

Evolutionary and Diagnostic Implications of Intragenomic Heterogeneity in the 16S rRNA Gene in *Aeromonas* Strains

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Sequencing 16S rRNA genes (SSU) cloned from *Aeromonas* strains revealed that strains contained up to six copies differing by $\leq 1.5\%$. The SSU copies from *Aeromonas veronii* LMG13695 clustered with sequences from four *Aeromonas* species. These results demonstrate intragenomic heterogeneity of SSU and suggest caution when using SSU to identify aeromonads.

Differences in virulence of *Aeromonas* species, ranging from environmental isolates and digestive-tract symbionts of blood-feeding animals to human pathogens (11, 12, 16, 24, 25), led to the increased use of 16S rRNA gene (SSU) sequences to identify *Aeromonas* species (4, 9, 10). A potentially complicating factor is that bacteria contain up to 15 copies of the ribosomal operon (17). Evidence suggests that concerted evolution does not always effectively homogenize rRNA operons in bacteria and that intragenomic heterogeneity exists (5, 7, 19, 23, 29, 31, 33). Two investigations of bacterial genomes revealed that differences in the SSU sequence occurred in 54 to 62% of organisms harboring at least two copies of the rRNA operon (1, 7). However, except in rare cases, the differences were less than 1% of the compared nucleotides, and both studies concluded that these minor differences did not have a significant effect on the determination of phylogenetic relationships.

A pioneering study by Martinez-Murcia et al. used the SSU sequence to reconstruct the phylogeny of *Aeromonas* species (20). This study revealed that some *Aeromonas* species, for example, *A. trola* and *A. caviae*, differ only by up to three nucleotides in the SSU sequence, even though DNA-DNA hybridization values for these species are low (30%), indicating that they are distantly related (20). In contrast, *Aeromonas sobria* and *Aeromonas veronii* differ by 12 nucleotides but have a DNA-DNA hybridization value of 60 to 65%, indicating that they are closely related (20). This contradiction and visual inspection of the SSU sequences led Sneath to suggest that lateral gene transfer and recombination could explain this lack of congruence (27). Previously, we undertook a restriction fragment length polymorphism-PCR analysis using multiple reference strains and discovered that several strains produced unexpected restriction patterns (10). In this report, we further analyzed *Aeromonas veronii* biovar *sobria* strains A132, A155,

A916, AER28, CDC0437-84, and LMG13695, *Aeromonas veronii* biovar *veronii* strain AER397, and *Aeromonas media* strains A6 and CDC0862-83 by sequencing cloned SSU (10), determining the phylogenetic relationship of the different copies of the SSU and evaluating whether the SSU evolved as a single unit.

Sequence analysis of cloned 16S rRNA genes. The strains were grown as previously described (10), and genomic DNA was isolated as described by Nelson and Selander (22), except that the resuspended sample was preheated for 3 min at 65°C. The 16S rRNA gene was amplified by using primers 27F and 1492R (15, 18). The PCR-amplified 16S rRNA genes were cloned into pGEM-T (Promega, Madison, WI) according to the manufacturer's instructions, and the cloned 16S rRNA genes were sequenced and analyzed (15).

In six *Aeromonas* strains, the copies of the 16S rRNA genes differed by up to 21 nucleotides out of 1,377 bp analyzed (Fig. 1). If differing sequences were detected in at least two plasmids containing SSUs from one strain, we considered them to be heterogeneous copies of SSU and for this discussion call them an allele without implying an associated functional difference. The analysis of the cloned SSUs suggested the presence of at least six alleles in the *A. media* strain CDC0862-83 and at least five different alleles in the *A. veronii* biovar *sobria* strain LMG13695 (Fig. 1). The alleles from *A. media* CDC0862-83 differed from each other by up to 19 bp and those from *A. veronii* LMG13695 by up to 21 bp out of 1,377 bp compared. In a recent publication, the presence of two heterogeneous copies of the SSU was suggested for *Aeromonas trola* but not substantiated (28). It is possible that any of the examined strains contains additional alleles.

