Major Trends in the Microbial Etiology of Nosocomial Infection

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To determine trends in the microbial etiology of nosocomial infections in the 1980s, surveillance data on the microbiology of documented nosocomial infection reported to the National Nosocomial Infections Surveillance System and from the University of Michigan Hospital were analyzed. Antimicrobial susceptibility data on selected pathogens from both sources were also reviewed. Overall, Escherichia coli decreased from 23% of infections in 1980 to 16% in 1986–1989, Klebsiella pneumoniae dropped from 7% to 5%, whereas coagulase negative staphylococci increased from 4% to 9% and Candida albicans increased from 2% to 5%. Staphylococcus aureus, Pseudomonas aeruginosa, Enterobacter species and enterococci had minor increases, but antimicrobial resistant strains for these pathogens as well as coagulase-negative staphylococci were seen more frequently. In contrast to the 1970s, major shifts in the etiology of nosocomial infection have occurred in the decade of the 1980s. Taken as a whole, the shifts are away from more easily treated pathogens toward more resistant pathogens with fewer options for therapy. These shifts underscore the continued need for prevention and control to accompany new developments in therapy.

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Nosocomial infection remains a significant consequence of hospitalization. Estimates are that from 3% to 5% of patients leave the hospital having acquired infection, depending on case mix, hospital size, and multiple other factors [1]. Examination of the microbial etiology of these infections provides important information in day-to-day decision making in individual hospitals regarding potential outbreaks, unusual pathogens, antimicrobial resistance, and local trends in the etiology of infection. It is also useful periodically to examine trends in the etiology of infection over more prolonged time periods to detect shifts in the cause of infection. This report uses two sources of data, pooled data from participants in the National Nosocomial Infections Surveillance (NNIS) System of the Centers for Disease Control (CDC) and single hospital data from a large tertiary referral teaching hospital, the University of Michigan Hospital, to examine trends in etiology of hospitalacquired infection in the decade of the 1980s.

METHODS

The NNIS system was established in January 1970 and is currently the only nationwide source of information on nosocomial infections in the United States [2,3]. This report examines surveillance among NNIS hospitals performing hospital-wide surveillance from 1980 to 1989. Patients were monitored for all nosocomial infections at all body sites. Standardized definitions for infection are provided to the participating hospitals [4] but a variety of case-finding methods were used. Various methods for performing antimicrobial susceptibilities, including disk diffusion, macrobroth, and microdilution are used, but each hospital must provide evidence of proficiency in the method being used. Data from each hospital are submitted monthly to the CDC.

The percentage of coagulase-negative staphylococci that were reported to be resistant to methicillin, oxacillin, or nafcillin (MRSE) was determined by estimating the mean percentage for 1989 for all hospitals that reported data. Hospitals that did not report susceptibility results for at least 20 coagulase-negative staphylococci in the year were ex-

TABLE I

Pathogen Distribution for Major Sites of Nosocomial Infection, National Nosocomial Infections Surveillance System, 1986–1989

Pathogen	Urinary Tract Infection	Wound	Pneumonia	Bloodstream	Total
E. coli	11,135 (26)*	1,951 (10)	946 (6)	733 (6)	14,765 (16)
Enterococci	6,720 (16)	2,645 (13)	342 (2)	1,037 (8)	10,744 (12)
P. aeruginosa	5,127 (12)	1,668 (8)	2,598 (17)	543 (4)	9,936 (11)
S. aureus	823 (2)	3,439 (17)	2,401 (16)	1,984 (16)	8,647 (10)
Coagulase negative staphylococci	1,634 (4)	2,472 (12)	293 (2)	3,384 (27)	7,783 (9)
Enterobacter sp.	2,339 (6)	1,529 (8)	1,625 (11)	610 (5)	6,103 (7)
K. pneumoniae	2,664 (6)	618 (3)	1.042 (7)	548 (4)	4,872 (5)
C. albicans	2,978 (7)	481 (2)	615 (4)	617 (5)	4,691 (5)
P. mirabilis	2,312 (5)	712 (4)	503 (3)	105 (1)	3,632 (4)
Streptococcal species	207 (0)	539 (3)	231 (1)	465 (4)	1,442 (2)
Citrobacter sp.	812 (2)	321 (2)	226 (1)	82 (1)	1,441 (2)
Candida sp.	853 (2)	81 (0)	109(1)	330 (3)	1,373 (2)
S. marcescens	367 (1)	271 (1)	579 (4)	152 (1)	1,369 (2)

*Number (percent). Note: a site may have up to four pathogens.

cluded from the analysis. Changes in percent of MRSE between two consecutive years were estimated using only the data from hospitals that reported to NNIS in both years. In this way, the estimated change in percent MRSE over the period 1980–1989 was not biased by the effects of sample migration, i.e., hospitals joining and leaving NNIS.

