

Editorial

Lessons From Outbreaks Associated With Bronchoscopy

David J. Weber, MD, MPH; William A. Rutala, PhD, MPH

Bronchoscopy is currently the most commonly employed invasive procedure in the practice of pulmonary medicine.¹ An estimated 497,000 bronchoscopy procedures were performed in the United States in 1996.² Current and new applications include bronchoscopic ultrasound, laser therapy, brachytherapy, electrocautery, cryotherapy, placement of airway stents, and balloon dilatation to relieve airway obstruction caused by airway lesions.³ Flexible endoscopes also are widely used in other medical disciplines. For example, more than 10,000,000 gastrointestinal endoscopies are performed each year.⁴

Endoscopes represent the medical devices most commonly linked to nosocomial outbreaks and pseudo-outbreaks.⁵ Flexible endoscopes present a challenge for low-temperature sterilization or high-level disinfection, because they have long narrow lumens, cross-connections, mated surfaces, sharp angles, springs and valves, occluded dead ends, absorbent material, and rough or pitted surfaces.^{6,7} Failure to eradicate contamination that occurred during use may lead to person-to-person transmission of pathogens (eg, *Mycobacterium tuberculosis*); failure to prevent contamination during disinfection or storage may lead to outbreaks or pseudo-outbreaks from environmental microbes (eg, nontuberculous mycobacteria, or *Rhodotorula rubra*). In this issue, Sorin and colleagues⁸ describe the nosocomial transmission of an imipenem-resistant strain of *Pseudomonas aeruginosa*, and Kressel and Kidd⁹ describe a pseudo-outbreak involving organisms relatively resistant to glutaraldehyde (ie, *Mycobacterium chelonae* and *Methylobacterium mesophilicum*) associated with the use of contaminated bronchoscopes.

Prevention of endoscope-related infections requires strict adherence to current guidelines for cleaning and disinfection. Guidelines for disinfection of flexible endoscopes, including bronchoscopes, have been published by the Association for Professionals in Infection Control and Epidemiology, Inc.^{10,11} To date, nosocomial outbreaks have not been reported in which all current recommendations were followed scrupulously. These guidelines are based on sound scientific principles generated from several sources of data: first, studies on the natural bioburden of endoscopes and efficacy of cleaning; second, studies on the in vitro efficacy of recommended high-level disinfectants and low-temperature sterilization methods; third, studies of disinfection of simulated endoscopes or experimentally inoculated endoscopes; fourth, studies of the effectiveness of current high-level disinfection and sterilization methods in actual practice; and finally, lessons learned from outbreaks and pseudo-outbreaks involving endoscopes.

Only limited data are available on the bioburden present on bronchoscopes following use. Alfa and Sitter reported the average load on bronchoscopes before cleaning was 6.4×10^4 colony-forming units (CFUs)/mL, with streptococci and normal upper respiratory flora being reported.¹² The bioburden on used gastrointestinal endoscopes is higher, ranging from 10^6 to 10^7 CFUs for upper gastrointestinal endoscopes and 10^8 to 10^{10} CFUs for colonoscopes.¹³ Cleaning has been demonstrated to reduce the bioburden on endoscopes in most studies by more than 4 logs.¹³ Cleaning also removes organic and inorganic debris that may compromise the disinfection and sterilization process. For example, Alfa and colleagues tested several low-temperature sterilization methods (ie, ethylene

From the Division of Infectious Diseases, University of North Carolina (UNC) School of Medicine, and the Department of Hospital Epidemiology, UNC Health Care System, Chapel Hill, North Carolina.

Address reprint requests to David J. Weber, MD, MPH, CB #7030 Burnett-Womack, 547, University of North Carolina at Chapel Hill, Chapel Hill, NC 27599-7030.

The authors have received no research funds from sterilizer manufacturers, but one of the authors (WAR) received honoraria from a sterilizer manufacturer in the past 12 months.

00-ED-056. Weber DJ, Rutala WA. Lessons from outbreaks associated with bronchoscopy. Infect Control Hosp Epidemiol 2001;22:403-408.

