates. Two other new clusters that were discovered from sequences from Pit 91 were represented by one and two clones, respectively, and also appeared to represent deep branches of genes encoding unknown types of dioxygenases.

## DISCUSSION

Previous research has suggested that bacteria in deep subsurface oil reservoirs have inhabited those environments since the oil was formed (25). The origin of the bacteria in the natural asphalts at Rancho La Brea is unknown, but their presence could reflect recent entry from dust deposition on the surface, bacteria originating from the subsurface oil reservoir that seeped to the surface, or the progeny of soil bacteria that were embedded in the asphalt matrix as heavy oil seeped to the surface. Regardless of their origin, life in asphalt poses extreme conditions in which microbial growth is limited by the lack of air and water, the presence of highly recalcitrant carbon sources, and high concentrations of potentially toxic metals and chemicals. The selectivity of this environment would be

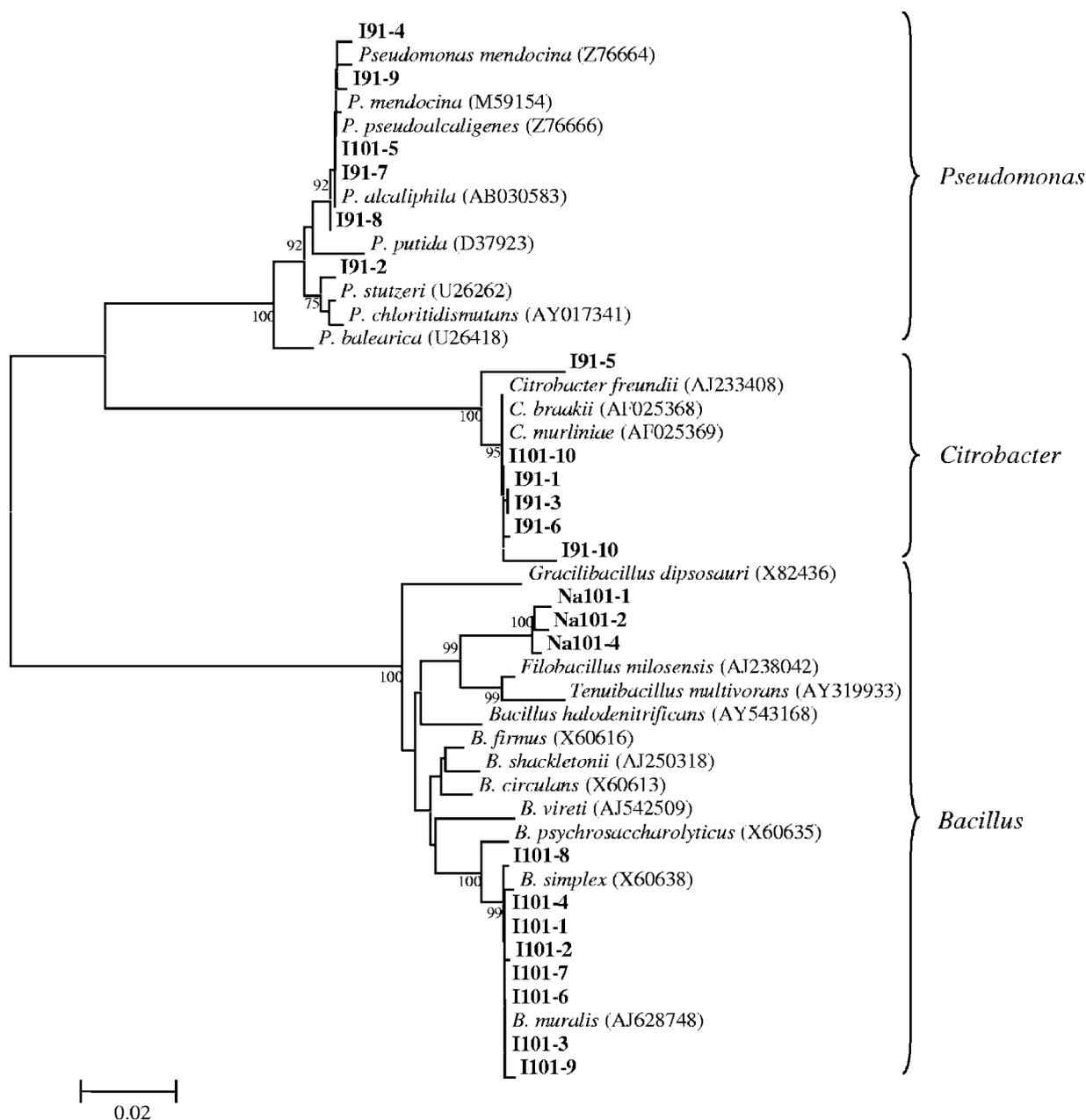


FIG. 4. Phylogeny of cultured bacteria isolated from the Rancho La Brea Tar Pits. Bootstrap values greater than 75 are shown. Bold letters indicate isolates were obtained from Pit 91 or Pit 101. Scale bar denotes 0.02 changes per nucleotide.

expected to require specialized adaptations. Here 235 clones were described, many of which appear to comprise new genera and families of *Proteobacteria* (Fig. 2). An analysis of two different pits differing in their chemical properties revealed very little overlap in diversity (Fig. 6A), indicating that site-specific differences in salinity and pH strongly influence selection within the asphalt communities.

The relatively simple community structures and low complexity of Pits 91 and 101 compared to soil suggest that this environment is highly selective. The phylogenetic trees within the *Proteobacteria* revealed considerable breadth in taxa at the levels of family and order but also manifested branches containing discrete clusters of related sequences. The occurrence

