

## Phylogeny and Molecular Identification of Vibrios on the Basis of Multilocus Sequence Analysis

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We analyzed the usefulness of *rpoA*, *recA*, and *pyrH* gene sequences for the identification of vibrios. We sequenced fragments of these loci from a collection of 208 representative strains, including 192 well-documented *Vibrionaceae* strains and 16 presumptive *Vibrio* isolates associated with coral bleaching. In order to determine the intraspecific variation among the three loci, we included several representative strains per species. The phylogenetic trees constructed with the different genetic loci were roughly in agreement with former polyphasic taxonomic studies, including the 16S rRNA-based phylogeny of vibrios. The families *Vibrionaceae*, *Photobacteriaceae*, *Enterovibrionaceae*, and *Salinivibrionaceae* were all differentiated on the basis of each genetic locus. Each species clearly formed separated clusters with at least 98, 94, and 94% *rpoA*, *recA*, and *pyrH* gene sequence similarity, respectively. The genus *Vibrio* was heterogeneous and polyphyletic, with *Vibrio fischeri*, *V. logei*, and *V. wodanis* grouping closer to the *Photobacterium* genus. *V. haliotocoli*-, *V. harveyi*-, *V. splendidus*-, and *V. tubiashii*-related species formed groups within the genus *Vibrio*. Overall, the three genetic loci were more discriminatory among species than were 16S rRNA sequences. In some cases, e.g., within the *V. splendidus* and *V. tubiashii* group, *rpoA* gene sequences were slightly less discriminatory than *recA* and *pyrH* sequences. In these cases, the combination of several loci will yield the most robust identification. We can conclude that strains of the same species will have at least 98, 94, and 94% *rpoA*, *recA*, and *pyrH* gene sequence similarity, respectively.

Vibrios are gram-negative, usually motile rods, are mesophilic and chemoorganotrophic, and have a facultatively fermentative metabolism (5). They are generally able to grow on marine agar and on the selective medium thiosulfate-citrate-bile salt-sucrose agar and are mostly oxidase positive. Vibrios belong to the *Gammaproteobacteria* according to 16S rRNA gene sequence analysis. These bacteria are found abundantly in aquatic habitats and in association with eukaryotes. Associations established by vibrios range from mutualistic, e.g., *Vibrio fischeri*-bobtail squid (26), to pathogenic, e.g., *V. cholerae*-humans (45). Probiotic *Vibrio* strains for fish and shellfish have also been documented (44).

The current family *Vibrionaceae* comprises the genera *Enterovibrio* (2 species), *Grimontia* (1 species), *Photobacterium* (7 species), *Salinivibrio* (1 species), and *Vibrio* (64 species). The novel species *Photobacterium rosenbergii* and *Enterovibrio corallii* have recently been proposed to encompass isolates associated with coral bleaching (41). Several new *Vibrio* species, mainly in the phylogenetic neighborhood of *V. harveyi*, *V. haliotocoli*, *V. splendidus*, *V. tubiashii*, and *V. fluvialis*, have been described in the last few years, with *V. neonatus*, *V. ezurae* (28), and *V. ponticus* (22) being the most recent ones. *V. harveyi*, *V.*

*splendidus*, and *V. tubiashii* are frequently associated with disease in different species of fish and shellfish worldwide, while the *V. haliotocoli* group comprises species that are potentially mutualist to abalones (28). These organisms may be promising probiotics for abalone rearing.

Accurate identification of vibrios at the family and genus levels is obtained by 16S rRNA gene sequencing, whereas identification at the species and strain levels requires the application of genomic analyses, including DNA-DNA hybridization, repetitive extragenic palindromic PCR, and amplified fragment length polymorphism (AFLP) analysis (40). These techniques are essential for reliable species identification, because several vibrios have nearly identical 16S rRNA sequences and similar phenotypic features (10). Unfortunately, their use is restricted to a few reference laboratories. Interlaboratory comparisons of fingerprint patterns are difficult. The sequencing of housekeeping genes is emerging as an alternative to overcome this problem. In addition, this type of data may improve the current pragmatic definition of bacterial species (28a). In silico whole-genome analysis applied to the *Gammaproteobacteria* identified a set of 203 genes that are most valuable for inferring bacterial phylogeny (18a). According to Gevers et al. (8), 135 of these genes are not conserved outside the *Gammaproteobacteria* and thus may not be appropriate for phylogenetic studies on a broader taxonomic scale.

Different loci, e.g., 23S rRNA (21), *gapA* (23), *gyrB* (20), *hsp60* (18), and *recA* (30), have been used for phylogenetic studies and the identification of *Vibrionaceae* species. So far,

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these genes (except for *recA*) have only been examined in a very limited number of species and strains. Thompson et al. (30) analyzed the *recA* sequences of most vibrios, but mainly using type strains, making it difficult to draw conclusions about the use of this gene as an identification marker. Alternative phylogenetic markers should fulfill several criteria, as put forward by Zeigler (46): (i) the genes must be widely distributed among genomes, (ii) the genes must be present as a single copy within a given genome, (iii) the individual gene sequences must be long enough to contain sufficient information but short enough to allow sequencing in a convenient way (900 to 2,250 nucleotides [nt]), and (iv) the sequences must predict whole-genome relationships with acceptable precision and accuracy that correlate well with the 16S rRNA data and with whole-genome similarities measured by, e.g., DNA-DNA hybridization. A combination of in silico analyses and recent experimental studies of different bacteria, including *Bacillus*, *Proteobacteria*, lactic acid bacteria, *Mycobacterium*, and *Mycoplasma*, suggested that the RNA polymerase alpha subunit gene (*rpoA*), *recA*, and the uridylylate kinase gene (*pyrH*) fulfill these requisites and could therefore be used for identification purposes (8, 18a, 46).

For the present study, we analyzed the *rpoA*, *recA*, and *pyrH* gene sequences of 192 well-documented *Vibrionaceae* strains comprising all currently known species (except for *Photobacterium profundum*, *V. agarivorans*, *V. calviensis*, *V. ruber*, *V. ponticus*, and *V. salmonicida*). We also included 16 presumptive *Vibrio* isolates associated with healthy and bleached corals. In order to determine the intraspecies variation among the three loci, we included at least three representative strains each of 37 *Vibrionaceae* species. The aim of this study was to evaluate the application of different genetic loci as phylogenetic and identification markers, both individually and as concatenated elements. The multigene-based phylogeny roughly confirmed the 16S rRNA gene-based grouping obtained in previous studies and revealed new interesting relationships among different *Vibrio* species.