TABLE 1
STEPS IN THE DISINFECTION PROCESS AND MECHANISMS FOR FAILURE

Disinfection Component	Reasons for Component	Mechanisms for Failure
Cleaning	Reduce bioburden	Inadequate policies
Appropriate disinfectant	Remove interfering substances: blood, salt	Inadequate staff training
	Inactivation of contaminating microbes (demonstrated efficacy and effectiveness)	Ineffective disinfectant
Contact between disinfectant and contaminating microbes	Requirement for killing	Inadequate concentration
		Inadequate duration
		AER: failure to use channel connectors
Sterilization of biopsy forceps	Eliminate contaminating microbes	AER: wrong channel connectors
		Occluded lumen
		Torn or damaged lumen
Rinse	Remove potentially toxic chemicals (eg, glutaraldehyde, hydrogen peroxide)	Inadequate policies
Prevention of recontamination	Prevent contamination with environmental microbes	Inadequate staff training
		Mucous membrane damage (eg, colitis)
		Tap water rinse without subsequent alcohol rinse
		Failure to air dry scope
		Contaminated AER
		Placement of scope in contaminated container

Abbreviation: AER, automatic endoscope reprocessor.

oxide, hydrogen peroxide gas plasma, and vaporized hydrogen peroxide) and reported that none could eradicate 10^6 CFUs of all bacterial strains inoculated on a carrier placed in a narrow lumen in the presence of 10% serum and 0.65% salt.¹⁴

Currently, the Food and Drug Administration has cleared several chemical sterilants listed as high-level disinfectants for reprocessing endoscopes.¹⁵ These include: $\geq 2.4\%$ glutaraldehyde, 0.55% ortho-phthalaldehyde, a 0.95% glutaraldehyde with 1.64% phenol/phenate, 1.0% hydrogen peroxide with 0.08% peracetic acid, 7.35% hydrogen peroxide with 0.23% peracetic acid, and 7.5% hydrogen peroxide.^{15,16} Although all of these products have excellent antimicrobial activity, 7.5% hydrogen peroxide and 1.0% hydrogen peroxide with 0.08% peracetic acid have limited use, because they cause cosmetic and functional damage to the endoscope. The two products most commonly used for reprocessing endoscopes in the United States are glutaraldehyde and the automated chemical sterilization process that uses peracetic acid (STERIS SYSTEM 1, Mentor, OH).¹⁷ The advantages and disadvantages of the chemical sterilant, peracetic acid (STERIS SYSTEM 1), and high-level disinfection methods have been reviewed.⁷

The importance of allowing the sterilant to come into contact with an inoculated carrier has been demonstrated by two studies that investigated the peracetic acid immersion system (ie, STERIS SYSTEM 1). Alfa and coworkers demonstrated excellent activity of the peracetic acid immersion system against three test organisms using a narrow-lumen device.¹⁸ In these experiments, the lumen test object was connected to channel connectors, which ensured that the sterilant had direct contact with contami-

nated carriers. The effectiveness was achieved by the combination of organism wash-off and peracetic acid inactivation of the test organisms. Data reported by Rutala and colleagues demonstrated failure of the peracetic acid immersion system to eliminate *Bacillus stearothermophilus* spores completely from an inoculated carrier placed in a stainless steel lumen test unit.¹⁹ In these experiments, the lumen test unit was not connected to channel connectors. The failure of the peracetic immersion system was felt to be attributed to the inability of the peracetic acid to diffuse into the center of a 40-cm-long, 3-mm-diameter tube, possibly due to an air lock or air bubble formed in the lumen that would impair flow.^{20,21} We have since repeated our experiments using a channel connector specially designed for our 1-, 2-, and 3-mm lumen test units, with the result that the STERIS SYSTEM 1 was completely effective in eliminating an inoculum of 10^6 *B. stearothermophilus* spores (WAR, unpublished data, October 2000). Both Sorin and colleagues⁸ and Kressel and Kidd⁹ demonstrate the clinical relevance of these findings.

Experimental contamination of flexible bronchoscopes with *Mycobacterium gordonae*²² and gastrointestinal endoscopes with duck hepatitis B virus²³ has demonstrated the importance of cleaning and validated current disinfection recommendations. The importance of cleaning also has been demonstrated in studies evaluating gastrointestinal endoscopes contaminated with *Helicobacter pylori*.²⁴ Simulated-use trials with the STERIS SYSTEM 1 have demonstrated excellent microbicidal activity, and three clinical trials have demonstrated both excellent microbial killing and no clinical failure leading to infection.⁷

Failure to follow current disinfection recommendations (Table 1) has led to multiple outbreaks^{8,25-35} (Table 2)

