



University of Pennsylvania
ScholarlyCommons

Cartographic Modeling Lab Papers

Cartographic Modeling Lab

1-1-2004

Hospital-Reported Pneumococcal Susceptibility to Penicillin

Joshua P. Metlay

Charles Branas

University of Pennsylvania, cbranas@upenn.edu

Neil O. Fishman

University of Pennsylvania, neil.fishman@uphs.upenn.edu

Follow this and additional works at: http://repository.upenn.edu/cml_papers

 Part of the [Medicine and Health Sciences Commons](#)

Metlay, Joshua P.; Branas, Charles; and Fishman, Neil O., "Hospital-Reported Pneumococcal Susceptibility to Penicillin" (2004).
Cartographic Modeling Lab Papers. 4.
http://repository.upenn.edu/cml_papers/4

Metlay, J. P., Branas, C. C., & Fishman, N. O. (2004). Hospital-reported pneumococcal susceptibility to penicillin. *Emerging Infectious Diseases [serial online]*, 10(1), 54-59. doi: 10.3201/eid1001.030140
Available from: <http://wwwnc.cdc.gov/eid/article/10/1/03-0140.htm>

This paper is posted at ScholarlyCommons. http://repository.upenn.edu/cml_papers/4
For more information, please contact libraryrepository@pobox.upenn.edu.

Hospital-Reported Pneumococcal Susceptibility to Penicillin

Abstract

Geographic variation in drug susceptibility among isolates of *Streptococcus pneumoniae* has influenced national treatment guidelines for community-acquired pneumonia. Whether individual hospital susceptibility data provide reliable and valid information for providers is unclear. We examined the geographic and temporal variability in hospital-reported rates of pneumococcal susceptibility. We surveyed all 52 hospitals that provided acute adult care in the five counties surrounding Philadelphia and collected data on levels of penicillin susceptibility among all pneumococcal blood isolates from 1998 to 2000. In 1998, pneumococcal nonsusceptibility to penicillin varied from 0% to 67% of all blood isolates across the 33 hospitals with ≥ 10 isolates in that year. Hospital location did not correlate with the level of reported pneumococcal susceptibility ($p = 0.8$). In addition, correlations were not significant in reported pneumococcal susceptibility to penicillin within individual hospitals during the 3 years.

Disciplines

Medicine and Health Sciences

Comments

Metlay, J. P., Branas, C. C., & Fishman, N. O. (2004). Hospital-reported pneumococcal susceptibility to penicillin. *Emerging Infectious Diseases [serial online]*, 10(1), 54-59. doi: 10.3201/eid1001.030140

Available from: <http://wwwnc.cdc.gov/eid/article/10/1/03-0140.htm>

Hospital-reported Pneumococcal Susceptibility to Penicillin¹

Joshua P. Metlay,* Charles C. Branas,† and Neil O. Fishman†

Geographic variation in drug susceptibility among isolates of *Streptococcus pneumoniae* has influenced national treatment guidelines for community-acquired pneumonia. Whether individual hospital susceptibility data provide reliable and valid information for providers is unclear. We examined the geographic and temporal variability in hospital-reported rates of pneumococcal susceptibility. We surveyed all 52 hospitals that provided acute adult care in the five counties surrounding Philadelphia and collected data on levels of penicillin susceptibility among all pneumococcal blood isolates from 1998 to 2000. In 1998, pneumococcal nonsusceptibility to penicillin varied from 0% to 67% of all blood isolates across the 33 hospitals with ≥ 10 isolates in that year. Hospital location did not correlate with the level of reported pneumococcal susceptibility ($p = 0.8$). In addition, correlations were not significant in reported pneumococcal susceptibility to penicillin within individual hospitals during the 3 years.

Antimicrobial drug treatment of patients with community-acquired pneumonia is largely empirical because the pathogen is not frequently identified. Even when a bacterial pathogen is identified, antimicrobial drug susceptibility information is frequently delayed for a few days, necessitating empiric treatment decisions. Guidelines for empirical treatment have been provided by several organizations, emphasizing the selection of agents based on the probability of specific pathogens in different clinical settings (1–3). One consistent recommendation of all guidelines for all clinical settings is that antimicrobial therapy cover *Streptococcus pneumoniae* because it is the most common bacterial pathogen and among the most virulent.