of many closely related species within the *Gammaproteobacteria* in both pits and the *Archaea* in Pit 101 suggested either an evolutionary radiation of species or an initial selection of closely related bacteria that share specific traits that enable them to survive in this environment.

Important questions arising from this research are the functional properties and adaptations of the taxa that inhabit the tar pits. It is very difficult to infer functionality at the level of phylum, and there is sparse information on many of the bacterial species that were found here. Among the predominant bacteria were 60 clones representing three unknown families from the *Gammaproteobacteria* in the order *Chromatiales*. This order has previously been characterized by two families, the

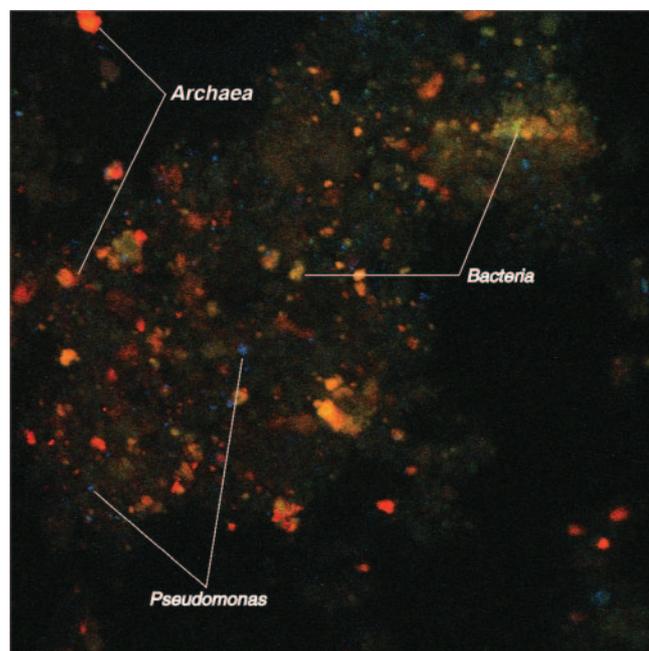


FIG. 5. Microorganisms associated with heavy oil-soil as revealed by FISH using a combination of dyes that stain different organisms. Cells of the *Archaea* are stained red, the *Bacteria* are yellow, and *Pseudomonas* (a genus of *Bacteria*) is blue. The microbial colonies can be seen as clumps of different-color cells that are growing together, which suggests coexistence in diverse consortia.

*Ectothiorhodospiraceae* and *Chromatiaceae*, both of which comprise phototrophic anaerobic bacteria that produce sulfur from hydrogen sulfide gas (14, 15). The former produce granules of sulfur on the outside of their cells, while the latter produce internal sulfur granules. The three new families discovered here are not likely to derive energy from photosynthesis given that they live in complete darkness within the asphalt-soil matrix. Nonetheless, an ability to utilize electrons from hydrogen sulfide is consistent with life in the tar pits, where hydrogen sulfide and methane are produced during anaerobic metabolism of hydrocarbons contained in the asphalt.