TABLE 2
NOSOCOMIAL OUTBREAKS VIA BRONCHOSCOPES DUE TO EXOGENOUS CONTAMINATION OR PERSON-TO-PERSON TRANSMISSION

Reference	Year	Pathogen*	Mechanism of Contamination
Webb et al ²⁵	1975	<i>Serratia marcescens</i>	Inadequate disinfectant (70% alcohol)
Hussain ²⁶	1978	<i>Pseudomonas</i>	Contaminated biopsy suction attachment (soaked in antiseptic)
Markovitz ²⁷	1979	<i>Pseudomonas pseudomallei</i>	Not specified
Leers ²⁸	1980	<i>Mycobacterium tuberculosis</i>	Inadequate cleaning and disinfectant (povidone-iodine)
Nelson et al ²⁹	1983	<i>M tuberculosis</i>	Inadequate disinfectant (povidone-iodine/70% ethanol)
Pappas ³⁰	1983	<i>Mycobacterium chelonae</i>	Two bronchoscopes with punctured suction channels
Wheeler et al ³¹	1989	<i>M tuberculosis</i>	Contaminated suction valve
Agerton et al ³²	1997	MDR <i>M tuberculosis</i>	Inadequate cleaning, failure to use leak-test equipment, no potency testing of glutaraldehyde, failure to immerse scope fully, terminal tap water without subsequent alcohol rinse
Blanc et al ³³	1997	<i>Pseudomonas aeruginosa</i>	AER: contaminated unit
Michele et al ³⁴	1997	<i>M tuberculosis</i>	Failure to use enzymatic cleaner, immerse scope fully, or sterilize biopsy forceps
Kramer et al ³⁵	2001	<i>P aeruginosa</i>	AER: contaminated disinfectant (0.04% glutaraldehyde) due to inadequate concentration (concentration mistakenly set too low)
Sorin et al ⁸	2001	<i>P aeruginosa</i>	AER: inappropriate channel connectors

Abbreviations: AER, automatic endoscope reprocessor; MDR, multidrug-resistant.
* Species as listed by investigator; may not reflect current taxonomy.

and pseudo-outbreaks^{9,36-64} (Table 3) involving bronchoscopes. The pathogen most commonly associated with outbreaks has been *M tuberculosis*, a finding that is not surprising in that only bacteria endospores are relatively more resistant than mycobacteria to disinfectants. Outbreaks associated with automatic endoscope reprocessors (AERs) commonly involve *P aeruginosa*, as was the case with the report by Sorin and colleagues.⁸ Pseudo-outbreaks most commonly involve nontuberculous mycobacteria or other water-derived environmental microbes such as *Legionella*, *R rubra*, and *P aeruginosa*. Pseudo-outbreaks also have resulted from use during bronchoscopy of contaminated medications or devices.⁶⁵ For example, pseudo-outbreaks have resulted from the use of an anesthetic contaminated with *M gordonae*⁶⁶ or fungi,⁶⁷ and atomizers contaminated with nontuberculous mycobacteria⁶⁸ or *M tuberculosis*.⁶⁹

Lessons learned from outbreaks reported in the literature include the following. First, cleaning must precede disinfection or sterilization. Second, ineffective disinfectants such as iodophors, 30% to 70% alcohol, or inadequate concentrations of disinfectant may result in outbreaks. Third, contact of all internal and external surfaces with the disinfectant is crucial. Outbreaks have resulted from failure to immerse the scope fully, disassemble valves, or repair rips or tears in internal channels. The outbreak reported by Sorin and coworkers⁸ and pseudo-outbreaks reported in the literature^{61,62,64} suggest that the proper use of channel connectors to ensure flow through an endoscope's inner channels is essential. If an AER is used, one must ensure that all channel connectors are attached according to the AER's manufacturer. Fourth, following disinfection, a sterile water rinse followed by forced-air drying or a tap water rinse followed by forced-air drying and a 70% alcohol rinse must be used to prevent recontamination. The disinfected endoscope must be stored so as to prevent recontamina-

tion. Failure to rinse the scope fully also may result in mucositis following use of the scope on another patient, if either glutaraldehyde⁷⁰ or hydrogen peroxide is used as the disinfectant. AERs offer several advantages to manual reprocessing, including automation and standardization of several important reprocessing steps,⁷¹⁻⁷³ which reduce the likelihood that an essential reprocessing step will be skipped, and reduction of personnel exposure to high-level disinfectants. However, failure of AERs has been linked to bronchoscopy-related outbreaks (Table 2) and pseudo-outbreaks (Table 3), in part because the water filtration system may not reliably be able to provide sterile rinse water.⁷⁴ It is critical that personnel rigorously adhere to the current recommendations for the use of AERs.¹¹ We agree with Sorin and colleagues that random bacterial surveillance cultures of endoscopes to assure appropriate disinfection should be done as part of a comprehensive program in quality assurance.