#### MATERIALS AND METHODS

The strains used for this study are listed in Table 1. All strains included in this study are deposited in the BCCM/LMG Bacteria Collection or the Research Collection at Ghent University (Ghent, Belgium). A detailed list of these strains can be found elsewhere (31, 40). This study also included fresh isolates associated with bleached and healthy corals of different species collected in 2002 in Australia and the United States. Isolates R-21409, R-21422, and R-21431 originated from different bleached *Pachyseris speciosa* colonies (Magnetic Island, Australia). R-21410 originated from a healthy *Montipora capitata* sample, and isolates R-21416 and R-21432 originated from a healthy *P. speciosa* sample, both in Kaneohe Bay (United States). R-21415, R-21419, and R-21433 were isolated from healthy *Merulina ampliata* samples (Magnetic Island, Australia), and R-21413 was isolated from bleached *M. ampliata* coral (Magnetic Island, Australia). R-21427 and R-21434 originated from bleached *Acropora millepora* coral (Davies Reef, Australia). R-21426, R-21435, and R-21439 were isolated from healthy *A. millepora* (Davies Reef, Australia), *Pocillopora damicornis* (Davies Reef, Australia), and *Barbattoia amicornum* (Magnetic Island, Australia) samples, respectively. R-23286 was isolated from a bleached *Montipora* sp. sample (Magnetic Island, Australia) in 2003.

Bacterial genomic DNAs were extracted according to the methodology described by Pitcher et al. (24). PCRs were performed essentially as described previously (30). The sequences of the primers used for amplification and sequencing are listed in Table 2. These primers were designed using 17 gene sequences of each locus of *Vibrio cholerae* (n16961\_o1), *V. parahaemolyticus* (o3k6\_rimd2210633), *V. vulnificus* (cmcp6), *Escherichia coli* (cft073, o157h7\_edl933, o157h7\_rimd0509952, and k12\_mg1655), *Shigella flexneri*

(2a\_2457t and 2a\_301), *Salmonella enterica* (ct18 and ty2\_typhi 3), *S. enterica* serovar Typhimurium (lt2sgsc1412\_atcc700720), *Yersinia pestis* (co92 and kim), *Haemophilus influenzae* (rd), *Pasteurella multocida* (pm70), and *Shewanella oneidensis* (mr1), which originated from publicly available data from whole-genome sequencing projects. All of the primers specifically amplified the target fragments of all currently known strains of vibrios.

PCR mixtures were composed of 29.5  $\mu$ l sterile MilliQ water, 5.0  $\mu$ l PCR buffer (10 $\times$ ), 5.0  $\mu$ l deoxynucleoside triphosphates (2 mM each), 2.5  $\mu$ l forward primer *rpoA*-01-F (10  $\mu$ M), 2.5  $\mu$ l reverse primer *rpoA*-03-R (10  $\mu$ M), 0.5  $\mu$ l AmpliTaq DNA polymerase (1 U/ $\mu$ l), and 5.0  $\mu$ l template DNA (0.01  $\mu$ g/ $\mu$ l). PCRs were performed using a GeneAmp PCR System 9600 thermocycler (Applied Biosystems). The thermal program consisted of (i) 5 min at 95°C; (ii) 3 cycles of 1 min at 95°C, 2 min 15 s at 55°C, and 1 min 15 s at 72°C; (iii) 30 cycles of 35 s at 95°C, 1 min 15 s at 55°C, and 1 min 15 s at 72°C; and (iv) a final 7 min at 72°C. PCR products with the expected size and intensity were purified using the Nucleofast 96 PCR cleanup system (Macherey-Nagel, Germany). Purified PCR products were eluted in 30 to 200  $\mu$ l sterile MilliQ water. Subsequently, 3.0  $\mu$ l of purified PCR product was mixed with 1.0  $\mu$ l ABI Prism Big Dye Terminator ready reaction mix, version 3.1 (Applied Biosystems), 3.0  $\mu$ l sequencing primer (4  $\mu$ M), 1.5  $\mu$ l dilution buffer (5 $\times$ ), and 1.5  $\mu$ l MilliQ water. The thermal program consisted of 30 cycles of 15 s at 96°C, 1 s at 35°C, and 4 min at 60°C. Sequencing products were purified using a Montage SEQ<sub>96</sub> sequencing reaction cleanup kit (Millipore). Purified sequencing reactions were eluted in 20  $\mu$ l of injection solution and mixed with 20  $\mu$ l of deionized formamide. Subsequently, separation of the DNA fragments was obtained in an ABI PRISM 3100 genetic analyzer (Applied Biosystems). The time and voltage of sample injection were 20 s and 1.25 kV. Each run was performed at 50°C for 6,500 s at 0.1 mA and 12.2 kV. Raw sequence data were transferred to the Gene Builder module within Kodon package 2.03 (Applied Maths, Belgium), where consensus sequences were determined using the two to four reads. Consensus sequences were imported into BioNumerics 3.0 software (Applied Maths, Belgium), where a similarity matrix and phylogenetic trees were created based on the maximum parsimony and neighbor joining methods (27). Splits tree decomposition analysis was done using software available on the Internet (<http://bibiserv.techfak.uni-bielefeld.de/splits/>) (13), while the GC content, ratio of mean synonymous substitutions per synonymous site to mean nonsynonymous substitutions per nonsynonymous site ( $d_s/d_N$ ), and Sawyer's test were calculated using the software package START obtained from <http://pubmlst.org/software/analysis/start/> (15).

