- Cloney DL, Donowitz LG. Overgown use for infection control in nurseries and neonatal intensive care units. Am J Dis Child 1986;140:680-683.
- Evans HE, Solomon OA, Baki A. Bacteriologic and clinical evaluation of gowning in a premature nursery. J Pediatr 1971;78:883-886.
- Donowitz LG. Failure of the overgown to prevent nosocomial infection in a pediatric intensive care unit. *Pediatrics* 1986;77:35-38.
- Haque KN, Chagla AH. Do gowns prevent infection in neonatal intensive care units? J Hosp Infect 1989;14:159-162.
- 36. Agbayani M, Rosenfeld W, Evans H, Salazar D, Jhaveri R, Braun J. Evaluation of modified gowning procedures in a neonatal intensive care unit. Am J Dis Child 1981;135:650-652.
- Klein BS, Perloff WH, Maki DG. Reduction of nosocomial infection during pediatric intensive care by protective isolation. N Engl J Med 1989;320:1714-1721.
- Leclair JM, Freeman J, Sullivan BF, Crowley CM, Goldmann DA. Prevention of nosocomial respiratory syncytial virus infections through compliance with glove and gown isolation precautions. N Engl J Med 1987;317:329-334.
- Madge P, Paton JY, McColl JH, Mackie PLK. Prospective controlled study of four infection-control procedures to prevent nosocomial infection with respiratory syncytial virus. *Lancet* 1992;340:1079-1083.
- Boyce JM, Opal SM, Chow JW, Zervos MJ, Potter-Bynoe G, Sherman CB, et al. Outbreak of multidrug-resistant *Enterococcus faecium* with transferable vanB class vancomycin resistance. J Clin Microbiol 1994;32:1148-1153.
- Boyce JM, Mermel LA, Zervos MJ, Rice LB, Potter-Bynoe G, Giorgio C, et al. Controlling vancomycin-resistant enterococci. *Infect Control Hosp Epidemiol* 1995;16:634-637.
- 42. Slaughter S, Hayden MK, Nathan C, Hu T, Rice T, Van Voorhis J, et al. A comparison of the effect of universal use of gloves and gowns with that of glove use alone on acquisition of vancomycin-resistant enterococci in a medical intensive care unit. Ann Intern Med 1996;5:448-456.
- Boyce JM. Vancomycin-resistant Enterococcus: detection, epidemiology, and control measures. Infect Dis Clin North Am 1997;11:367-384.
- 44. McDowell JW. An environmental, economic, and health comparison of single-use and reusable drapes and gowns. Asepsis 1993:1-15.
- Smith JW, Nichols RL. Barrier efficiency of surgical gowns: are we really protected from our patients' pathogens? Arch Surg 1991;126:756-763.
- DiGiacomo JC, Odom JW, Ritota PC, Swan KG. Cost containment in the operating room: use of reusable versus disposable clothing. Am Surg 1992;58:654-656.