In conclusion, there is a need for further development and redesign of AERs⁷⁵ and endoscopes,⁶ so that they do not represent a potential source for infection. Newly developed disposable-component endoscope systems may be able to improve the ease of cleaning and disinfection and so reduce the risk of infection. Recommendations for the cleaning and disinfection of endoscopic equipment should be followed strictly. Unfortunately, audits have shown endoscopic personnel often fail to adhere to guidelines on disinfection.⁷⁶⁻⁷⁸ To ensure that persons responsible for reprocessing are properly trained, there should be initial and annual competency testing for such personnel.^{79,80}

REFERENCES

- Ahmad M, Dweik RA. Future of flexible bronchoscopy. *Clin Chest Med* 1999;20:1-17.
- Center for Disease Control and Prevention. Vital and health statistics: ambulatory and inpatient procedures in the United States, 1996. DHHS

TABLE 3
NOSOCOMIAL PSEUDO-OUTBREAKS VIA BRONCHOSCOPES DUE TO EXOGENOUS CONTAMINATION OR PERSON-TO-PERSON TRANSMISSION

Reference	Year	Pathogen*	Mechanism of Contamination
Weinstein et al ³⁶	1977	<i>Proteus</i> species	Inadequate disinfection (30% alcohol)
Dawson et al ³⁷	1982	<i>Mycobacterium intracellulare</i>	Inadequate disinfection of plastic tubing for collecting specimens
Sammartino et al ³⁸	1982	<i>Pseudomonas aeruginosa</i>	Inadequate disinfectant (povidone-iodine)
Goldstein and Abrutyn ³⁹	1985	<i>Bacillus</i> species	Contaminated automatic suction valve
Siegman-Igra et al ⁴⁰	1985	<i>Serratia marcescens</i>	Inadequate disinfection (alcohol)
Richardson et al ⁴¹	1986	<i>Bacillus</i> species	Contaminated suction valves, terminal tap water rinse
Hoffmann et al ⁴²	1989	<i>Rhodotorula rubra</i>	Contaminated channel cleaning brushes and leak-test tub water
Wheeler et al ³¹	1989	<i>Mycobacterium avium</i>	Contaminated suction valve
Nye et al ⁴³	1990	<i>Mycobacterium chelonae</i>	Contaminated tap water rinse
Fraser et al ⁴⁵	1992	<i>M. chelonae</i>	AER: contaminated AER. No terminal ethanol rinse and scopes not forced-air dried
Gubler et al ⁴⁶	1992	<i>M. chelonae</i>	AER: contaminated AER
Nicolle et al ⁴⁷	1992	<i>Blastomyces dermatitidis</i>	Inadequate disinfection of bronchoscope
Whitlock et al ⁴⁸	1992	<i>R. rubra</i>	Failure to air dry scope, contamination of suction and biopsy valves
Bryce et al ⁴⁹	1993	<i>Mycobacterium tuberculosis</i>	AER: contaminated suction valves and faulty wash/disinfect switch
Vandernbroucke-Grauls et al ⁵⁰	1993	<i>S. marcescens</i>	Inadequate immersion time (2 min), terminal tap water rinse, stored without drying
Bennett et al ⁵¹	1994	<i>Mycobacterium xenopi</i>	Inadequate disinfectant (0.13% glutaraldehyde-phenate) and exposure time, rinsed with contaminated tap water, inadequate drying
Campagnaro et al ⁵²	1994	<i>M. chelonae</i>	AER: contaminated suction valve, terminal tap water rinse
Kolmos et al ⁵³	1994	<i>P. aeruginosa</i>	Failure to clean suction and biopsy channels, inexperienced bronchoscopy staff
Maloney et al ⁵⁴	1994	<i>M. abscessus</i>	AER: contaminated AER
Petersen et al ⁵⁵	1994	<i>M. abscessus</i>	AER: contaminated AER
Hagan et al ⁵⁶	1995	<i>R. rubra</i>	Contaminated suction channel, inadequate drying
Takigawa et al ⁵⁷	1995	<i>M. chelonae</i>	AER
Wang et al ⁵⁸	1995	<i>M. chelonae</i>	AER: contaminated suction channel
Mitchell et al ⁵⁹	1997	<i>Legionella pneumophila</i>	Use of contaminated tap water for rinse, failure of 70% ethanol flush
Wallace et al ⁶⁰	1998	<i>M. abscessus</i>	AER and manual disinfection procedure
Wallace et al ⁶⁰	1998	<i>M. abscessus</i>	AER
Wallace et al ⁶⁰	1998	<i>Mycobacterium fortuitum</i>	AER
CDC ⁶¹	1999	<i>M. tuberculosis</i>	AER: failure to replace biopsy port cap before loading in AER
CDC ⁶¹	1999	<i>Mycobacterium avium-intracellulare</i>	AER: use of channel connectors provided by bronchoscope manufacturer rather than connector kit produced by AER manufacturer
Strelczyk ⁶²	1999	Acid-fast bacilli	AER: inadequate channel connectors provided by bronchoscope manufacturer
Wilson et al ⁶³	2000	<i>Aureobasidium</i> species	Reuse of single-use stopcocks disinfected by an AER
Larson et al ⁶⁴	2001	<i>M. tuberculosis</i>	AER: errors in cleaning, incompatible AER
Kressel and Kidd ⁹	2001	<i>M. chelonae</i> , <i>Methylobacterium mesophilicum</i>	AER: biofilm buildup in AER, no alcohol flush, organisms relatively resistant to glutaraldehyde