The University of Michigan Hospital is a 588 bed adult medical and surgical tertiary care hospital. Separate hospitals house the obstetrics service, the neonatal unit, and pediatric patients, which are not included in the data in this report. Definitions for nosocomial infection identical to those in NNIS are used [4]. Antimicrobial susceptibilities are performed by microbroth dilution.

RESULTS

When the overall pathogen distribution for the four major sites of hospital-acquired infections reported to NNIS from 1986-1989 is examined, Escherichia coli remains the most common isolate (**Table I**). The other pathogens accounting for greater than 10% of infections are enterococci, Pseudomonas aeruginosa, and Staphylococcus aureus. These same four pathogens also dominate at the University of Michigan Hospital but the order differs slightly. E. coli also is most common, accounting for 23% of infections, but P. aeruginosa is second at 20% followed by the other two. Although E. coli still is the most frequent isolate, it has dropped in overall frequency comparing NNIS data from 1980 with 1986–1989 (**Figure 1**). Two of the top 10 pathogens have dropped in percentage, E. coli from 23% to 16% and Klebsiella pneumoniae from 7% to 5% of isolates. In contrast, coagulasenegative staphylococci have increased from 4% to 9%, almost all as a function of increase in bloodstream infections. Candida albicans also increased from 2% in 1980 to 5% in 1986-1989, with the increase occurring across all major sites of infection. Slight increases have occurred with *S. aureus*, *P. aeruginosa*, enterococci and *Enterobacter* species, with the overall change of 1–2% in each instance.

In addition to the overall changes noted, there have been changes as well within a given genus, in particular in antimicrobial resistance phenotype. For example, 1982 was the first full year of susceptibility data for cefotaxime at the University of Michigan with 91% of isolates of P. aeruginosa susceptible in contrast to only 65% in 1989 and 93% of Enterobacter cloacae susceptible in 1982 versus 76% in 1989. Similarly, 97% of P. aeruginosa were sensitive to gentamicin in 1982 versus 88% in 1989. Changes in phenotype in the grampositive nosocomial pathogens have been seen as well. At the University of Michigan, in 1980 less than 1% of S. aureus isolates were methicillinresistant, but in 1989 17% were resistant. In 1981, a single clinical isolate of high-level gentamicinresistant enterococci was identified in our hospital. In 1989, 20% of all enterococcal isolates displayed this phenotype. Currently, more than 60% of coagulase-negative staphylococci at the University hospital are methicillin-resistant. This increasing resistance is also reflected in the pooled data from NNIS (Figure 2).

COMMENTS

Surveillance, including bacteriologic surveillance to provide background information in order to recognize nosocomial infection problems requiring control measures or interventions, remains important to current infection control efforts in hospitals. As has been noted by Stamm and coworkers [5], the bacteriology of epidemic nosocomial infection is dramatically different from endemic infection. The trends noted in this report pool bacteriology from both types of infections but endemic infections overwhelm any impact of out-

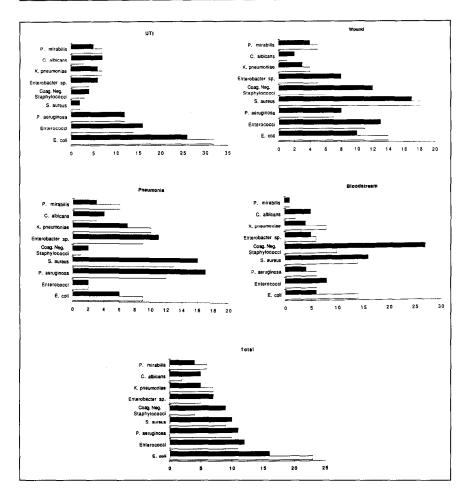


Figure 1. Comparative percentage of pathogen distribution for nosocomial infections, National Nosocomial Infections Surveillance System, 1980 (white bars) versus 1986-89 (black bars). UTI = urinary tract infection.

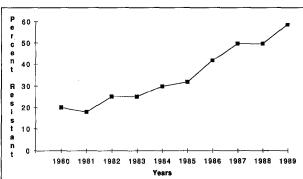


Figure 2. Proportion of coagulase-negative staphylococci from the National Nosocomial Infections Surveillance System reported as resistant to methicillin, oxacillin, or nafcillin, 1980-1989 (n = 27,150).

breaks. The changes noted thus appear to be gradual shifts in the pathogens isolated from hospital-acquired infection. It should be noted that these shifts are in contrast to similar analysis over the last decade where no secular or periodic trends were found [6].