Another important cluster in Pit 101 was classified within the *Rubrobacteraceae*. There were 16 closely related sequences from this family. The *Rubrobacteraceae* are in the phylum *Actinobacteria* and have been previously reported to occur in high-ionizing-radiation environments and in Australian desert soils (11). *Rubrobacter* strains have received considerable attention as being among the bacteria most resistant to ionizing radiation (7). Whether the bacteria from Pit 101 are radiation resistant is unknown, but it can be speculated that this trait could be of importance as an adaptation to protection from DNA damage in mixtures of heterocyclic aromatic hydrocarbons which are potent mutagens (48).

The occurrence of *Pseudomonas stutzeri* sequences, which included eight new genomovars, is consistent with prior reports on the distribution of this species. *P. stutzeri* is a well-known petroleum hydrocarbon degrader (8) and appears to be a cosmopolitan species that is readily isolated from various petroleum-contaminated environments (20, 29). Previously 17 genomovars have been described (20); this research has added

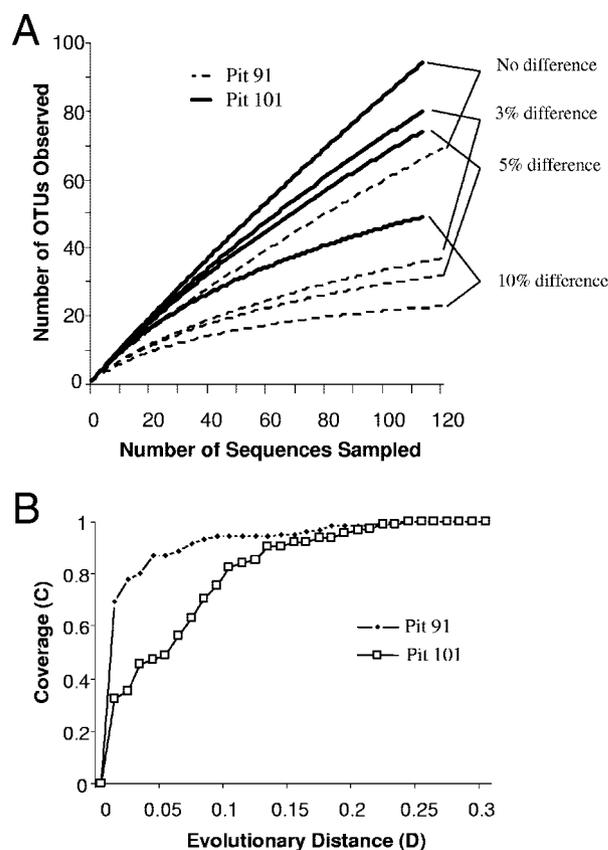


FIG. 6. Rarefaction curves of observed OTU richness using DOTUR (A) and coverage calculated using LIBSHUFF (B) from the two bacterial 16S rRNA gene libraries in Pit 91 and Pit 101 of Rancho La Brea.

another 8 candidate genomovars. As noted for several taxa in the communities from the tar pits, the existence of closely related sequences of *P. stutzeri* could be explained either by selection for bacteria that shared essential characteristics that allow them to survive in the asphalt or as an evolutionary radiation of ecotypes. With the exception of *Pseudomonas* spp., all of the identified taxa were dissimilar to those reported earlier for two studies investigating a high-temperature oil reservoir, which are the only other studies in the literature reporting a bacterial survey of a natural petroleum habitat using culture-independent methods (3, 26).

The discovery of many halophilic *Archaea* sequences in Pit 101 is particularly intriguing and provides an opportunity for future studies on the role of *Archaea* in petroleum hydrocarbon degradation. Based on studies conducted with enrichment cultures of bacterial petroleum degraders, biodegradation typ-

TABLE 2. Richness and diversity estimations in 16S rRNA gene libraries from Pit 91 and Pit 101 of La Brea Tar Pits

Library	Richness					Diversity index value		Coverage
	No. of OTUs	ACE value	Boot value	Chao1 value	Jack value	Shannon	Simpson	
Pit 91	37	64	46	58	58	2.8	0.139	0.8
Pit 101	80	268	105	242	265	4.2	0.014	0.45