Abbreviations: AER, automatic endoscope reprocessor; CDC, Centers for Disease Control and Prevention.

* Species as listed by investigator; may not reflect current taxonomy.

- publication no. 99-1710. Hyattsville, MD: US Department of Health and Human Services, National Center for Health Statistics; 1998.
- Prakash UB. Advances in bronchoscopic procedures. *Chest* 1999;116:1403-1408.
 - American Society for Gastrointestinal Endoscopy. *Reprocessing of Flexible Gastrointestinal Endoscopes*. Manchester, MA: American Society for Gastrointestinal Endoscopy; 1995.
 - Spach DH, Silverstein FE, Stamm WE. Transmission of infection by gastrointestinal endoscopy and bronchoscopy. *Ann Intern Med* 1993;118:117-128.
 - Bond WW. Endoscopy reprocessing: problems and solutions. In: Rutala

- WA, ed. *Disinfection, Sterilization and Antisepsis in Health Care*. Champlain, NY: Polyscience Publications; 1998:151-163.
- Rutala WA, Weber DJ. Disinfection of endoscopes: review of new chemical sterilants used for high-level disinfection. *Infect Control Hosp Epidemiol* 1999;20:69-76.
 - Sorin M, Segal-Maurer S, Mariano N, Urban C, Combest A, Rahal JJ. Nosocomial transmission of imipenem-resistant *Pseudomonas aeruginosa* following bronchoscopy associated with an improper connection to the STERIS SYSTEM 1 processor. *Infect Control Hosp Epidemiol* 2001;22:409-413.
 - Kressel AB, Kidd F. A pseudo-outbreak of *Mycobacterium chelonae*