- Murphy L. Cost/benefit study of reusable and disposable OR draping materials. J Health Mater Manage 1993;11:44-48.
- Tyler DS, Lyerly HK, Nastala CL, Shadduck PP, Fitzpatrick KT, Langlois AJ, et al. Barrier protection against the human immunodeficiency virus. *Curr Surg* 1989;46:301-304.
- Shadduck PP, Tyler DS, Lyerly HK, Sebastian MW, Farnitano C, Fitzpatrick KT, et al. Commercially available surgical gowns do not prevent penetration by HIV-1. Surgical Forum 1990;41:77-80.
- Beck WC, Collette TS. False faith in the surgeon's gown and surgical drape. Am J Surg 1952;83:125-126.
- Bernard HR, Beck WC. Operating room barriers—idealism, practicality and the future. Bull Am Coll Surgeons 1975;60:16.
- Smith JW, Tate WA, Yazdani S, Garcia RY, Muzik AC, Nichols RL. Determination of surgeon-generated gown pressures during various surgical procedures in the operating room. *Am J Infect Control* 1995;23:237-246.
- Granzow JW, Smith JW, Nichols RL, Waterman RS, Muzik AC. Evaluation of the protective value of hospital gowns against blood strikethrough and methicillin-resistant Staphylococcus aureus penetration. Am J Infect Control 1998;26:85-93.
- Quebbeman EJ, Telford GL, Hubbard S, Wadsworth K, Hardman B, Goodman H, et al. In-use evaluation of surgical gowns. *Gynecol Obstet* 1992;174:369-375.
- Pissiotis CA, Komborozos V, Papoutsi C, Skrekas G. Factors that influence the effectiveness of surgical gowns in the operating theatre. *Eur J* Surg 1997;163:597-604.
- 56. Ahmad FK, Sherman SJ, Hagglund KH. The use and failure rates of protective equipment to prevent blood and bodily fluid contamination in the obstetric health care worker. *Obstet Gynecol* 1998;92:131-136.
- Flaherty AL, Wick TM. Prolonged contact with blood alters surgical gown permeability. Am J Infect Control 1993;21:249-256.
- Werner HP, Feltgen M. Quality of surgical drape and gown materials. Hyg Med 1998;23:9-29.
- Lovitt SA, Nichols RL, Smith JW, Muzik AC, Pearce PF. Isolation gowns: a false sense of security? Am J Infect Control 1992;20:185-191.
- McCullough EA, Schoenberger LK. Liquid barrier properties of nine surgical gown fabrics. INDA Journal of Nonwovens Research 1991;3:14-20.
- McCullough EA. Methods for determining the barrier efficacy of surgical gowns. Am J Infect Control 1993;21:368-374.
- Leonas KK, Jinkins RS. The relationship of selected fabric characteristics and the barrier effectiveness of surgical gown fabrics. Am J Infect Control 1997;25:16-23.
- Belkin NL. Evaluating the protective value of hospital gowns. Am J Infect Control 1999;27:178-179.

Vancomycin Use and VRE Colonization in Dialysis Patients

Gina Pugliese, RN, MS Martin S. Favero, PhD

Vancomycin is widely used with hemodialysis patients as empirical therapy for dialysis-associated infections. To determine the relation of this practice to the generation of vancomycin-resistant enterococcal (VRE) colonization, Atta and coinvestigators from Johns Hopkins in Baltimore, Maryland, conducted a 2-year prospective cohort study. Cultures for VRE were taken from rectal swabs obtained from patients at the start and finish of the study period and during interim hospitalizations.

The results showed that 10 of 124 patients initially grew VRE. Twenty-four of the remaining patients had no follow-up cultures because of patient death (62%), transfer to another dialysis facility

(17%), patient's refusal (7%), and transplantation (4%), and thus were excluded. The remaining patients (n=90) had a median age of 54.3 years and were 92% African American and 50% male. Fifty-eight percent were treated by hemodialy-sis. They received 403 g of intravenous vancomycin over 157.2 patient-years of follow-up, 73% as outpatients.

Sixteen (17.8%) of 90 patients became colonized with VRE, an incidence rate of one case per 9.8 patient-years of follow-up. None of the 29 patients who did not receive vancomycin developed VRE, compared with 26% of those treated with vancomycin (*P*=.001). The odds ratio (95% confidence interval) for the association of outpatient vancomycin (*g*/*y*) with VRE colonization was 1.23 (1.05, 1.44, *P*=.008). The association remained significant following adjustment in separate logistic regression analyses for relevant demographic, clinical, antimicrobial (inpatient vancomycin, oral or intravenous cephalosporins, aminoglycosides, quinolones, or anti-anaerobics), and hospitalization exposures. The unadjusted relative risk of death in patients growing VRE was significantly higher than in those not colonized with VRE (P=.005).

The authors concluded that VRE colonization is a relatively common and underrecognized problem among chronic dialysis patients and is strongly and independently associated with the outpatient use of vancomycin, which should be avoided whenever possible.

FROM: Atta MG, Eustace JA, Song X, Perl TM, Scheel PJ. Outpatient vancomycin use and vancomycin-resistant enterococcal colonization in maintenance dialysis patients. *Kidney Int* 2001;59: 718-724.