ticularly in *S agalactiae* infections. In the early postpartum period, fever in the mothers was significantly less likely in the patients offered vaginal disinfection, a reduction from 7% in those douched using saline compared with 3% in those disinfected using chlorhexidine. A lower occurrence of urinary tract infections also was observed: 6% in the saline group as compared with 3% in the chlorhexidine group (*P*<.01).

This prospective controlled trial demonstrated that vaginal douching with 0.2% chlorhexidine during labor can significantly reduce both maternal and early neonatal infectious morbidity. The squeeze bottle procedure was simple, quick, and well-tolerated.

FROM: Stray-Pedersen B, Bergan T, Hafstad A, Normann E, Grogaard J, Vangdal M. Vaginal disinfection with chlorhexidine during childbirth. *Int J Antimicrob Agents* 1999;12:245-251.

Effectiveness of Live, Attenuated Intranasal Influenza Virus Vaccine

A recent study by Nichol and colleagues concluded that, among healthy adults, a live, attenuated influenza vaccine delivered intranasally not only helps prevent serious illness but also saves money.

In a randomized, double-blinded, placebo-controlled trial of 4,561 healthy adults aged 18 to 64, investigators found that recipients of intranasally administered trivalent, live, attenuated influenza virus (LAIV) vaccine were as likely to experience one or more febrile illnesses as placebo recipients during peak outbreak periods (13.2% for vaccine vs 14.6% for placebo). However, vaccination significantly reduced the numbers of severe febrile illnesses (18.8% reduction) and febrile upper respiratory tract illnesses (23.6% reduction). Vaccination also led to fewer days of illness across all illness syndromes (22.9% reduction for febrile illnesses; 27.3% reduction for severe febrile illnesses), fewer days of work lost (17.9% reduction for severe febrile illnesses; 28.4% reduction for febrile upper respiratory tract illnesses), and fewer days with healthcareprovider visits (24.8% reduction for severe febrile illnesses; 40.9% reduction for febrile upper respiratory tract illnesses). Use of prescription antibiotics and over-the-counter medications was also reduced across all illness syndromes. Vaccine recipients were more likely to experience runny nose or sore throat during the first 7 days after vaccination, but serious adverse events between the groups were not significantly different.

The match between the type A(H3N2) vaccine strain and the predominant circulating virus strain (A/Sydney/05/97[H3N2]) for the 1997/98 season was poor, suggesting that LAIV provided substantial crossprotection against this variant influenza A virus strain. The authors concluded that intranasal trivalent LAIV vaccine

was safe and effective in healthy, working adults in a year in which a drifted influenza A virus predominated.

FROM: Nichol KL, Mendelman PM, Mallon KP, Jackson LA, Gorse GJ, Belshe RB, et al. Effectiveness of live, attenuated intranasal influenza virus vaccine in healthy, working adults: a randomized controlled trial. *JAMA* 1999;282:137-144.

Gastrointestinal Endoscopic Reprocessing Practices in the United States

Patient infection from contaminated gastrointestinal (GI) endoscopes generally can be attributed to failure to follow appropriate reprocessing guidelines. Recently, the Food and Drug Administration recommended a 45-minute exposure of GI endoscopes to 2.4% glutaraldehyde solutions heated to 25°C. Simultaneously, the American Society for Gastrointestinal Endoscopy (ASGE), the American Gastroenterological Association, and the Society of Gastroenterology Nurses and Associates endorsed a reprocessing guideline that emphasized manual precleaning and recommended a 20-minute exposure to a 2.4% glutaraldehyde solution at room temperature. Since then, little information has become available regarding actual reprocessing practices in the United States.

Cheung and colleagues mailed a questionnaire regarding endoscopic disinfection practices to 730 randomly selected members of the ASGE; 294 (40%) responded. Appropriate manual cleaning (suctioning detergent through the accessory channel and brushing the channel and valves) was reported by 91% of respondents; 70% then used automated reprocessors for disinfection or sterilization. Glutaraldehyde was the most widely used chemical disinfectant; 85% used glutaraldehyde as one of their primary disinfectants. The most commonly used disinfection time with 2.4% glutaraldehyde was 20 minutes (83.9%) followed by 45 minutes (11.4%). Only 24% of users of 2.4% glutaraldehyde heated their solution; 60% of centers tested disinfectant concentration daily or more frequently; 74% sterilized nondisposable forceps before use; 29% of centers reused disposable endoscopic accessories (which are more frequently disinfected rather than sterilized). Twelve respondents reported cases of endoscopic cross-infection.