A number of factors guide the selection of empirical therapy for patients with community-acquired pneumonia, including drug efficacy, side effect profile, and ease of administration. In addition, the recent emergence of antimicrobial drug resistance among clinical isolates of *S. pneumoniae* has raised considerable concern that a number of drugs historically endorsed by national treatment guide-

lines may no longer be effective. The recommendations to avoid β -lactams and macrolides in high-risk settings has not been preceded by clinical studies demonstrating that drug resistance in vitro translates into clinical treatment failures (4). Regardless, emerging resistance to penicillins, tetracyclines, and macrolides has prompted newer versions of treatment guidelines to recommend abandoning these therapies in favor of newer therapies, such as fluoroquinolones, in settings where the risk for a drug-resistant pneumococcal infection is high. Given that penicillin-resistant *S. pneumoniae* are increasingly multidrug resistant (5), if the risk for penicillin-resistant *S. pneumoniae* is high, the risk for resistance to macrolides and tetracyclines is correspondingly high.

Specific risk factors for infection with penicillin-resistant *S. pneumoniae* have not been clearly identified. Individual risk factors include prior exposure to antibiotics and exposure to young children in daycare (6), although no specific rules guide the interpretation of these risk factors. In addition, awareness of substantial geographic variation in the proportion of drug-resistant *S. pneumoniae* has led some groups to recommend consideration of regional susceptibility patterns in the choice of empirical therapy for community-acquired pneumonia (2). Unfortunately, routine isolation of *S. pneumoniae* is highly dependent on the clinical setting. For example, blood and sputum samples are rarely sent from outpatient settings (7); thus, local rates of pneumococcal drug resistance are primarily determined by the phenotypes of bacteria isolated from sicker patients who require hospitalization. Whether results from local hospital microbiology laboratories provide valid information for guiding treatment is unclear.

The specific aims of this study were to compare the rates of penicillin resistance among pneumococcal blood isolates across all acute-care hospitals within a five-county region of eastern Pennsylvania. We examined the relationship between pneumococcal resistance, rates of pneu-

*VA Medical Center, Philadelphia, Pennsylvania, USA; and
†University of Pennsylvania, Philadelphia, Pennsylvania, USA

¹Presented in part at the 3rd International Symposium on Pneumococci and Pneumococcal Disease, Anchorage, Alaska, May 2002.

mococcal isolation, and geographic location to determine the reliability of individual hospital rates for guiding treatment decisions. Our underlying hypothesis was that individual hospital rates of penicillin resistance among clinical isolates of *S. pneumoniae* would be poorly correlated across time and space, emphasizing the inherent dangers in using individual hospital rates to guide empirical treatment decisions in the community.

Methods

Design

This study was a cross-sectional study of penicillin susceptibility among clinical blood isolates reported at all acute-care hospitals in the Delaware Valley Case Control Network (DVCCN). DVCCN is a population-based network of all adult acute-care hospitals in the five Pennsylvania counties around the metropolitan Philadelphia region. These counties (Bucks, Montgomery, Chester, Delaware, and Philadelphia) contain 52 adult acute-care hospitals serving a population of 3.4 million residents.

Data Collection

Data were collected during two periods. During the first data collection, microbiology laboratory supervisors were contacted by telephone at each area hospital during January 1999 to collect data on all pneumococcal blood isolates during the preceding year. Local laboratory personnel reported cumulative numbers of pneumococcal blood isolates and the proportion of isolates that were penicillin nonsusceptible for the 12-month period. When available, laboratories separately reported susceptibility percentiles as susceptible, intermediate susceptible, and resistant; otherwise, the laboratories reported the combined percentile nonsusceptible (intermediate plus resistant). We collected information on unique isolates but did not confirm that these represented unique episodes of patient infections. However, we think hospitals generally reported only one isolate per patient because guidelines recommend that hospitals report cumulative susceptibility data on the basis of single isolates per patient per reporting period (8).