#### RESULTS AND DISCUSSION

We sequenced fragments of the *rpoA* (931 nt), *recA* (613 to 783 nt), and *pyrH* (443 nt) genes of 192 vibrios corresponding to 60 to 93% of the coding regions of these genes (EMBL accession no. AJ842347 to AJ842743). The GC contents of *rpoA* (46%  $\pm$  0.9%), *recA* (46.6%  $\pm$  1.9%), and *pyrH* (48.7%  $\pm$  0.9%) are within the average for the total genomes of vibrios (34–39). We compared the *rpoA*, *recA*, *pyrH*, and 16S rRNA pairwise similarities using Pearson's product-moment correlation coefficient. A significant correlation was obtained between 16S rRNA and the three loci ( $R = 0.81$  for *rpoA* and 0.68 for *recA* and *pyrH*). The *rpoA* and 16S rRNA data had a linear relationship, whereas the *pyrH* and *recA* data were best fitted with a polynomial regression of the second degree. These significant correlations show that *rpoA*, *recA*, and *pyrH* are indeed phylogenetic markers in vibrios. The gene sequences of these three loci have high proportions of synonymous mutations ( $d_s/d_N = 29$  for *rpoA* and *recA* and 39 for *pyrH*), suggesting that there is selection against amino acid changes in these genes (6).

The families *Vibrionaceae*, *Photobacteriaceae*, *Enterovibrionaceae*, and *Salinivibrionaceae* were clearly separated from one another on the basis of the four loci (Fig. 1). The five currently known genera of vibrios were all differentiated on the basis of the gene sequences (Fig. 1). *Enterovibrio* strains had 95.4 to 100%, 88.6 to 100%, and 85.6 to 99.8% *rpoA*, *recA*, and *pyrH* gene sequence similarity, respectively. *Grimontia hollisae* and

TABLE 1. Gene sequence variation within species

Species (strains)	% Sequence variation		
	rpoA	recA	pyrH
<i>Enterovibrio norvegicus</i> (LMG 19839T, LMG 19840, LMG 19842, R-3717)	0.2	0.4	1.0
<i>E. corallii</i> (LMG 22228T)			
<i>Grimontia hollisae</i> (LMG 17719T, LMG 21416, LMG 21538)	0.1	0.0	0.1
<i>Photobacterium angustum</i> (LMG 8455T)			
<i>P. damsela</i> subsp. <i>damsela</i> (LMG 7892T, LMG 10940, LMG 19445)	0.1	0.1	1.0
<i>P. rosenbergii</i> (LMG 22223T, LMG 22224, LMG 22225, LMG 22226, LMG 22227, R-21419)	0.4	5.5	3.0
<i>P. leiognathi</i> (LMG 4228T, LMG 10944, LMG 11221)	0.2	1.0	2.0
<i>P. phosphoreum</i> (LMG 4233T)			
<i>Salinivibrio costicola</i> subsp. <i>costicola</i> (LMG 11651T)			
<i>Vibrio aerogenes</i> (LMG 19650T)			
<i>V. aestuarianus</i> (LMG 7909T)			
<i>V. alginolyticus</i> (LMG 4409T, LMG 2174, LMG 4407, LMG 19993, R-14876)	0.6	1.0	1.5
<i>V. anguillarum</i> (LMG 4437T, LMG 10861)	0.2		0.5
<i>V. brasiliensis</i> (LMG 20546T, LMG 20010)	0.0		0.1
<i>V. campbellii</i> (LMG 11216T, LMG 11256, LMG 16835, LMG 20369, R-14899, R-14902, R-21413, R-21427, R-21434)	0.4	3.0	2.0
<i>V. chagasii</i> (LMG 21353T, LMG 13219, LMG 13237)	0.1		2.0
<i>V. cholerae</i> (LMG 21698T, R-18244, R-18258, R-18297, R-18303, R-20544, R-20545, R-20546, R-20548)	0.1	6.0	5.0
<i>V. cincinnatiensis</i> (LMG 7891T)			
<i>V. corallilyticus</i> (LMG 20984T, LMG 10953, LMG 19270, LMG 20538, LMG 20548, LMG 21350, R-14978, R-14968, R-21432, R-23286)	0.9	4.0	3.0
<i>V. cyclitrophicus</i> (LMG 21359T, LMG 20001, R-14874)	0.0	3.0	
<i>V. diabolicus</i> (LMG 19805T)			
<i>V. diazotrophicus</i> (LMG 7893T, LMG 11217, LMG 13218, LMG 20033, R-3706)	0.5	1.6	1.5
<i>V. ezurae</i> (LMG 19970T, LMG 19979, R-15766)	0.0	0.4	0.5
<i>V. fischeri</i> (LMG 4414T, LMG 11653)	0.2	4.0	2.0
<i>V. fluvialis</i> (LMG 7894T, LMG 11654, R-15090)	0.2	1.8	1.0
<i>V. fortis</i> (LMG 21557T, LMG 20547, R-15037, R-21409, R-21431)	0.3	6.0	6.0
<i>V. furnissii</i> (LMG 7910T, LMG 11655, LMG 11656, LMG 11758)	0.2	0.7	1.5
<i>V. gallicus</i> (LMG 21330T)			
<i>V. gazogenes</i> (LMG 19540T)			
<i>V. halitocoli</i> (LMG 18542T, LMG 19700, LMG 19963, LMG 19975)	0.1	1.5	2.0
<i>V. harveyi</i> (LMG 4044T, LMG 7890, LMG 11226, LMG 19643, LMG 19714, LMG 20977, R-14947, LMG 11659, LMG 20370, R-14913, R-21410, R-21426, R-21435, R-21439)	2.0	4.2	1.5
<i>V. hepatarius</i> (LMG 20362T)			
<i>V. hispanicus</i> (LMG 13240T)			
<i>V. ichthyenteri</i> (LMG 19664T, R-3774, R-3789, R-3911)	0.1	1.0	0.1
<i>V. kanaloaei</i> (LMG 20539T, R-15010)	0.0	0.2	0.1
<i>V. lentus</i> (LMG 21034T, R-3884, R-3895, R-3912)	0.2	2.6	1.5
<i>V. logei</i> (LMG 19806T)			
<i>V. mediterranei</i> (LMG 11258T, LMG 11663, LMG 16836, LMG 19703T, R-14988, R-14989, R-14990, R-21415)	0.2	0.4	4.0
<i>V. metschnikovii</i> (LMG 11664T, LMG 4416, LMG 4426, R-22290)	0.0	3.2	3.5
<i>V. mimicus</i> (LMG 7896T, R-20564, R-20565, R-20568)	0.1	2.5	3.0
<i>V. mytili</i> (LMG 19157T)			
<i>V. natriegens</i> (LMG 10935T)			
<i>V. navarrensis</i> (LMG 15976T)			
<i>V. neonatus</i> (LMG 19972T, LMG 19976, LMG 19978)	0.1	1.8	2.0
<i>V. neptunius</i> (LMG 20536T, LMG 20613, R-1575)	0.2	0.6	1.0
<i>V. nereis</i> (LMG 3895T)			
<i>V. nigripulchritudo</i> (LMG 3896T)			
<i>V. ordalii</i> (LMG 13544T, LMG 10951, R-15101, R-15107)	0.0	0.7	2.0
<i>V. orientalis</i> (LMG 7897T)			
<i>V. pacinii</i> (LMG 19999T, LMG 13245, R-15016)	0.1	1.1	1.5
<i>V. parahaemolyticus</i> (LMG 2850T, LMG 11670, RIMD 2210633)	0.3	6.0	6.0
<i>V. pectenica</i> (LMG 19642T, LMG 20549, LMG 20550)	0.0	0.0	1.5
<i>V. pelagius</i> (LMG 3897T, LMG 19995)	0.1		0.5
<i>V. penaeicida</i> (LMG 19663T)			
<i>V. pomeroyi</i> (LMG 20537T, R-14805)	0.2	2.2	1.5
<i>V. proteolyticus</i> (LMG 3772T, R-15065)	0.0	0.0	0.5
<i>V. rotiferianus</i> (LMG 21460T, R-14935, R-21416, R-21422, R-21433)	0.0	0.6	0.5
<i>V. rumoiensis</i> (LMG 20038T, LMG 20039)	0.0	0.3	0.5
<i>V. scopthalmi</i> (LMG 19158T, LMG 20023, R-15029)	0.0	0.8	
<i>V. splendidus</i> (LMG 19031T, R-14789, LMG 16748, LMG 16751, LMG 16752)	0.8	6.0	1.0
<i>V. superstes</i> (LMG 21323T)			
<i>V. tapetis</i> (LMG 19706T, LMG 19704, LMG 19705)	0.0	0.0	1.0
<i>V. tasmaniensis</i> (LMG 20012T, R-14842, R-14846)	0.0	0.6	1.0
<i>V. tubiashii</i> (LMG 10936T, LMG 16851, R-14825)	1.0	3.3	3.7
<i>V. vulnificus</i> (LMG 13545T, R-15063, CMCP6)	0.3	1.3	2.4
<i>V. wodanis</i> (K16, K26)	0.1	0.3	1.1
<i>V. xuii</i> (LMG 21346T, LMG 20011, R-15053)	0.1	1.7	1.2