One explanation for the shifts noted would be selective prevention of infections due to E. coli and K. pneumoniae. These organisms do share in common a similar reservoir, the gastrointestinal tract, and endogenous acquisition as mode of infection. Antimicrobial prophylaxis or therapy with agents active against these organisms but not more resistant organisms, such as Enterobacter species, if used widely might explain the shift away from them as causes of infection. An antimicrobial group with such a spectrum is the so-called firstgeneration cephalosporins, which are often chosen as prophylaxis for a variety of procedures in hospitals, but whether this explains the observed shift away from *Klebsiella* and *E. coli* is speculation.

Another major change is the increasing importance of coagulase-negative staphylococci. This change is almost entirely due to the increase in bloodstream infection. This may in part be due to the growing appreciation of this organism as a pathogen [7] and thus a willingness to report positive cultures as true infections. It also may in part be due to the increasing reliance on a variety of devices, especially intravascular devices, to aid in the care of seriously ill hospital patients. A report by Gaynes et al [2] examines this secular trend.

The most disturbing trend occurring in the 1980s was a move away from more susceptible toward more resistant pathogens. This occurred both between and within genera. Genera that tend to be more susceptible to antimicrobials, Proteus mirabilis, E. coli, and K. pneumoniae, all decreased in prevalence, whereas Enterobacter, Pseudomonas, Enterococcus, and Candida increased. Within genera, traditional etiologies in some instances changed little, but phenotype did. Several studies have already pointed out the growing problem of methicillin resistance in S. aureus [8,9] and similar changes in coagulase-negative staphylococci were found in this analysis. The abrupt increase in the proportion of coagulase-negative staphylococci resistant to semisynthetic penicillins in 1986 is likely the result of changes in recommended testing methodologies [10], but the trend upward was clear both before and after these recommendations. High-level gentamicin-resistant enterococci are a new and prominent nosocomial infection in some hospitals [11], and beta-lactamase production and glycopeptide resistance are also being encountered in this pathogen. More frequent isolation of aminoglycoside-resistant gram-negative pathogens was hinted at in the review of data from the 1970s [6] and a similar problem with the new extended spectrum beta-lactams is apparent from the limited data presented here. Overall, the remarkable ability of nosocomial bacterial pathogens to persist in the hospital environment despite new therapies continues to present important challenges to infection control. More than ever, prevention rather than treatment after occurrence of

nosocomial infection should remain a primary goal.

REFERENCES

- 1. Haley RW, Culver DH, White JW, Morgan WM, Emori TG. The nationwide nosocomial infection rate: a new need for vital statistics. Am J Epidemiol 1985; 121: 159–66.
- 2. Gaynes RP, Culver DH, Emori TG, et al. The National Nosocomial Infections Surveillance System; plans for the 1990s and beyond. Am J Med 1991; 91(Suppl 3B): 116–120.
- **3.** Emori TG, Culver DH, Horan TC, *et al.* National Nosocomial Infections Surveillance System (NNIS): description of surveillance methods. Am J Infect Control 1991; 19: 19–35.
- 4. Garner JS, Jarvis WR, Emori TG, Horan TC, Hughes JM. CDC definitions for nosocomial infections, 1988. Am J Infect Control 1988; 16: 128–40.
- 5. Stamm WE, Weinstein RA, Dixon RS. Comparison of endemic and epidemic nosocomial infections. Am J Med 1981; 70: 393–7.
- **6.** Allen JR, Hightower AW, Martin SM, Dixon RE. Secular trends in nosocomial infections: 1970–79. Am J Med 1981; 70: 389–92.
- 7. Hamory BH, Parisi JT. Staphylococcus epidermidis: a significant nosocomial pathogen. Am J Infect Control 1987; 15: 59–74.
- **8.** Haley RW, Hightower AW, Khabbaz RF. The emergence of methicillin-resistant Staphylococcus aureus infections in United States hospitals. Possible role of the housestaff-patient transfer circuit. Ann Intern Med 1982; 97: 297–308.
- Boyce JM, Causey WA. Increasing occurrence of methicillin-resistant Staphylococcus aureus in the United States. Infect Control 1982; 3: 377–83.
- 10. National Committee for Clinical Laboratory Standards. Methods for dilution-antimicrobial susceptibility tests for bacteria that grow aerobically. Approved Standard M7-A. National Committee for Clinical Laboratory Standards, Villanova, Pa., 1985.
- 11. Zervos M, Dembinski S, Mikesell T, Schaberg DR. High-level gentamicin resistance in Streptococcus faecalis: risk factors and evidence for exogenous acquisition. J Infect Dis 1986; 153: 1075–83.