- and *Methylobacterium meophilicum* caused by contamination of an automated endoscope washer. *Infect Control Hosp Epidemiol* 2001;22:414-418.
10. Rutala WA. APIC guideline for selection and use of disinfectants. *Am J Infect Control* 1996;24:313-342.
 11. Alvarado CJ, Reichelderfer M, APIC Guidelines Committee. APIC guideline for infection prevention and control in flexible endoscopy. *Am J Infect Control* 2000;28:138-155.
 12. Alfa MJ, Sitter DL. In-hospital evaluation of ortho-phthalaldehyde as a high level disinfectant for flexible endoscopes. *J Hosp Infect* 1994;26:15-26.
 13. Roberts CG. Studies on the bioburden of medical devices and the importance of cleaning. In: Rutala WA, ed. *Disinfection, Sterilization and Antisepsis: Principles and Practices in Healthcare Facilities*. Champlain, NY: Polyscience Publications; 2001:63-69.
 14. Alfa MJ, DeGagne P, Olson N, Puchalski T. Comparison of ion plasma, vaporized hydrogen peroxide, and 100% ethylene oxide sterilizers to the 12/88 ethylene oxide gas sterilizer. *Infect Control Hosp Epidemiol* 1996;17:92-100.
 15. US Food and Drug Administration. Sterilants and high level disinfectants cleared by FDA in a 510(k) as of June 29, 2001, with general claims for processing reusable medical and dental devices. <http://www.fda.gov/cdrh/ode/germlab.html>. updated July 2, 2001.
 16. Rutala WA, Weber DJ. Infection control: the role of disinfection and sterilization. *J Hosp Infect* 1999;43(suppl):S43-S55.
 17. Cheung RJ, Ortiz D, Dimarino AJ Jr. GI endoscopic reprocessing practices in the United States. *Gastrointest Endosc* 1999;50:362-368.
 18. Alfa MJ, Olson N, DeGagne P, Hizon R. New low temperature sterilization technologies: microbiocidal activity and clinical efficacy. In: Rutala WA, ed. *Disinfection, Sterilization, and Antisepsis in Health Care*. Washington DC: Association for Professionals in Infection Control and Epidemiology; 1998.
 19. Rutala WA, Gergen MF, Weber DJ. Comparative evaluation of the sporicidal activity of new low-temperature sterilization technologies: ethylene oxide, 2 plasma sterilization systems, and liquid peracetic acid. *Am J Infect Control* 1998;26:393-398.
 20. Alfa MJ. Importance of lumen flow in liquid chemical sterilization (letter). *Am J Infect Control* 1999;27:373-374.
 21. Rutala WA, Gergen MF, Weber DJ. Importance of lumen flow in liquid chemical sterilization (reply). *Am J Infect Control* 1999;27:374-375.
 22. Jackson J, Leggett JE, Wilson D, Gilbert DN. *Mycobacterium gordonae* in fiberoptic bronchoscopes. *Am J Infect Control* 1996;24:19-23.
 23. Deva AK, Vickery K, Zou J, West RH, Harris JP, Cossart YE. Establishment of an in-use testing method for evaluating disinfection of surgical instruments using the duck hepatitis B model. *J Hosp Infect* 1996;33:119-130.
 24. Wu MS, Wang JT, Yang JC, Wang HH, Sheu JC, Chen DS, et al. Effective reduction of *Helicobacter pylori* infection after upper gastrointestinal endoscopy of mechanical washing of the endoscope. *Hepatology* 1996;43:1660-1664.
 25. Webb SF, Vall-Spinosa A. Outbreak of *Serratia marcescens* associated with the flexible bronchoscope. *Chest* 1975;68:703-708.
 26. Hussain SA. Fiberoptic bronchoscope-related outbreak of infection with *Pseudomonas*. *Chest* 1978;74:483.
 27. Markovitz A. Inoculation by bronchoscopy. *West J Med* 1979;131:550.
 28. Leers W-D. Disinfecting endoscopes: how not to transmit *Mycobacterium tuberculosis* by bronchoscopy. *Can Med Assoc J* 1980;123:275-280.
 29. Nelson KE, Larson PA, Schraufnagel DE, Jackson J. Transmission of tuberculosis by flexible fiberbronchoscopes. *Am Rev Respir Dis* 1983;127:97-100.
 30. Pappas SA, Schaaf DM, DiCostanzo MB, King FW, Sharp JT. Contamination of flexible fiberoptic bronchoscopes. *Chest* 1983;127:391-392.
 31. Wheeler PW, Lancaster D, Kaiser AB. Bronchopulmonary cross-colonization and infection related to mycobacterial contamination of suction valves of bronchoscopes. *J Infect Dis* 1989;159:954-958.
 32. Agerton T, Walway S, Gore B, Pozsik C, Plikaytis B, Woodley C, et al. Transmission of a highly drug-resistant strain (strain W1) of *Mycobacterium tuberculosis*. Community outbreak and nosocomial transmission via a contaminated bronchoscope. *JAMA* 1997;278:1073-1077.
 33. Blanc DS, Parret T, Janin B, Raselli P, Francioli P. Nosocomial infections and pseudo-infections from contaminated bronchoscopes: two-year follow up using molecular markers. *Infect Control Hosp Epidemiol* 1997;18:134-136.
 34. Michele TM, Cronin WA, Graham NM, Dwyer DM, Pope DS, Harrington S, et al. Transmission of *Mycobacterium tuberculosis* by a fiberoptic bronchoscope. Identification by DNA fingerprinting. *JAMA* 1997;278:1093-1095.
 35. Kramer MH, Krizek L, Gebel J, Kirsch A, Wegan E, Marklein G, et al. Bronchoscopic transmission of *Pseudomonas aeruginosa* due to a contaminated disinfectant solution from an automated dispenser unit. In: Final Program of the 11th Annual Scientific Meeting of the Society of Healthcare Epidemiology of America; Toronto, Ontario, Canada; April 1-3, 2001. Abstract 118.