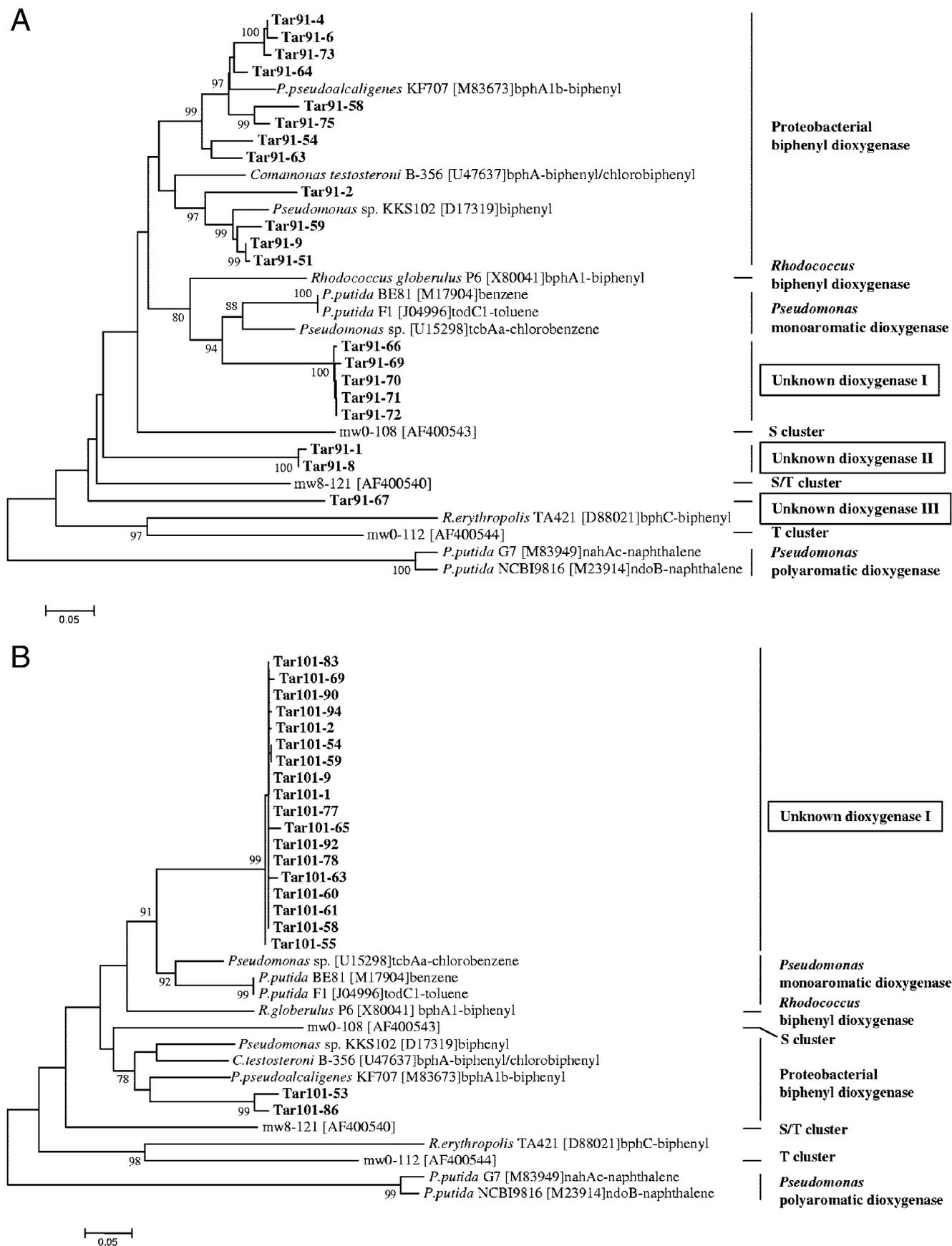


FIG. 7. Phylogeny of aromatic ring hydroxylating dioxygenases identified from Rancho La Brea Tar Pits. (A) Pit 91; (B) Pit 101. Bootstrap values greater than 75 are shown. Bold letters denote clones sampled from the asphalt-soil mixtures. Sequences shown in regular typeface are known dioxygenases from GenBank used here for classification. Enzyme classes are indicated by vertical bars at the right of the trees. Scale bar indicates 0.05 changes per nucleotide.