Verification of sequence differences by Southern analysis. To confirm the presence of the different SSU sequences in the same genome in a PCR-independent manner, we used Southern analysis of genomic DNA from strain CDC0862-83. Oligonucleotide probes were designed to recognize allele 1 from CDC0862-83, (16S-230-232-A [5'-GGG TTC ATC CAA TCG CG-3']), alleles 2 through 6 from CDC0836-83 (16S-230-232-B [5'-GGG CAT ATC CAA TCG CG-3']), and, as a control, all alleles (27F [5'-AGA GTT TGA TCM TGG CTC AG-3']). The hybridization was done by using a sodium chloride-sodium

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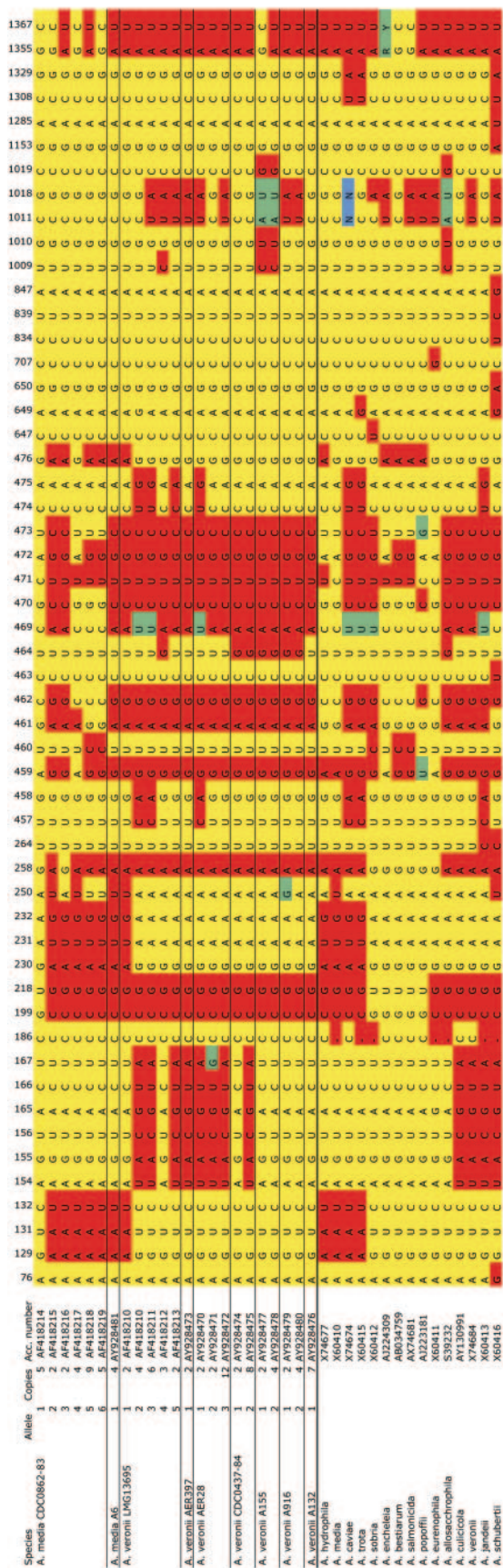


FIG. 1. Sequence differences among 16S rRNA gene alleles cloned from *A. media* and *A. veronii* strains. The diagram shows the variable sites detected in our study and in the 16S rRNA gene from *Aeromonas* species (modified from reference 27). The species names and the GenBank accession (Acc.) numbers are shown. The allele numbers are shown (Allele), as is how often each allele was cloned (Copies). The numbers in the top row correspond to the *Escherichia coli* numbering scheme. Yellow shading indicates the sequence of allele 1 from CDC0862-83 and red, green, or blue shading indicates a nucleotide substitution.

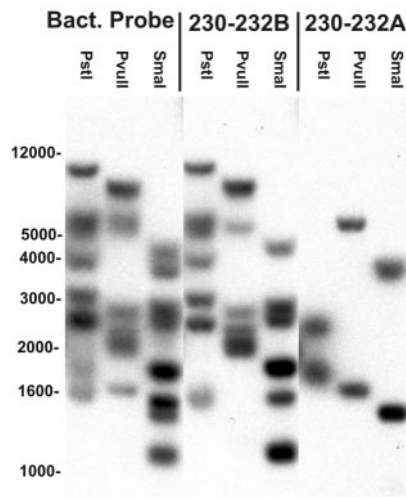


FIG. 2. Southern analysis of CDC0826-83 demonstrates that the observed variation is not a PCR artifact but encoded on the chromosome. The bacterial (Bact.) probe recognizes most bacterial 16S rRNA genes and each of the 16S rRNA gene copies in *Aeromonas*. Probe 230-232A anneals only to allele 1 from CDC0862-83, and probe 230-232B anneals to alleles 2 through 6. The hybridization patterns of the membrane probed with 230-232A show two bands hybridizing with the genomic DNA digested with PstI, PvuII, or SmaI, indicating that two copies of this polymorphism characteristic of allele 1 are present in the genome of CDC0862-83.

citrate protocol (3). Oligonucleotide probes (MWG, Münchenstein, Germany) were labeled with [α - 32 P]dCTP (3). The membranes were prehybridized for 1.5 h at 37°C, hybridized overnight at 37°C, washed in 6× SSC (1× SSC is 0.15 M NaCl plus 0.015 M sodium citrate) with 0.05% pyrophosphate for 30 min at 48°C, and used to expose BioMax MS film (Kodak, Rochester, N.Y.).

