## MOLECULAR EPIDEMIOLOGY OF *SERRATIA MARCESCENS* BACTERMIC ISOLATES PRODUCING AN EXTENDED-SPECTRUM β-LACTAMASE OR A NOVEL NATURAL CEPHALOSPORINASE (S4) IN SOUTHERN TAIWAN

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**BACKGROUND/AIMS:** Serratia marcescens is a gram-negative pathogen of nosocomial infections. Resistance to extended-spectrum  $\beta$ -lactams is usually mediated by a hyperproduced AmpC cephalosporinase or an extended-spectrum  $\beta$ -lactamase (ESBL), particularly CTX-M-3 in central Taiwan. We aimed to investigate the detailed ESBL and AmpC epidemiology of bacteremic *S. marcescens* isolated from one medical center in southern Taiwan.

**METHODS:** From August 1999 through July 2003, 69 non-repetitive isolates were investigated. ESBL- and AmpC-related genes including TEM, SHV, CTX-M and SRT families were performed by polymerase chain reaction. Genotype was analyzed by pulsed-field gel electrophoresis (PFGE).

**RESULTS:** ESBL production occurred in 11 (15.9%) isolates, including CTX-M-3 (n=10) and SHV-12 (n=1). All the remaining 58 isolates produced an AmpC, including a novel cephalosporinase (designated S4, n=50), SRT-2 (n=3), SST-1 (n=1), S4-like (n=2), AF384203-like (n=1) and AY538705-like (n=1) cephalosporinases. Isolates with S4 or CTX-M-3 may exhibit a similar phenotype of cefotaxime resistance but were susceptible to ceftazidime. ESBL-producers demonstrated diverse genotypes, whereas 47 (94%) of the 50 S4-producers belonged to 3 epidemic clones, including type A (n=28), type B (n=17), and type C (n=2). Strains with subtypes A<sub>1</sub> and B<sub>3</sub> have unexpectedly existed for 3 and 2 years in the surgical intensive care units and oncology wards, respectively.

**DISCUSSION/CONCLUSIONS:** Ceftotaxime resistance of our bacteremic *S. marcescens* population is mainly conferred by CTX-M-3 or AmpC (S4). The novel S4-producing strains with prolonged dissemination are first reported. Continuous active surveillance cultures with molecular analysis may assist in detecting the unnoticed epidemic clones.

Keyword: Serratia marcescens, CTX-M-3, AmpC cephalosporinase