The authors note that a significant minority of endoscopy centers still do not completely conform to recent ASGE, American Gastroenterological Association, and the Society of Gastroenterology Nurses and Associates guidelines on disinfection, and they may not be appropriately disinfecting GI endoscopes. Rigid adherence to recommended guidelines is strongly encouraged to ensure patient safety.

FROM: Cheung RJ, Ortiz D, DiMarino AJ Jr. GI endoscopic reprocessing practices in the United States. *Gastrointest Endosc* 1999;50:362-368.

The School of Medicine at the University of Washington

is recruiting a full-time faculty member at the Assistant or Associate Professor level to lead the program in Infection Control at the University of Washington Medical Center. The closing date for applications is December 10, 1999.

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1. Publication Title	2. Publication Number	3. Filing Date	
Infection Control and Hospital Epidemiology	0899-823X	October 1, 1999	
4. asue Frequency	5. Number of Issues Published Annually	6. Annual Subscription Price	
Monthy	12	\$105.00	
7. Complete Meiling Address of Known Office of Publication	on (Not Printer) (Street.city.county.state.and ZIP+4)	Contact Person	
8900 Grove Road		Lester Robeson	
Thorofare, Gloucester County, New Jersey 08086-9	447	Telephone	
& Complete Mailing Address of Headquarters or General	Business Office of Publisher (Not printer)	(856) 848-1000 extension 245	
Slack, Incorporated 6900 Grove Road Thorofare, New Ju			
Full Names and Complete Mailing Addresses of Publish		1	
Publisher (Name and complete mailing address)			
Richard N. Rossh, Stack Incorporated, 6900 Grove Rosk	d Thorofare, New Jersey 08086-9447		
Editor (Name and complete mailing address)			
Michael D. Decker, MD,. MPH	Same as above		
Managing Editor (Name and complete mailing address)			
Strinley Strunk	Same as above	•	
hill Hame		Complete Malting Address	
PHS & WSS, Incorporated dba Slack, Incorporated	6900 Grove Road, Thorofare, New Jerse	y 08096-9447	
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13. Publication Title	14. Issue Date for Circulation Date Be	low
Infection Control and Hospital Epidemiology	September 1999	
15. Extent and Nature of Circulation	Average No. Copies Each leave During Preceding 12 Months	Actual No. Copies of Single labue Published Nearest to Filing Date
a. Total Number of Copies (Net Print Run)	6,487	6,633
Paid and/or Requested Circulation (1) Sales Through Dealers and Carriers, Street Vendors, and Counter Sales (Not Mailed)	0	0
(2) Paid or Requested Mail Subscriptions (include advertiser's proof copies and exchange copies)	5,312	5,362
c. Total Paid and/or Requested Circulation (Sum of 15b(1) and 15c(2))	5,312	5,362
d. Free Distribution by Mall (Samples, complimentary, and other free)	433	442
s. Free Distribution Outside the Mail (Carriers or other means)	272	352
f. Total Free Distribution (Sum of 15d and 15e)	705	794
g. Total Distribution (Sum of 15c and 15f)	6,017	6,158
h. Copies not Distributed (1) Office Use, Leftovers, Spolled	470	477
(2) Returns from News Agents	0	0
. Total (Sum of 15g, 15h(1), and 15h(2))	6,487	6,633
Percent Paid and/or Requested Circulation (15o/15g x 100)	88%	87%
Publication Statement of Ownership Publication required. Will be printed in the November Publication not required.	k 1999 issue of this publication.	
17. Signature and Title of Editor, Publisher, Business Manager, or Owner		Date
Richard N. Roash, VP/Group Publisher Chich and N. Coash		October 1, 1999