TABLE 2. Amplification and sequencing primers for *rpoA*, *recA*, and *pyrH*

Gene product	Primer	Sequence (5'→3')	Position
Uridylate kinase ( <i>pyrH</i> ; 750 nt)	pyrH-02-R	GTRAABGCNGMYARRTCCA	599
	pyrH-04-F	ATGASNACBAAAYCCWAAACC	1
RNA polymerase alpha chain ( <i>rpoA</i> ; 1,000 nt)	rpoA-01-F	ATGCAGGGTTCTGTDACAG	1
	rpoA-03-R	GHGGCCARTTTTCHARRCGC	951
	rpoA-05-F	GCAGCDCGTGTWGARCARGC	568
	rpoA-06-R	CGYTGYTCWACACGHGCTGC	568
	recA-01-F	TGARAARCARTTYGGTAAAGG	222
RecA protein ( <i>recA</i> ; 1,300 nt)	recA-02-R	TCRCNTRTAGCTRTACC	1040
	recA-03-F	TYGGBGTGATGTTYGGTAACC	767
	recA-04-R	GGGTTACCRAACATCACVCC	769

*E. corallii* had 96.2%, 85%, and 86% *rpoA*, *recA*, and *pyrH* sequence similarity, respectively. The new species *E. corallii* was tentatively allocated to the genus *Enterovibrio* on the basis of polyphasic taxonomic analysis (41), but we anticipate that this organism may be allocated to a new genus in future studies. *Photobacterium* species had 90.4 to 100%, 80 to 100%, and 79 to 99.6% *rpoA*, *recA*, and *pyrH* sequence similarity, respectively. Strains of the genus *Vibrio* showed the highest gene sequence variation, with ca. 19% variation for *rpoA* and 27% variation for *recA* and *pyrH*. *V. logei*, *V. fischeri*, and *V. wodanis* grouped apart from the other genus members, suggesting that *Vibrio* species are polyphyletic. The two squid symbionts had 96.6, 84, and 86% *rpoA*, *recA*, and *pyrH* sequence similarity, respectively.

