VanB Phenotype-vanA Genotype Enterococcus faecium with Heterogeneous Expression of Teicoplanin Resistance

Il Joong Park, Lee, Jong Hee Shin, Kyung Won Lee, and Gun Jo Woo

Department of Laboratory Medicine, Ajou University School of Medicine, Suwon, Korea; Department of Laboratory Medicine, Chonnam National University Medical School, Gwangju, Korea; Department of Laboratory Medicine, Yonsei University College of Medicine, Seoul, Korea; and Center for Food Safety Evaluation, Korea Food and Drug Administration, Seoul, Korea

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VanA glycopeptide resistance is characterized by acquired inducible resistance to both vancomycin and teicoplanin, whereas the VanB phenotype is characterized by variable levels of resistance to vancomycin but susceptibility to teicoplanin (1, 5). During the past 5 years, vancomycin-resistant enterococci (VRE) have become increasingly prevalent in Asia (8, 11, 12). In 2005, we experienced an outbreak of VanB phenotype-vanA genotype Enterococcus faecium strains with heterogeneous expression of teicoplanin resistance isolated from six patients at a tertiary care teaching hospital. To our knowledge, this is the first description of IS1216V inserted in the coding region of vanS. This could be the underlying cause of the VanB phenotype-vanA genotype with heterogeneous expression of teicoplanin resistance.

From November to December 2005, six VanB phenotype-vanA genotype E. faecium isolates (AJJ2, AJJ3, AJJ4, AJJ5, AJJ8, and AJJ10) were collected from a tertiary care teaching hospital. The organisms were identified by using conventional biochemical reactions and a Vitek identification system (bioMérieux, Hazelwood, MO). To evaluate the genetic relatedness of the isolates, pulse-field gel electrophoresis (PFGE) was performed with SmaI-digested genomic DNA (Gibco BRL, Gaithersburg, MD) as described by Murray et al. (13), with pulse times beginning with 1 s and ending with 20 s at 6 V/cm for 24 h. Dendrograms based on Dice coefficients and an arithmetic averages algorithm were generated with Bio-Gene software (Vilber Lourmat, Marne-la-Vallée, France). All six VanB phenotype-vanA genotype E. faecium isolates revealed identical or closely related PFGE patterns according to the method of Tenover et al. (14). The MICs of vancomycin and teicoplanin for the isolates were determined by Etest (AB Biodisk North America, Inc., Culver City, CA). Previously characterized VRE strains, E. faecium BM4147 (3) and VanB phenotype-vanA genotype E. faecium JC03 (12) without heterogeneous expression of teicoplanin resistance, served as controls.

All isolates displayed various levels of vancomycin resistance (MICs, 64 to 128 μg/ml) and teicoplanin susceptibility (MICs, 4 to 12 μg/ml) with heterogeneous expression of teicoplanin resistance. The heterogeneous expression of resistance was characterized by the growth of colonies in the elliptic inhibition zone during Etest susceptibility testing (Fig. 1A). Colonies in the elliptic inhibition zone displayed a homogeneous phenotype of resistance to teicoplanin when retested by Etest (Fig. 1B). VanB phenotype-vanA genotype E. faecium JC03 (12) without heterogeneous expression of teicoplanin resistance presented teicoplanin susceptibility and displayed no growth of colonies in the inhibition zone (Fig. 1C), and E. faecium BM4147 (3) displayed a homogeneous resistance to teicoplanin (Fig. 1D).

Bacterial DNA was extracted with a Qiagen DNeasy kit (Qiagen GmbH, Hilden, Germany) according to the manufacturer’s instructions. For structural analysis of Tn1546-like elements, an overlapping PCR amplification of internal regions of Tn1546 was performed as described previously (9). PCR amplification with IS1216V primer and IS1216V primer were also purified and subsequently sequenced to determine the exact integration site and orientation of the IS1216V insertion. Nucleotide sequences were compared to the reference sequence of transposon Tn1546 (GenBank accession number M97287) (3).

Filter matings were performed with Enterococcus faecalis JH2-2 (10) as the recipient and all six isolates as the donors, as previously described (15). Transconjugants were selected on brain heart infusion (BHI) agar plates containing 50 μg/ml of rifampin, 20 μg/ml of fusidic acid, and 10 μg/ml of vancomycin. The transconjugants were examined for the structural analysis of Tn1546.

