(continued from page 206)

tween IV drug administration and surgery should not exceed 1 hr.^{4,5}

REFERENCES

- 1. Faden H. Prophylactic antibiotics in pediatric orthopedic surgery: Current practices. Infection Control 1981, 2:107-9.
- 2. Pavel A, Smith RL, Ballard A et al. Prophylactic antibiotics in elective orthopedic surgery: A prospective study of 1,591 cases. South Med J 1977, 70:50-5.
- 3. Veterans Administration Ad hoc Interdisciplinary Advisory Committee on Antimicrobial Drug Usage. 1. Prophylaxis in surgery. JAMA 1977, 237:1003-7.
- Polk H, Lopez-Mayor J. Post-operative wound infections: A prophylactic study of determinant factors for prevention. Surgery 1969, 66:97-103.
- 5. Griffith D, Simpson R, Shoney B et al. Single-dose preoperative antibiotic prophylaxis in gastrointestinal surgery. Lancet 1976. 2:325-328.

Milap C. Nahata, Pharm. D. Ashok K. Chawla, B.S. Valerie Bookwalter, B.S. Joseph Talarico, B.S. Dwight A. Powell, M.D. Colleges of Pharmacy and Medicine The Ohio State University Children's Hospital Department of Pediatrics Columbus, Ohio

Filter Use for Hyperalimentation Therapy

To the Editor:

I would like to inquire about your recommendations regarding the use of filters for hyperalimentation therapy.

The current policy and procedure for parenteral therapy at our 246-bed hospital includes changing the intravenous tubing and the .22 micron filter every 24 hours.

In my clinical practice, I have found that before the 24 hours is complete, by the process of elimination, occlusion is traced to the filter. Therefore, either complete tubing change or just the filter change is necessary. Obviously, only changing the filter breaks the system, which is not acceptable. Do you recommend a larger size filter or none at all?

I would appreciate your recommendations on this subject, as I am the nurse on our Nutrition and Metabolic Support Service Team.

> Rosemary Blevins, R.N. Nutrition and Metabolic Service Medical Center Hospital Largo, Florida.

This letter was referred to Richard A. Garibaldi, M.D., for his comments.

A great deal of confusion still exists regarding the need for bacteria-tight filters with hyperalimentation therapy. In the early 1970s, high rates of bacterial and fungal sepsis were associated with the administration of hyperalimentation.¹ Microbiologic studies suggested that hyperalimentation solution was a nutrient media for the growth of certain fungi and gramnegative bacteria.² At that time it was felt, on a theoretic basis, that filters could prevent intrinsic contaminants from gaining access to the patient's bloodstream. Subsequently, as more stringent methods for hyperalimentation administration were developed the incidence of hyperalimentationassociated sepsis has decreased.3 Currently, it is thought that organisms causing sepsis are more likely to gain access to the blood stream by migrating along the outside of the catheter or by contamination of the infusion apparatus secondary to breaks in the closed system.⁴ Thus, some groups have felt that bacteria-tight filters are unnecessary from the point of view of infection control, and might actually increase

the risk of infection because their use necessitates frequent filter or tubing changes.

Unfortunately, no large scale, prospective, blinded trial is available which evaluates the efficacy of filters preventing hyperalimentationin associated infections. Thus, the decision to recommend or not recommend filters must be gleaned from indirect testimonials and subjective impressions. Each hospital must weigh potential risks against potential benefits. It is even more difficult to calculate costs associated with using and not using filters because data on efficacy are not available. In view of the lack of supportive data. I think that it is reasonable to forego the routine use of bacteria-tight filters for hyperalimentation infusions.

For the purposes of infection control, I would place a greater emphasis on the mechanics of infusate preparation, catheter insertion, wound care, maintenance of a closed system and avoidance of other uses for the hyperalimentation line such as blood sampling, medication administration or transfusions. Clearly, this is a subject for which more information is needed.

REFERENCES

- 1. Curry CR, Quie PG. Fungal septicemia in patients receiving parenteral hyperalimentation. New Engl J Med 285:1221-1225, 1971.
- 2. Goldman DA et al. Growth of bacteria and fungi in total parenteral nutrition solutions. Amer J Surg 126:314-318, 1973.
- Sanderson I, Deitel M. Intravenous hyperalimentation without sepsis. Surg Gyn & Obs 136:577-585, 1973.
- 4. Ryan JA et al. Catheter complications in total parenteral nutrition. New Engl J Med 290:757-761, 1974.

Richard A. Garibaldi, M.D. Associate Professor of Medicine Hospital Epidemiologist University of Connecticut Health Center Farmington, Connecticut

LONZA

Control gram positive and gram negative bacteria, fungi and viruses under the most demanding hard water conditions and organic soil contamination. . .



With New Tougher Quaternary Ammonium Compounds

Now you can have hard surface sanitizers and disinfectants that have the clout of phenolics under tough conditions—without the negative aspects of phenolics.

Ask about the new generation of synergistic Twin ChainTM quaternary ammonium compounds. Compounds that keep their muscle even under hard water conditions of up to 400 ppm and pass the EPA required microbiological tests in the presence of 5% organic soil contamination.

These new compounds also provide greater germicidal efficiency against Pseudomonas aeruginosa, Staphylococcus aureus, Salmonella choleraesuis and other gram positive and gram negative bacteria. And they are compatible with nonionic, and cationic detergents. . .even have a high tolerance for residual anionic detergents.

To get the disinfectants that are strong on germ control and pleasant to use, ask your manufacturer/supplier for complete information about these new quaternaries.