The hybridization patterns of the allele-specific probes demonstrated that the polymorphism at nucleotides 230 through 232 of allele 1 was present in two copies and that the remaining copies hybridized with the probe for alleles 2 through 6. These results indicate that at least two different alleles were present in the genome of *A. media* strain CDC0862-83 (Fig. 2). The banding pattern suggests the presence of at least eight copies of SSU in CDC0862-83 (Fig. 2).

Phylogenetic analysis. We assessed the significance of the sequence differences by aligning sequences in ClustalW version 1.83 (30) and constructing phylogenetic trees using the PhyML program version 2.4.1 (13) under the HKY85 substitution model (14) with among-site rate variation described through a discrete approximation of the gamma distribution with four rate categories (32). One hundred bootstrap samples were generated and evaluated in PhyML under the same model, and the consensus tree was generated by using the CONSENSE program from the PHYLIP package (8).

The phylogenetic analysis shows that the phylogenetic placement of the SSU is affected by the intragenomic heterogeneity (Fig. 3). The SSU alleles from the *A. media* strain CDC0862-83 clustered either within the *Aeromonas hydrophila*-*A. media* clade that has a 77% bootstrap support value or within a large clade that contained *Aeromonas encheleia*, *Aeromonas eucrenophila*, and other species with a 75% bootstrap value. Al-

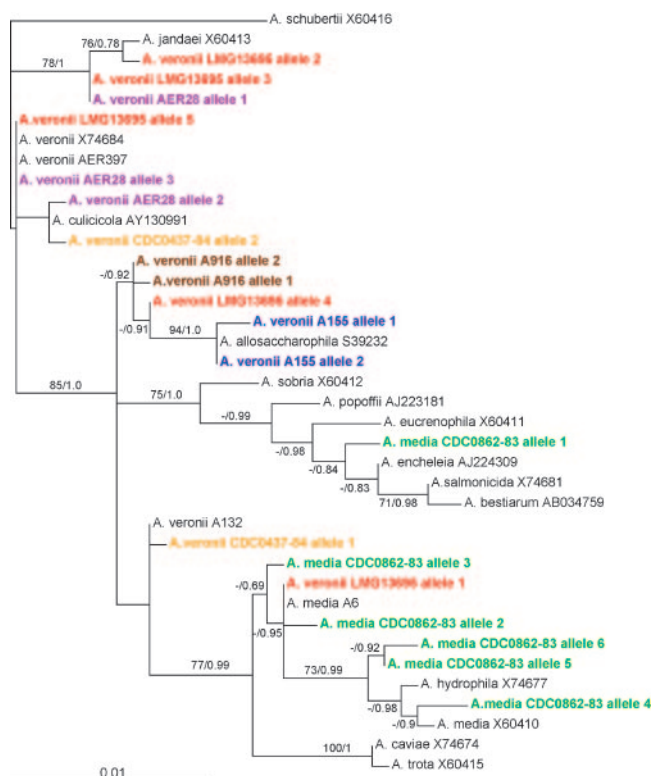


FIG. 3. Maximum-likelihood tree from published 16S rRNA genes of *Aeromonas* species and cloned alleles from the CDC0862-83 and LMG13695 strains. Selected bootstrap support values and posterior probabilities are shown as the first and second numbers, respectively. The tree is calculated from the full-length alignment using PhyML. Individual alleles of the same strain sometimes form sister groups with other *Aeromonas* species. Moreover, the analyses of bipartitions show that alleles have different phylogenetic relationships among each other if different fragments of the alignment are analyzed (Table 2).

les from the *A. veronii* strain LMG13695 grouped with sequences of *A. veronii*, *Aeromonas jandaei*, and *Aeromonas allosaccharophila* and with sequences of the *A. hydrophila*-*A. media* group. Another *A. veronii* biovar *sobria* strain, A155, clustered close to *A. allosaccharophila*, and *A. veronii* biovar *sobria* strain AER28 clustered near *A. jandaei* and *A. veronii*. All of these strains are well characterized and were previously identified to the species level by DNA-DNA hybridization (2, 6, 10). Restriction fragment length polymorphism analysis of 82 surveyed *Aeromonas* strains (10, 21) suggests that at least 21% had intragenomic heterogeneous SSU alleles as determined by the presence of faint bands that could not be removed by adding more restriction enzyme (data not shown) indicating that SSU heterogeneity is not a rare phenomenon in this genus. Our results suggest that the SSU sequences may not properly reflect the taxonomic relationship of these strains and are not the best choice for identifying *Aeromonas* species (10).