The induction of resistance to teicoplanin in VanB pheno-
type-vanA genotype *E. faecium* AJJ5 with heterogeneous expression was studied by the determination of growth rates after overnight incubation with or without teicoplanin. The isolate was grown overnight at 37°C in BHI broth with or without teicoplanin (6 μg/ml). The cultures were diluted 1:20 into 20 ml of BHI with or without teicoplanin (6 μg/ml), grown at 37°C with shaking, and monitored for their optical density at 600 nm. Previously characterized VRE strains, *E. faecium* BM4147 (3) and VanB phenotype-vanA genotype *E. faecium* JC03 (12) without heterogeneous expression of teicoplanin resistance, served as controls.

A total of 12 isolates (six original isolates and six resistant derivatives obtained from the elliptic inhibition zone on an Etest strip, respectively) were characterized by one copy of IS1216V in the left end of Tn1546, a second in the coding region of vanS, and a third in the vanX-vanY intergenic region, as well as IS1542 in the orf2-vanR intergenic region. IS1216V in vanS was integrated at nucleotide 4900 with an 8-bp duplication of the target sequence (GTCATTAG) in the forward orientation. Although the VanS sensor peptide was reported earlier to have no functional effect due to the disruption of vanS by IS1216V (6), IS1216V in the coding region of vanS in our cases might affect the expression of teicoplanin resistance. The VanRSA system activates transcription of the resistance genes in response to vancomycin and teicoplanin, whereas VanB-type enterococci remain susceptible to teicoplanin, which is not an inducer (2, 7). Amino acid substitutions due to the three point mutations of vanS are responsible for impaired teicoplanin resistance among vanA genotype VRE strains (8). On the other hand, the modification of signal transduction due to substitutions in the putative linker of VanS has been shown to be the most common mechanism of acquisition of inducible resistance to teicoplanin among vanB genotype VRE strains (4). Similarly, the genetic alteration of vanS due to the IS1216V insertion in our isolates might be associated with heterogeneous teicoplanin resistance.

All isolates transferred vancomycin resistance, at frequencies of $6 \times 10^{-9}$ to $2 \times 10^{-8}$ transconjugants per donor. The transconjugants revealed the same heterogeneous resistance to teicoplanin.

In the growth rate studies, after overnight incubation of AJJ5 with teicoplanin, growth was delayed at the beginning of the subculture in the presence of teicoplanin, with growth resuming after a lag phase of 4 h (Fig. 2A). The growth of VanB phenotype-vanA genotype *E. faecium* JC03 without heterogeneous expression of teicoplanin resistance was suppressed throughout the subculture in the presence of teicoplanin, regardless of prior overnight incubation (Fig. 2B). The growth of BM4147 did not show any lag phase throughout the subcultures either in the absence or presence of teicoplanin, regardless of prior overnight incubation (Fig. 2C). The resumption of growth of AJJ5, which is characterized by an IS1216V insertion in the coding region of the vanS gene, in the presence of teicoplanin after overnight incubation with teicoplanin and the homogeneous expression of teicoplanin resistance in a colony within the inhibition zone suggest that a portion of bacteria that retained teicoplanin resistance were fully induced after incubation with teicoplanin.

The heterogeneous teicoplanin resistance of our isolates was clinically important because the conventional test cannot detect heterogeneous teicoplanin resistance. These isolates might potentially give rise to homogeneous resistance during teicoplanin therapy. The results of PFGE showed genetically identical or closely related patterns for all isolates, indicating clonal distribution (14). When VanB phenotype-vanA genotype *E. faecium* strains are found, molecular typing and MIC determi-
nated by Etest are recommended during teicoplanin therapy to prevent a nosocomial outbreak.

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FIG. 2. Effect of overnight growth in the presence of teicoplanin (6 μg/ml; TP6) on E. faecium AJJ5 with heterogeneous expression of teicoplanin resistance (A), VanB phenotype-vanA genotype E. faecium JC03 without heterogeneous expression of teicoplanin resistance (B), and E. faecium BM4147 (C) in the absence (0) or presence of teicoplanin. Teicoplanin concentration in the overnight culture/in the subculture medium. O.D., optical density.

REFERENCES