 36. Weinstein HJ, Bone RC, Ruth WE. Contamination of a fiberoptic bronchoscope with a *Proteus* species. *Am Rev Respir Dis* 1977;116:541-543.
 37. Dawson DJ, Armstrong JG, Blacklock ZM. Mycobacterial cross-contamination of bronchoscopy specimens. *Am Rev Respir Dis* 1982;126:1095-1097.
 38. Sammartino MT, Israel RH, Magnussen CR. *Pseudomonas aeruginosa* contamination of fiberoptic bronchoscopes. *J Hosp Infect* 1982;3:65-71.
 39. Goldstein B, Abrutyn E. Pseudo-outbreak of *Bacillus* species: related to fiberoptic bronchoscopy. *J Hosp Infect* 1985;6:194-200.
 40. Siegmund-Igra Y, Inbar G, Campus A. A 'outbreak' of pulmonary pseudo-infection by *Serratia marcescens*. *J Hosp Infect* 1985;6:218-220.
 41. Richardson AJ, Rothburn MM, Roberts C. Pseudo-outbreak of *Bacillus* species: related to fiberoptic bronchoscopy. *J Hosp Infect* 1986;7:208-210.
 42. Hoffmann KK, Weber DJ, Rutala WA. Pseudoepidemic of *Rhodotorula rubra* in patients undergoing fiberoptic bronchoscopy. *Infect Control Hosp Epidemiol* 1989;10:511-514.
 43. Nye K, Chadha DK, Hodgkin P, Bradley C, Hancox J, Wise R. *Mycobacterium chelonae* isolation from broncho-alveolar lavage fluid and its practical implications. *J Hosp Infect* 1990;16:257-260.
 44. Centers for Disease Control and Prevention. Nosocomial infection and pseudo-infection from contaminated endoscopes and bronchoscopes—Wisconsin and Missouri. *MMWR* 1991;40:675-678.
 45. Fraser VJ, Jones M, Murray PR, Medoff G, Zhang Y, Wallace RJ. Contamination of flexible fiberoptic bronchoscopes with *Mycobacterium chelonae* linked to an automated bronchoscope disinfection machine. *Am Rev Respir Dis* 1992;145:853-855.
 46. Gubler JGH, Salfinger M, von Graevenitz A. Pseudoepidemic of nontuberculous mycobacteria due to a contaminated bronchoscope cleaning machine. *Chest* 1992;101:1245-1249.
 47. Nicolle LE, McLeod J, Romance L, Parker S, Paraskevas M. Pseudo-outbreak of blastomycosis associated with contaminated bronchoscopes. *Infect Control Hosp Epidemiol* 1992;13:324.
 48. Whitlock WL, Dietrich RA, Steimke EH, Tenholder MF. *Rhodotorula rubra* contamination in fiberoptic bronchoscopy. *Chest* 1992;102:1516-1519.
 49. Bryce EA, Walker M, Bevan C, Smith JA. Contamination of bronchoscopes with *Mycobacterium tuberculosis*. *Canadian Journal of Infection Control* 1993;8:35-36.
 50. Vandenberghe-Grauls CMJE, Baars ACM, Visser MR, Hulstaert PF, Verhoef J. An outbreak of *Serratia marcescens* traced to a contaminated bronchoscope. *J Hosp Infect* 1993;23:263-270.
 51. Bennett SN, Peterson DE, Johnson DR, Hall WN, Robinson-Dunn B, Dietrich S. Bronchoscopy-associated *Mycobacterium xenopi* pseudo-infections. *Am J Respir Crit Care Med* 1994;150:245-250.
 52. Campagnaro RI, Teichtahl H, Dwyer B. A pseudoepidemic of *Mycobacterium chelonae*: contamination of a bronchoscope and auto-cleaner. *Aust N Z J Med* 1994;24:693-695.
 53. Kolmos HJ, Lerche A, Kristoffersen K, Rosdahl VT. Pseudo-outbreak of *Pseudomonas aeruginosa* in HIV-infected patients undergoing fiberoptic bronchoscopy. *Scand J Infect Dis* 1994;26:653-657.
 54. Maloney S, Welbel S, Daves B, Adams K, Becker S, Bland L, et al. *Mycobacterium abscessus* pseudo-infection traced to an automated endoscope washer: utility of epidemiologic and laboratory investigation. *J Infect Dis* 1994;169:1166-1169.
 55. Peterson K, Bus N, Walter V, Chenoweth C. Pseudoepidemic of *Mycobacterium abscessus* associated with bronchoscopy. *Infect Control Hosp Epidemiol* 1994;15(suppl):P30. Abstract S32.
 56. Hagan ME, Klotz SA, Bartholomew W, Potter L, Nelson M. A pseudoepidemic of *Rhodotorula rubra*: a marker for microbial contamination of the bronchoscope. *Infect Control Hosp Epidemiol* 1995;16:727-728.
 57. Takigawa K, Fujita J, Negayama K, Terada S, Yamaji S, Kawanashi K, et al. Eradication of contaminating *Mycobacterium chelonae* from bronchoscopes and an automated bronchoscope disinfection machine. *Respir Med* 1995;89:423-427.
 58. Wang HC, Liaw YS, Yand PC, Kuo SH, Luh KT. A pseudoepidemic of *Mycobacterium chelonae* infection caused by contamination of a fiberoptic bronchoscope suction channel. *Eur Respir J* 1995;8:1259-1262.
 59. Mitchell DH, Hicks LJ, Chiew R, Montanaro JC, Chen SC. Pseudoepidemic of *Legionella pneumophila* serogroup 6 associated with contaminated bronchoscopes. *J Hosp Infect* 1997;37:19-23.
 60. Wallace RJ Jr, Brown BA, Griffith DE. Nosocomial outbreaks/pseudo-outbreaks caused by nontuberculous mycobacteria. *Annu Rev Microbiol* 1998;52:453-490.
 61. Centers for Disease Control and Prevention. Bronchoscopy-related infections and pseudo-infections—New York, 1996 and 1998. *MMWR* 1999;48:557-560.
 62. Strelczyk K. Pseudo-outbreak of acid-fast bacilli. *Am J Infect Control*