Mosaic evolution of SSU. Visual inspection of the alignment (Fig. 1) suggested that different parts of the sequence from one allele were most similar to SSU sequences from distantly related *Aeromonas* species. These conflicting phylogenetic relationships suggest that some of the SSU sequences are mosaic. We tested the hypothesis by dividing the alignment into three

TABLE 1. Results of AU test^a

Fragment	Topology	P value
W1	W1	0.997
	W2	2×10^{-13}
	W3	4×10^{-80}
	F	0.003
W2	W1	9×10^{-62}
	W2	1.000
	W3	1×10^{-40}
	F	3×10^{-49}
W3	W1	5×10^{-07}
	W2	6×10^{-52}
	W3	1.000
	F	5×10^{-06}
F	W1	1×10^{-48}
	W2	7×10^{-05}
	W3	1×10^{-118}
	F	1.000

^a The fit of four tree topologies (calculated from full alignment [F] and three fragments) was evaluated for full-length alignment and three fragments. The significantly better topology per analyzed fragment is shown in bold. In all cases, the tree topology calculated for each data set is significantly better than the alternatives.

fragments: nucleotides 1 through 264 (fragment W1), 457 through 476 (fragment W2), and 1009 through 1492 (fragment W3), and we compared the phylogenies derived from each fragment and from the entire sequence. The resulting trees were incongruent (topologies not shown). The approximately unbiased (AU) test (26) demonstrated that each of the trees calculated as optimal for one fragment had a significantly better fit to the data than the alternative trees calculated from the other fragments (Table 1). These results support the concept that the SSU evolved in a mosaic manner. This was further supported by a bipartition analysis that focuses on all significantly supported bipartitions (34) (Table 2). The different fragments strongly supported conflicting bipartitions, i.e., bipartitions that cannot coexist in a single phylogenetic tree.

One intriguing result is the clustering of *A. media* CDC0862-83 allele 1 in the branch containing *A. encheleia* and *A. eucrenophila*, because in the *rpoD*-derived phylogenetic tree, *A. media* clusters with these two species (28), while in the 16S rRNA-derived phylogenetic tree, *A. media* clusters with *A. hydrophila* (20). This discrepancy and the previously noted discrepancy between the phylogeny derived by SSU sequences and by DNA-DNA reassociation constants can be explained by the intragenomic heterogeneity. Two possible scenarios can explain the observed data. First, ancient polymorphisms could have preceded the speciation events in the genus *Aeromonas*, and these polymorphisms could have persisted throughout the history of the species. Our analysis shows that the different alleles do not cluster but intersperse in the reconstructed phylogeny. Although there was sufficient time for alleles from other lineages to be lost over evolution, for some reason, other alleles that are very similar to distantly related species were maintained. The alternative scenario suggested by Sneath (27), horizontal gene transfer, appears more likely. In either case,

TABLE 2. Supported bipartitions for relationships among alleles^a

Fragment	Supported bipartitions for indicated strain (bootstrap value [%])	
	CDC0862-83	LMG13695
Full alignment	1, 2, 3, 4 5, 6 (70)	1, 4 2, 3, 5 (97)
	1, 2, 3 4, 5, 6 (77)	1, 4, 5 2, 3 (89)
W1	1, 3 2, 4, 5, 6 (85)	1, 4 2, 3, 5 (100)
W2	1, 4, 5, 6 2, 3 (100)	1, 4, 5 2, 3 (100)
	1, 2, 3, 4 5, 6 (99)	
	1, 4 2, 3, 5, 6 (69)	
W3	1, 2, 4, 6 3, 5 (83)	1, 2 3, 4, 5 (94)
		1, 2, 4 3, 5 (75)

^a Alleles are numbered using Arabic numerals. Taxa on the opposite sides of bipartition are separated by the vertical bar: e.g., the bipartition "1, 4 | 2, 3, 5" stands for alleles 1 and 4 on one side of the bipartition and 2, 3, 5 on the other. The bootstrap support values for bipartitions are shown in parentheses. Only nontrivial bipartitions with bootstrap support values greater than 68% are shown. Note that each fragment supports only bipartitions that are compatible with the other bipartitions supported by this fragment; however, the sets of bipartitions supported by different fragments frequently are incompatible with one another.

the alleles present today resulted from recombination between divergent SSUs, and thus, one would not expect that the SSU phylogeny always reflects the evolutionary history of the strain in this genus.

For the identification of *Aeromonas* spp., it is critical to use multiple molecular markers, such as *rpoD* and *gyrB*, in the phylogenetic analysis (28) or an even broader approach utilizing multilocus sequence typing. Whole-genome sequence comparisons of several strains from each species might provide the insight needed to resolve the complex evolutionary history that probably included frequent gene duplication, gene transfer, and recombination events.

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