A second data collection was conducted in early 2001. The same hospital laboratories were contacted by mail with follow-up telephone calls requesting the same information on cumulative numbers of pneumococcal blood isolates and the proportion of isolates that were penicillin nonsusceptible for the 12-month period. Data were requested for 1999 and 2000 separately, if available.

Analysis

Annual number of pneumococcal blood isolates and

penicillin-susceptibility rates were summarized for each hospital for each year. Standard National Committee for Clinical Laboratory Standards (NCCLS) criteria define three categories of penicillin susceptibility for *S. pneumoniae*: susceptible (MIC <0.1 µg/mL), intermediate susceptible (MIC 0.1–1.0 µg/mL) and resistant (MIC ≥2.0 µg/mL) (9). Penicillin-nonsusceptible isolates are defined as intermediate susceptible and resistant isolates combined. For all analyses, we examined the proportion of isolates that were nonsusceptible. We specifically excluded any hospital site reporting <10 isolates per year as too unreliable to provide point estimates of resistance rates. This level is consistent with NCCLS guidelines recommending hospital laboratories report cumulative susceptibility data only for organisms for which there is a minimum of 10 isolates per reporting period (8). Descriptive statistics were developed for the distributions of susceptibilities and annual pneumococcal isolation rates. Similar to prior studies of hospital-level variation (10,11), we report the proportion of hospitals with >5% deviation from the overall regional mean rate of resistance because lower levels of deviation are unlikely to influence prescribing decisions.

To analyze the reliability of individual hospital rates of pneumococcal resistance, we examined the correlation of rates across each of the 3 years of data collection. We restricted these analyses to those hospitals reporting ≥10 pneumococcal blood isolates in 1998, 1999, and 2000. Correlations between any 2 years were calculated with Spearman's correlation coefficient, testing the assumption that pneumococcal susceptibility levels at individual hospitals should be correlated year to year, regardless of whether overall susceptibility levels for the region were rising, falling, or remaining constant. In addition, we calculated an intra-class correlation coefficient across all 3 years of data at the hospital level by using a general linear model for analysis of variance (12).

Next, we tested the assumption that the proportion of nonsusceptible *S. pneumoniae* at each hospital should demonstrate an underlying geographic clustering. We tested the geographic clustering of pneumococcal resistance rates with two methods. First, we analyzed whether the location of each hospital at the county level was associated with the proportion of nonsusceptible pneumococci by using the nonparametric Kruskal-Wallis test. Second, we conducted Geographic Information System (GIS) mapping to visually judge the spatial distribution of hospital susceptibility rates. Hospitals were geographically assigned longitude and latitude coordinates on the basis of the hospital address by using GIS mapping software (Mapitude, Caliper Corporation, Newton, MA) to display the proportion of nonsusceptible pneumococci at each hospital location on the map. Using data from 1998 only, we graphical-

ly represented the proportion of nonsusceptible pneumococci as a proportional symbol map (bubble plot), where each hospital location is represented by a circle whose radius is proportional to the level of penicillin nonsusceptibility among pneumococcal blood isolates.

We tested the null hypothesis that no spatial autocorrelation of the nonsusceptibility rates occurred with Moran's I statistic, based on a proximity matrix calculated from the city-block distances between the longitude-latitude coordinate points of hospital addresses. Moran's I is a spatial cross-product coefficient that is interpreted similarly to the Pearson correlation coefficient (13,14).

Results

For the survey of 1998 susceptibility results, we received responses from 47 of 52 hospitals in the DVCCN. However, in two instances, the results from multiple hospitals were combined at a single laboratory and could not be separated for reporting purposes. Of the 45 laboratories reporting results, 33 laboratories reported ≥ 10 pneumococcal blood isolates during 1998. Among the 33 sites, the median number of pneumococcal blood isolates during 1998 was 19 (range 10-92).

Figure 1 displays the frequency distribution of the proportion of isolates with nonsusceptibility to penicillin as reported by each hospital in 1998. The proportion of nonsusceptible isolates ranged from 0% to 67%, with a mean proportion of 21%. Sixty-one percent of hospitals were $>5\%$ above or below the mean. If we only included hospitals with ≥ 20 pneumococcal isolates in 1998, a total of 16 hospitals would be included in the analysis; the proportion of nonsusceptible isolates ranged from 0% to 36%, with a mean proportion of 20%. Fifty-six percent of hospitals were $>5\%$ above or below the mean proportion of nonsusceptible isolates.