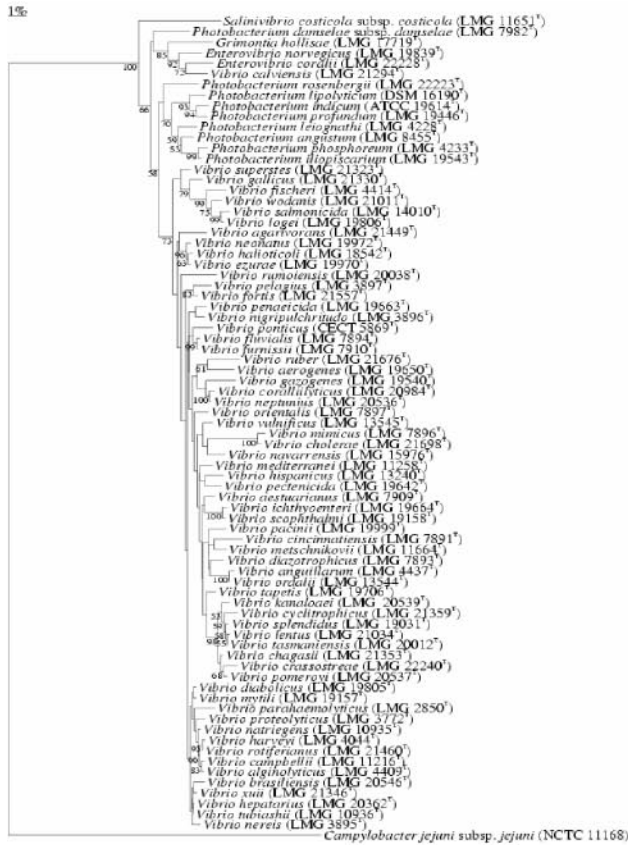
Within the *Vibrionaceae*, some pairs of highly related species, i.e., *V. aerogenes*-*V. gazogenes*, *V. fluvialis*-*V. fumissii*, *V. cholerae*-*V. mimicus*, *V. diazotrophicus*-*V. hispanicus*, *V. ichthyocenteri*-*V. scopthalmi*, and *V. anguillarum*-*V. ordalii*, appeared in the different phylogenetic trees. These pairs are indeed known to have highly related genomes, with about 70% DNA-DNA similarity. *V. haliotocoli*-, *V. harveyi*-, *V. splendidus*-, and *V. tubiashii*-related species formed groups in each tree. Roughly, the grouping of *Vibrio* species obtained with different genes is in agreement with that in previous polyphasic taxonomy studies (31–38, 40). *V. harveyi*-related species had at least 96.5%, 97%, and 95% *rpoA*, *recA*, and *pyrH* gene sequence similarity, respectively. *V. harveyi* was closely related to *V. parahaemolyticus* (98.8 to 99%), but it had only 96.5%, 97.4 to 97.7%, and 91 to 95% *rpoA*, *recA*, and *pyrH* gene sequence similarity, respectively, to its sister species *V. campbellii* and *V. rotiferianus* (9). *V. harveyi* LMG 11659, LMG 20370, and R-14913 clustered apart from the other conspecific strains, having 98.3% similarity to the other *V. harveyi* strains. These three strains were found in the former AFLP clusters A30 and A31 (31) and had about 70% DNA-DNA similarity with *V. harveyi* LMG 4044<sup>T</sup>, suggesting that they may belong to a new species (10). The gut abalone vibrios were all grouped together with at least 91.2, 97, and 93% *rpoA*, *recA*, and *pyrH* similarity, respectively. *V. haliotocoli*, *V. ezuriae*, and *V. neonatus* grouped to-

gether (93 to 99.5%) in all trees. The gut abalone vibrios have similar 16S rRNA sequences (≥98%) (12, 28), indicating that the three loci studied here are alternatives for the identification of these organisms. It is also evident that the three genetic loci are useful for the identification of the species *V. nigripulchritudo*, *V. penaeicida*, *V. tapetis*, *V. mediterranei*, *V. rumoiensis*, *V. proteolyticus*, *V. metschnikovii*, and *V. cincinnatiensis*, as they had <96% gene sequence similarity with their closest neighbors.

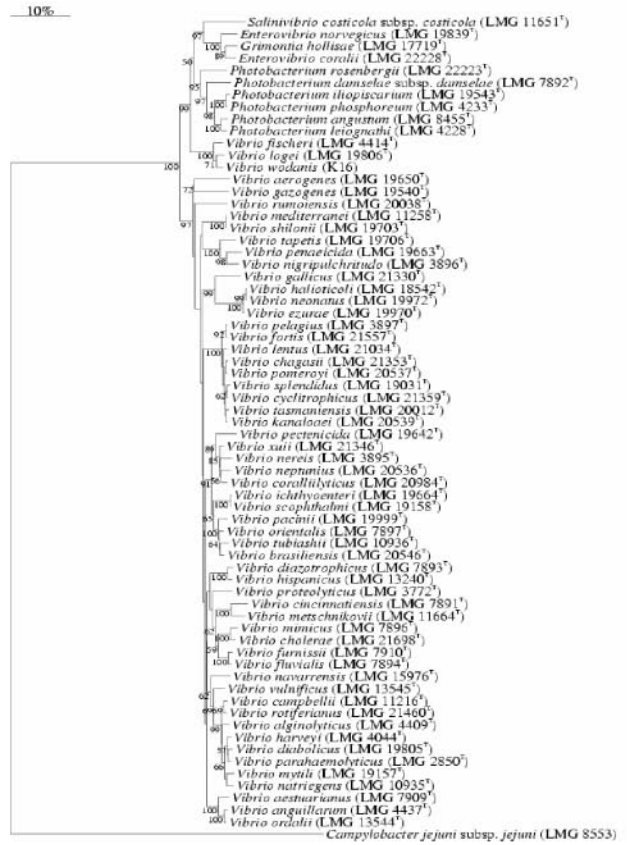
The high similarity of genomes found among different species of vibrios, e.g., in the *V. splendidus*- and *V. tubiashii*-related groups, may be explained by niche adaptation (4). *V. splendidus*-related species probably coexist in the same environment. These organisms have been associated with bivalve mollusks and with the so-called summer mortality syndrome (19). According to Cohan (4), natural selection is the main driving force in the evolution of bacterial species. He suggests that sexual isolation is not a milestone in the origin and maintenance of bacterial species. Indeed, candidate gene transfers, including paralog acquisition or the displacement and acquisition of new genes, have been detected in 5.6% (215 genes) of the genome of *V. cholerae* N16961 (17). Horizontal gene transfer (HGT) may indeed be an important force in the evolution of vibrios. The positions of various pairs of species, e.g., *V. hispanicus* and *V. diazotrophicus*, *V. mytili* and *V. diabolicus*, *V. coralliilyticus* and *V. neptunius*, and *V. brasiliensis* and *V. tubiashii*, changed in the trees constructed from the different genetic loci, suggesting that recombination might have occurred among these vibrios. Although *rpoA* and *recA* are thought to belong to the bacterial core genome and for this reason may be refractory to HGT (11), a recent study reported the HGT of *rpoA* in *Aquifex*, *Thermotoga*, and *Fusobacterium* (14). Our study did provide clear evidence of gene conversion events in the *rpoA*, *recA*, and *pyrH* ( $P < 0.05$ ) genes of *V. splendidus*- and *V. tubiashii*-related species using Sawyer's test. In addition, our splits-tree decomposition analysis revealed a network-like tree for these groups (Fig. 2). The presence of parallelograms in splits trees is a hallmark of recombination (6). Thus, we may conclude that recombination is a rather common evolutionary

FIG. 1. Phylogenetic trees based on neighbor-joining method using 16S rRNA (1,300 nt), *rpoA* (928 nt), *pyrH* (443 nt), and *recA* (613 nt) gene sequences of vibrios. Distance estimations were obtained by the model of Jukes and Cantor (16). Bootstrap percentages (≥50) after 1,000 simulations are shown. Bars, 1% (16S rRNA) and 10% (*rpoA*, *pyrH*, and *recA*) estimated sequence divergence. The *Campylobacter* NCTC 11168 sequence was used as an outgroup.