- 1999;27:18. Abstract.
63. Wilson SJ, Everts RJ, Kirkland KB, Sexton DJ. A pseudo-outbreak of *Aerobasidium* species lower respiratory tract infections caused by reuse of single-use stopcocks during bronchoscopy. *Infect Control Hosp Epidemiol* 2000;21:470-472.
 64. Larson J, Lambert L, Stricof R, Ridzon R, Navin T. *Mycobacterium tuberculosis* contamination and potential exposure from a bronchoscope, Pennsylvania—2000. In: Final Program of the 11th Annual Scientific Meeting of the Society for Healthcare Epidemiology of America; Toronto, Ontario, Canada; April 1-3, 2001.
 65. Mehta A, Minai OA. Infection control in the bronchoscopy suite. A review. *Clin Chest Med* 1999;20:19-32.
 66. Steere AC, Corrales J, von Graevenitz A. A cluster of *Mycobacterium gordonae* isolates from bronchoscopy specimens. *Am Rev Respir Dis* 1979;120:214-216.
 67. Schleupner CJ, Hamilton JR. A pseudoepidemic of pulmonary fungal infections related to fiberoptic bronchoscopy. *Infect Control* 1980;1:38-42.
 68. Cox R, deBorja K, Bach MC. A pseudo-outbreak of *Mycobacterium chelonae* infections related to bronchoscopy. *Infect Control Hosp Epidemiol* 1997;18:136-137.
 69. Southwick KL, Hoffmann K, Ferree K, Matthews J, Salfinger M. Cluster of tuberculosis cases in North Carolina: possible association with atomizer use. *Am J Infect Control* 2001;29:1-6.
 70. Weber DJ, Rutala WA. Occupational risks associated with the use of selected disinfectants and sterilants. In: Rutala WA, ed. *Disinfection, Sterilization and Antisepsis in Health Care*. Champlain, NY: Polyscience Publications; 1998:211-226.
 71. Bradley CR, Babb JR. Endoscope decontamination: automated vs. manual. *J Hosp Infect* 1995;30(suppl):537-542.
 72. Muscarella LF. Advantages and limitations of automatic flexible endoscope reprocessors. *Am J Infect Control* 1996;24:304-309.
 73. Muscarella LF. Automatic flexible endoscope reprocessors. *Gastrointest Endosc Clin N Am* 2000;10:245-257.
 74. Cooke RP, Rhyment-Morris A, Umasankar RS, Goddard SV. Bacteria-free water for automatic washer-disinfectors: an impossible dream? *J Hosp Infect* 1998;39:63-65.
 75. Lynch DA, Porter C, Murphy L, Axon AT. Evaluation of four commercial automatic endoscope washing machines. *Endoscopy* 1992;24:766-770.
 76. Jackson FW, Ball MD. Correction of deficiencies in flexible fiberoptic sigmoidoscope cleaning and disinfection technique in family practice and internal medicine offices. *Arch Intern Med* 1997;6:578-582.
 77. Orsi GB, Filocamo A, Di Stefano L, Tittobello A. Italian national survey of digestive endoscopy disinfection practices. *Endoscopy* 1997;29:732-738.
 78. Honeybourne D, Neumann CS. An audit of bronchoscopy practice in the United Kingdom: a survey of adherence to national guidelines. *Thorax* 1997;52:709-713.
 79. Food and Drug Administration, Centers for Disease Control and Prevention. FDA and CDC public health advisory: infections from endoscopes inadequately reprocessed by an automated endoscope reprocessing system. September 10, 1999. <http://fda.gov/cdrh/safety/endoreprocess.html>.
 80. Society of Gastrointestinal Nurses and Associates. Standards for infection control and reprocessing of flexible gastrointestinal endoscopes. *Gastroenterology Nursing* 2000;23:172-179.