For the 1999 and 2000 survey results, we received responses from 31 hospital laboratories. Of these, 23 hospital laboratories reported ≥ 10 pneumococcal isolates in 1 of the 2 study years. The mean proportion of isolates with reduced susceptibility to penicillin was 19% in 1999 and 24% in 2000. Fifty-three percent of sites were $>5\%$ above or below the mean in 1999, and 65% of sites were $>5\%$ above or below the mean in 2000.

The Table summarizes the relationship between each hospital's annual proportion of nonsusceptible pneumococcal blood isolates during the 3 years of the study. For hospitals reporting ≥ 10 isolates in 1998, 1999, and 2000 ($N = 15$), the proportion of nonsusceptible isolates at each hospital was poorly correlated between any 2 years. For 1998 to 1999, the Spearman correlation coefficient was -0.07 ($p = 0.82$), for 1999 to 2000, the Spearman correlation coefficient was 0.28 ($p = 0.32$), and for 1998 to 2000, the Spearman correlation coefficient was -0.03 ($p = 0.91$).

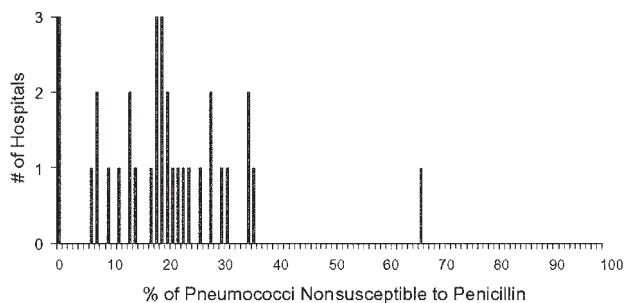


Figure 1. Proportion of penicillin-nonsusceptible pneumococcal bloodstream infections at hospitals in the Delaware Valley. The number of hospitals with each reported level of penicillin nonsusceptibility among all pneumococcal bloodstream isolates at each hospital in 1998 are shown. Penicillin nonsusceptibility was defined as any isolate with a penicillin MIC >0.1 $\mu\text{g/mL}$. Hospitals with <10 isolates in 1998 were excluded.

Across all 3 years, the overall intraclass correlation coefficient was 0.18.

Location of hospital by county did not correlate with the proportion of pneumococcal blood isolates nonsusceptible to penicillin ($p = 0.45, 0.37, 0.08$ in 1998, 1999, and 2000, respectively). Figure 2 is a proportional symbol map with each hospital location represented by a circle whose radius corresponds to the proportion of nonsusceptible pneumococcal blood isolates at each hospital in 1998. Overall, Figure 2 shows geographic clustering of the 33 hospitals reporting ≥ 10 pneumococcal blood isolates in 1998, 14 of which are gathered in Philadelphia, the most heavily populated county in our five-county study area. No geographic clustering in the proportion of nonsusceptible pneumococci isolated at each hospital was evident from our data. This finding is supported by a Moran correlation coefficient of -0.01 , demonstrating a lack of statistically significant spatial autocorrelation in the proportion of nonsusceptible hospital pneumococci at hospitals across the five-county region ($p = 0.80$).

Discussion

Pneumococcal drug resistance has increased at a fast rate. Most resistant isolates now demonstrate reduced susceptibility to multiple antimicrobial drug classes, including β -lactams, macrolides, and tetracyclines. However, substantial geographic variability in the proportion of pneumococci resistant to different antimicrobial drugs at the international and national levels suggests that the impact of resistance on empirical treatment decisions may vary across regions. Indeed, at least one guideline recommends that the selection of empirical therapy for outpatients with community-acquired pneumonia should be influenced by regional antimicrobial drug susceptibility patterns for *S. pneumoniae* (2).