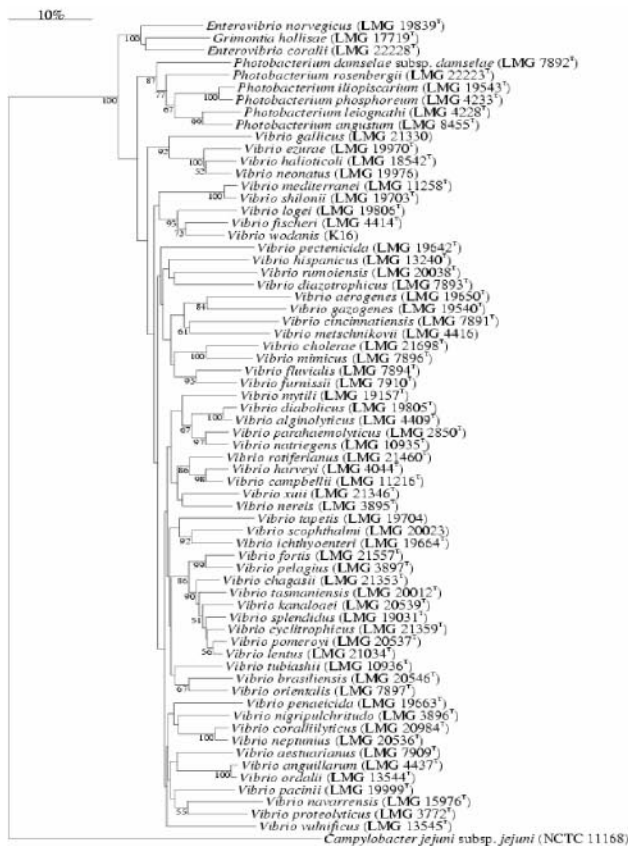
16S rDNA



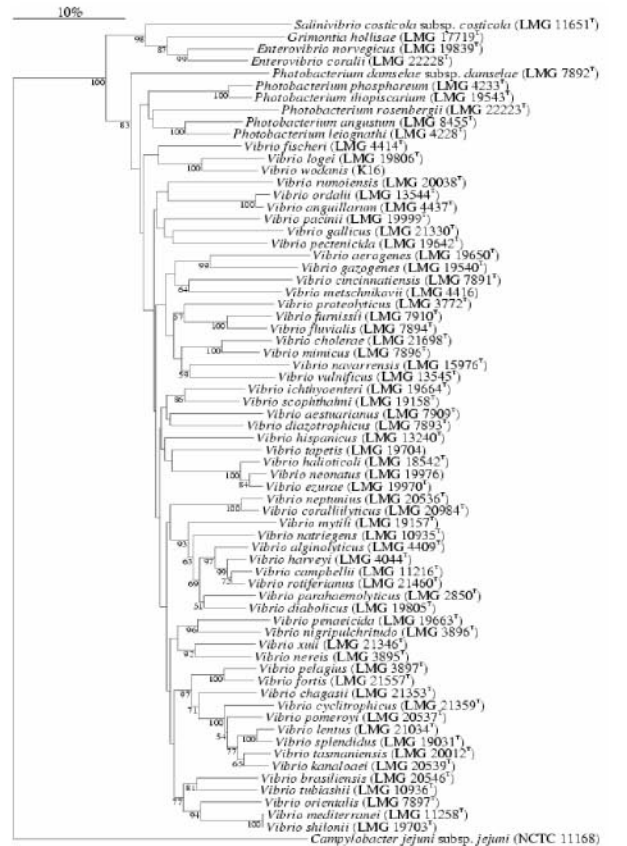
rpoA



pyrH



recA



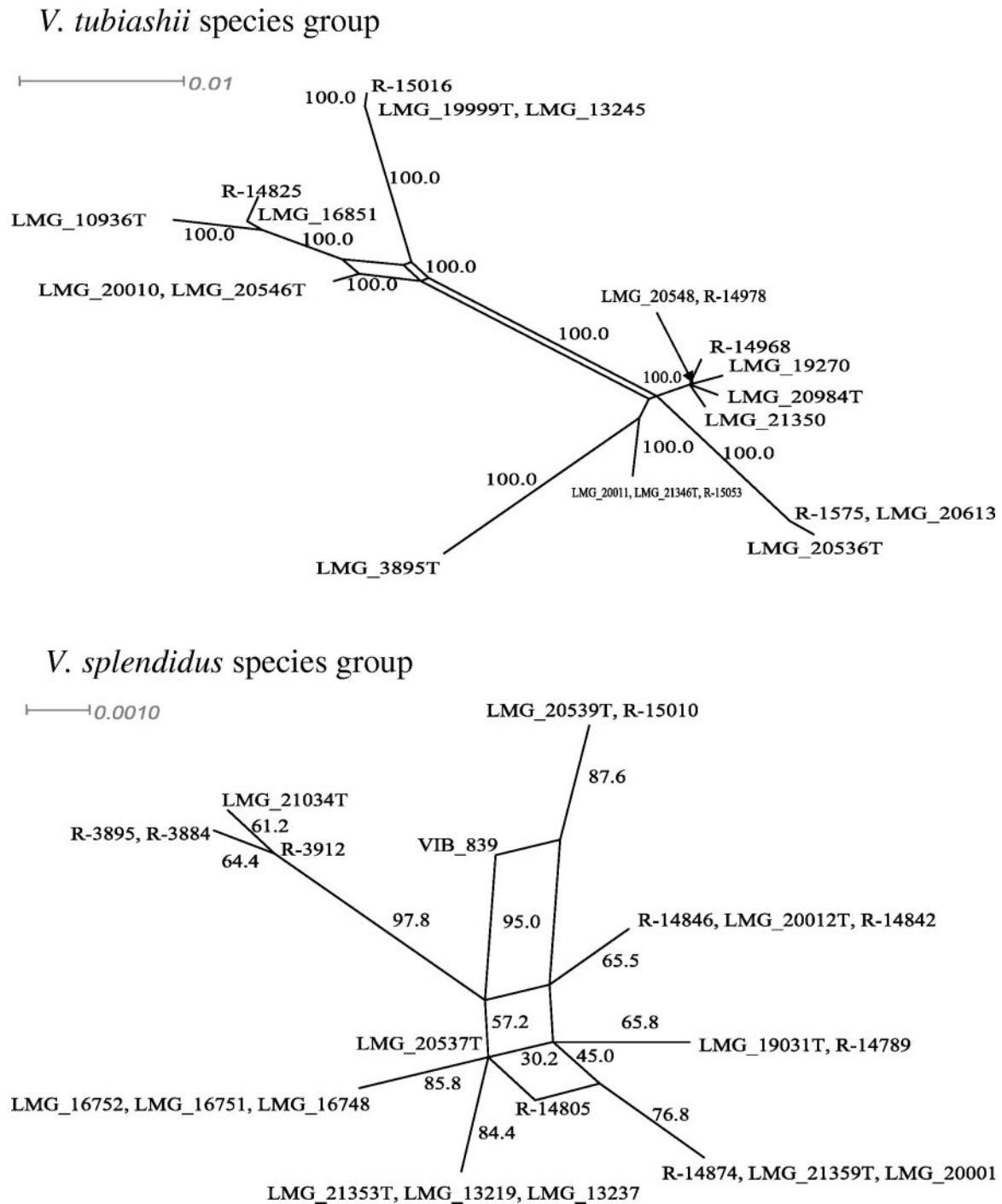


FIG. 2. Splits tree showing networks in *V. tubiashii* (A)- and *V. splendidus* (B)-related groups on the basis of *rpoA* gene sequences (931 bp). Bootstrap percentages ( $\geq 50$ ) after 500 simulations are shown. Bar, 10% estimated sequence divergence.

process within different species of vibrios. Due to such recombinational events, different vibrios, e.g., *V. tubiashii*- and *V. splendidus*-related species, may group close to one another, hampering their identification. In order to overcome the effects of gene conversion and recombination in species identification, several loci should be indexed simultaneously and used to construct multigene phylogenetic trees. The four families of vibrios were clearly apart from one another in our multigene

tree (Fig. 3). The groups obtained with this tree corresponded to those in the single-locus trees, except for the position of *V. mediterranei* along with those of *V. tubiashii*-related species. All species had  $< 97\%$  concatenated sequence similarity, with the exception of the pairs *V. coralliilyticus*-*V. neptunius* and *V. anguillarum*-*V. ordalii*.

So far, there is not a single gene that can differentiate well all species of vibrios. Different genes show different degrees of