ically involves consortia in which the species composition of the degrader community is strongly influenced by salinity (18, 44). Whether this selection also occurs with *Archaea* is not known. The *Archaea* have been reported to occur in crude oil sludge but not in crude oil samples in oil stockpiles in Japan (40) or in petroleum reservoirs in California (26). Various *Halobacteria* isolated from soil and sediments that are capable of degrading hydrocarbons under saline conditions have been described. These include species tentatively identified as *Halobacterium* (2), *Haloferax* (51), and *Haloarcula* (40). A prior report on *Archaea* that can degrade aromatic hydrocarbons under anaerobic conditions using Fe(III) as an electron acceptor was published in 2001 (39). Although not yet cultivable, genes from these bacteria potentially could be used for improving culturable hydrocarbon degraders used for bioaugmentation or could serve as a source of catabolic genes that could be seeded into the environment on plasmids to facilitate adaptation of indigenous strains for degradation of recalcitrant hydrocarbons (38). It will also be important to determine whether the large unknown cluster of ring-hydroxylating dioxygenases identified in Pit 101 are carried by the *Archaea*, which is suggested by their cooccurrence in this particular site.

As expected from prior experience with environmental samples, relatively few isolates were obtained using culture-based methods. The culturable bacteria included strains of *Pseudomonas* spp., *Citrobacter* spp., and *Bacillus* spp. that were similar to known strains from oil-contaminated environmental samples (Table 2). The ability to culture these strains, especially isolates of *Pseudomonas*, provides opportunities for full genome analysis and examination of genetic exchange that may have occurred within this group of bacteria, as well as determination of plasmid-borne genes or catabolic pathways.

From an applied perspective, the most practical aspect of this research may be the confirmation of heavy oil seeps and natural asphalts as sources of novel genes for biodegradation of petroleum hydrocarbons. Here we focused on genes encoding aromatic-ring-hydroxylating dioxygenases that are important for degradation of BTEX and aromatic chemicals, such as polychlorinated biphenyls (PCBs) and polycyclic aromatic hydrocarbons, that are common environmental pollutants. The maintenance of genes encoding dioxygenases by bacteria that have undergone selection for life in an anaerobic system is somewhat of a paradox but is common for known oil-degrading bacteria. Previously described toluene-degrading bacteria placed under anaerobic conditions have been shown to first use dioxygenases to degrade toluene until all of the oxygen is consumed, after which the cells switch to a benzylsuccinate pathway that is coupled to denitrification (33). Among the sequences that were obtained were a large number that were similar to those encoding known biphenyl dioxygenases that function for degradation of PCBs. Subtle variations in key regions of these genes can lead to large differences in substrate range and specificity for different PCB congeners. In the future, enrichment culture methods may be used to identify still other enzymes that can target specific substrates.

Relatively little is known yet about anaerobic petroleum hydrocarbon degradation, although there has been steady progress in this field (1, 23, 30, 50). Tentative mechanisms that function for anaerobic degradation of alkylbenzenes and non-aromatic hydrocarbons are proposed to involve hydrolases and

carboxylases and are coupled to sulfate or nitrate reduction (35, 46). In our survey, anaerobic bacteria that were identified included members of *Gammaproteobacteria* (60 clones of purple sulfur bacteria), *Bacteroidetes*, *Clostridia*, and *Acidobacteria*, none of which have been studied with respect to their possible contributions to anaerobic degradation of petroleum hydrocarbons. Homologous genes for the benzylsuccinate synthase (*bss*), which is the key enzyme for anaerobic toluene degradation, have been cloned (33) and may provide an entry point for future studies on the relevance of this pathway in the tar pit bacteria. In addition to direct catabolism of hydrocarbons, anaerobic bacteria may also contribute to hydrocarbon degradation by syntrophy, in which metabolically linked consortia function to consume fatty acids and degradation products of hydrocarbons to generate methane (46).

Looking toward future research on the Rancho La Brea microorganisms, the discovery of closely related bacterial clusters and genes encoding new dioxygenases is of particular interest for understanding the evolutionary biology of bacteria during adaptation to the extreme environment posed by life in asphalt. Detailed studies on efficient regulator-promoter pairs are now being conducted to understand and design improved operons for xenobiotic degrading bacteria (6). New approaches using high-throughput DNA sequencing will be the next step for obtaining insight into the function and diversity of oil-inhabiting bacteria and the catabolic pathways for degradation of petroleum hydrocarbons.

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