

Letters to the Editor

Molecular Analysis of Tn1546 in *vanA*-Containing *Enterococcus* spp. Isolated from Humans and Poultry

The genes encoding the VanA type of vancomycin resistance in enterococci are located on elements related to Tn1546 (1). Heterogeneity of Tn1546 in *Enterococcus faecium* has been previously reported (3, 4, 6, 9, 10, 12, 14, 15, 16) and includes deletions, insertions, and mutations. In this study, Tn1546 elements from 43 *vanA*-containing isolates of different enterococcal species with diverse pulsed-field gel electrophoresis (PFGE) patterns (criteria of Tenover et al. [13]) were analyzed: 30 *E. faecium* isolates (18 from chicken feces or products and 12 from human fecal samples) showing 25 unrelated PFGE patterns, 7 *Enterococcus durans* isolates with two unrelated PFGE patterns, (chicken feces or products), 5 *Enterococcus hirae* isolates (4 from chicken feces or products and 1 from a human fecal sample known to have an indistinguishable or a closely related PFGE pattern), and 1 *Enterococcus faecalis* isolate (from a chicken product).

PCR products were obtained from all isolates for the seven genes of Tn1546 (*vanR*, *vanS*, *vanH*, *vanY* [10], *vanA* [16], *vanX* [11], and *vanZ* [8]); IS1216V-related sequences (6) were also demonstrated by PCR for all 43 isolates. When the *vanXY* (9) region was amplified, the expected 1,947-bp fragment was obtained with 36 of the 43 isolates; however, in all 5 *E. hirae* isolates and in 2 *E. faecium* isolates (both from chickens, with different PFGE patterns) the fragment amplified was longer than expected. Results of hybridization of *vanXY* PCR products with an IS1216V probe indicated that IS1216V was located within the *vanXY* region in these seven isolates and outside this region in the other isolates. An IS1216V-like sequence was first described within the intergenic *vanXY* region (5) and later both within and outside Tn1546 (3, 6); disruption of *vanS* by IS1216V has been found in a clinical *E. faecium* isolate (2). Despite these reports for *E. faecium*, this is the first time that IS1216V has been reported within the *vanXY* region of *E. hirae*. The restriction of the IS1216V-*vanXY* association to *E. hirae* and two *E. faecium* strains isolated from chickens could suggest interspecies transmission of these transposons in animal gastrointestinal tracts.

IS1251 sequences (6) were detected in 18 *E. faecium* isolates (13 from chickens and 5 from humans; 15 unrelated PFGE patterns) but not in the other species tested. Analysis of *vanSH* (9) amplicons and hybridization showed that IS1251 was not included in this region. IS1251 has been previously found in the *vanSH* intergenic region and at other sites in *E. faecium* (4, 6).

IS1476, first found in the *vanY* gene of an *E. faecium* isolate (9), was not detected in our strains nor in two other studies (6, 15). Recently, a 1-bp difference in *vanX* was found at position 8234, with either a G (G type) or a T (T type) (6). All our *vanA* isolates, except one (*E. faecium* from ground chicken), belonged to the G type as determined by *DdeI* digestion of the *vanX* gene PCR fragment (315 bp) (6). The G type has been associated with poultry, and the T type has been associated with porcine *E. faecium* isolates (3, 7). Both types have been found among isolates from humans in different countries (7), although we found only the G type among vancomycin-resistant enterococci isolated from humans in Spain.

In conclusion, *vanA*-containing *E. faecium*, *E. faecalis*, *E.*

hirae, and *E. durans* strains of human and animal origins were found to contain similar genetic arrangements of the *vanA* gene cluster, suggesting either horizontal transfer, the existence of a common reservoir, or a predilection for insertion of certain elements at specific sites.

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