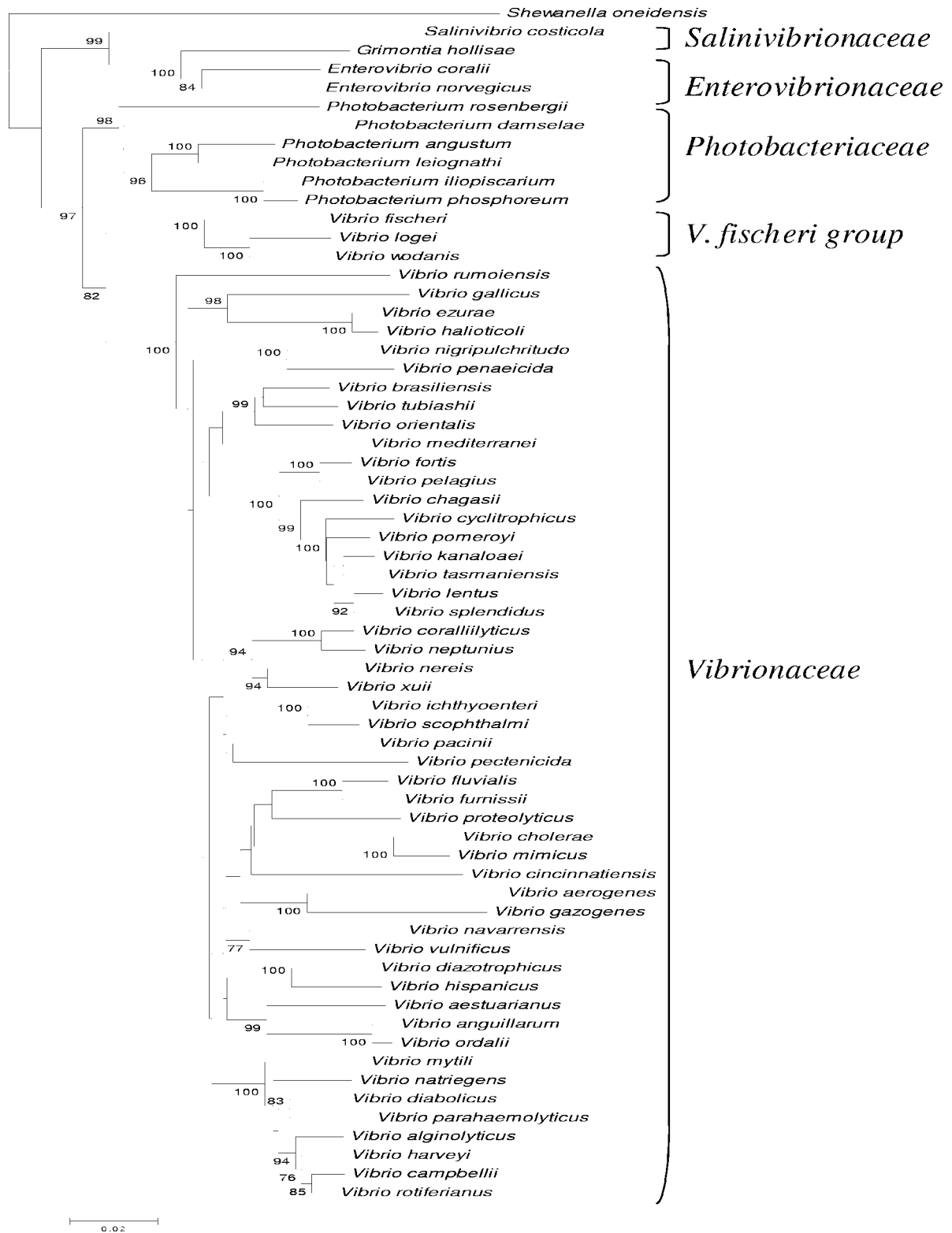


FIG. 3. Phylogenetic tree based on neighbor-joining method using the concatenated sequences (3,284 nt) of 16S rRNA, *rpoA*, *pyrH*, and *recA* of type strains. Distance estimations were obtained by the model of Jukes and Cantor (16). Bootstrap percentages ( $\geq 50$ ) after 1,000 simulations are shown. Bar, 2% estimated sequence divergence.

discrimination according to the group of vibrios under analysis. For instance, *V. cyclitrophicus*, *V. splendidus*, *V. pomeroyi*, and *V. chagasii* had about 99% *rpoA* sequence similarity. *V. splendidus*-related species are indeed highly related, having little variation (<2%) in *gyrB* and 16S rRNA gene sequences, phenotypic features, and DNA-DNA hybridization (19, 36). On the other hand, these species have <97% *recA* and *pyrH* gene sequence similarity (except for *V. lentus* and *V. splendidus*, which have 97.2% gene sequence similarity) (30). Nevertheless, the different *V. splendidus*-related species clearly formed separated clusters on the basis of the three loci, suggesting that these genes are indeed useful for the differentiation of highly related species.

Another interesting example of high gene sequence similarity in different loci is found within the *V. tubiashii*-related species. These organisms had at least 92.9% *rpoA* gene sequence similarity. *V. coralliilyticus* had 98.2, 97.9, 97.1, and 95.1% *rpoA* sequence similarity with *V. xuii*, *V. neptunius*, *V. nereis*, and *V. tubiashii*, respectively. The corresponding 16S rRNA similarity values were 96.9, 98.2, 96.7, and 97.1%, respectively (35), while those for *pyrH* (except those for *V. neptunius* and *V. coralliilyticus*, which had a 96% sequence similarity) and *recA* were below 87 and 97%, respectively. *V. tubiashii* and *V. brasiliensis* had a 98.2% gene sequence similarity for both *rpoA* and 16S rRNA (35). Overall, *rpoA* and 16S rRNA gene sequences have similar discriminatory powers for *V. tubiashii*-related species that are inferior to those of *recA* and *pyrH* (40). *V. pacinii*, *V. pectenicida*, and the pair *V. scophthalmi*-*V. ichthyenteri* had <96% *rpoA* gene sequence similarity to *V. tubiashii*-related species, suggesting that this gene is useful for the identification of these species. Overall, *rpoA* gene sequences were more discriminatory than 16S rRNA sequences. 16S rRNA and *recA* similarities above 97 and 94% corresponded to *rpoA* similarities above 88 and 97%, respectively. This fact highlights the need for future studies aiming at additional loci.

