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Endotoxins in Dialysate Cause Outbreak of Peritonitis

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In August 1996, five states identified outbreaks of culture-negative peritonitis in peritoneal dialysis patients. The CDC investigated these outbreaks to determine the cause and extent of culture-negative peritonitis among New York patients on either continuous ambulatory peritoneal dialysis (CAPD) or continuous cyclic peritoneal dialysis (CCPD). Information from patients was obtained by a telephone survey of all New York outpatient dialysis centers, using a standardized questionnaire. Dialysis solutions from one manufacturer were tested by the FDA for bacterial and

endotoxin contamination. The results of the investigation found that 95 dialysis centers, with a total of 2,471 peritoneal dialysis patients, reported 97 cases (3.9%) of culture-negative peritonitis. This included culture-negative peritonitis in 79 (8.7%) of 1,554 CCPD patients compared to 18 (1.2%) of 907 CAPD patients. Among the CCPD patients, 27.2% (73/268) using products from company A developed culture-negative peritonitis, compared with 0.9% (6/639) using products from company B. No association was found between products used and culturenegative peritonitis in CAPD patients. Two samples of company A dialysate retrieved from CCPD case patients had high levels of endotoxin (>1.25

endotoxin units/mL). The investigation implicated the use of dialysate from company A by CCPD patients in the largest reported outbreak of culture-negative peritonitis in peritoneal dialysis patients. The authors conclude that current standards for testing dialysate solutions prior to sterilization may not protect these patients from illness due to residual endotoxin.

FROM: Hopkins DP, Cicirello H, Dievendorf G, et al. An outbreak of culture-negative peritonitis in dialysis patients. Presented at the 46th Annual Epidemic Intelligence Service Conference; April 14-18, 1997; Atlanta, GA.