Vibrios have been implicated in the phenomenon of coral bleaching (2, 25). We used *rpoA* gene sequences to allocate taxonomically fresh isolates of vibrios associated with coral bleaching. These isolates were identified as *V. fortis*, *V. campbellii*, *V. coralliilyticus*, *V. harveyi*, *V. mediterranei*, and *V. rotiferianus*, suggesting that the process of coral bleaching may be carried out by different *Vibrio* species. *V. harveyi* has been implicated in diseases of a wide range of marine animals (1), including different coral species (10, 29). Although vibrios isolated in Hawaii were always associated with healthy corals, our data suggest that highly related strains of potentially pathogenic vibrios, e.g., R-21432 and R-23286, are present in both Kaneohe Bay (Hawaii) and the Great Coral Barrier (Australia). Environmental conditions in Australia may favor the prevalence of coral infection caused by vibrios, but this remains to be determined in future studies.

Considerable numbers of representative strains of the species *V. campbellii* ( $n = 9$ ), *V. cholerae* ( $n = 10$ ), *V. coralliilyticus* ( $n = 9$ ), *V. harveyi* ( $n = 14$ ), and *V. mediterranei* ( $n = 7$ ) were examined in this study in order to unambiguously determine the intraspecies variation of *rpoA*, *recA*, and *pyrH* gene sequences. The intraspecies gene sequence heterogeneity for most species was well below 2, 6, and 6%, respectively (Table 1). Overall, the 192 type and reference strains represent well

the currently known genomic diversity of most *Vibrionaceae* species. Strains within each of the examined species fulfill the criteria of a  $\geq 60\%$  mutual AFLP pattern similarity,  $\geq 70\%$  DNA-DNA similarity, and  $\geq 97.5\%$  16S rRNA gene sequence similarity. Representative strains were selected in order to represent the currently known genomic diversity of these species. The *V. cholerae* strains, for instance, comprise the serogroups O1, O139, and non-O1/non-O139 and represent the known genomic diversity of this species, as revealed by FAFLP analysis (39, 40). We could therefore conclude that strains of the same species will have at least 98%, 94%, and 94% sequence similarity in the *rpoA*, *recA*, and *pyrH* genes, respectively.

The data generated in this study are well suited to be used for the rapid detection and identification of pathogenic vibrios in the environment through, e.g., real-time PCR (3, 7). The data could also be an alternative to 16S rRNA gene sequences in studies of the ecology and community dynamics of vibrios in coastal waters (42, 43). The advantages of the loci studied here are that they belong to the bacterial core genome (11), have a high phylogenetic signal, and are single-copy genes and that different species of vibrios have different gene sequences that thus enable the reliable identification of these organisms. Our multilocus sequence analysis data will be used as the basis for the creation of a free-access online identification system for vibrios (<http://img.ugent.be/bnserver/MLSA/Vibrionaceae/>). Work is under way on other genes, including the *atpA*, *obg*, *uvrB*, *pheS*, and *serS* genes encoding tRNA synthases and *thd*, which together will enhance the discrimination of all currently recognized species of vibrios.

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