For most physicians, the obvious source of information

Table. Annual variation in proportion of penicillin-nonsusceptible isolates at hospitals in the Delaware Valley

Site	Annual no. blood isolates			% penicillin nonsusceptible		
	1998	1999	2000	1998	1999	2000
A	25	21	26	0	33	27
B	12	15	17	0	13	29
C	29	25	29	24	20	28
D	14	18	12	14	22	8
E	44	39	28	18	23	25
F	27	19	20	19	21	20
G	26	56	35	19	23	17
H	62	15	22	19	0	27
I	10	13	12	20	38	50
J	44	15	32	20	33	16
K	92	45	87	22	4	8
L	35	31	30	29	16	20
M	41	36	34	29	14	38
N	33	32	35	36	13	9
O	19	15	19	37	40	47

on community-acquired pneumonia is the microbiologic susceptibility results from their local hospital laboratory. Hospitals frequently provide this information in the form of antibiograms, specifically designed to aid in antimicrobial drug selection. The validity of this information is largely unknown. This study demonstrates that substantial variability exists in hospital-reported rates of pneumococcal drug resistance over a small geographic region and that the rates at each site are poorly correlated over time. In addition, using a variety of approaches, we were unable to demonstrate substantial geographic clustering of the data at the individual hospital level, suggesting that the individual hospital rates poorly reflect an underlying rate of resistance in the community.

Other studies have demonstrated similar heterogeneity in the proportion of drug-resistant pneumococci reported by hospitals over small geographic areas. The Centers for Disease Control and Prevention reported levels of pneumococcal resistance to penicillin among invasive isolates from all hospitals within the seven regions in the United States that comprise their Active Bacterial Core Surveillance program. The proportion of penicillin-nonsusceptible *S. pneumoniae* ranged from 15.3% to 38.3% across the seven regions. The proportion of hospitals within each region that were >5% below or above the average for the region ranged from 35% to 76%. No hospital characteristics, including proportion of isolates from children or from black patients, predicted deviation from the regional average (11). More recently, 16 hospitals in Brooklyn, New York, participated in boroughwide surveillance for *S. pneumoniae* in 1997 and 1999. Aggregating the hospitals into the western and eastern ends of the borough demonstrated a significant difference in the proportion of penicillin-susceptible isolates, 57% versus 75% ($p = 0.046$) (15).

Our hypothesis that individual hospital rates of drug-resistant *S. pneumoniae* should demonstrate geographic

clustering is based on the assumption that rates of resistance should reflect the underlying distribution of drug resistance in the source community. Prior research has demonstrated that patients with community-acquired pneumonia are hospitalized on average <5 miles from their place of residence (16). Thus, hospitals within the same geographic region should admit patients with similar underlying rates of drug resistance. Moreover, if the pneumococcal resistance rates reported by individual hospitals should be used to guide physician antibiotic prescribing decisions for patients with community-acquired pneumococcal infections, hospitals serving similar communities should report similar rates of resistance.



Figure 2. Geographic distribution of penicillin nonsusceptibility among pneumococcal isolates at 33 hospitals in the Delaware Valley in 1998. The figure is a proportional symbol map (bubble plot). Each hospital location is represented by a circle with an H in the center at the corresponding longitude and latitude of the hospital. The radius of the circle is directly proportional to the proportion of penicillin-nonsusceptible pneumococci at each hospital in 1998. The range is 0% to 67%. Hospitals with <10 isolates in 1998 were excluded. An insert magnifies the geographic distribution of hospitals clustered in the center of Philadelphia.

Beyond reflecting true variability in the underlying community levels of drug resistance, hospital-level variation in the proportion of penicillin-susceptible pneumococcal isolates reflects at least four phenomena. First, chance error can create significant variability in hospital-level results particularly since many hospitals have only a small number of invasive pneumococcal isolates per year. We excluded hospitals with <10 isolates in each study year from our analyses to reduce the role of chance in our findings, but the relatively small number of isolates at each site undoubtedly contributed to the significant year-to-year variability and lack of geographic clustering. However, a minimum of 10 isolates per year is the NCCLS-recommended minimum number of isolates for reporting cumulative antimicrobial susceptibility results for any species in any reporting period (8).

Second, variation in testing strategies at individual hospitals may create bias in the proportion of isolates identified as nonsusceptible if the thresholds for sending microbiologic tests vary according to characteristics that are likely to influence the probability of a drug-resistant infection. While this may be an important determinant of variability in drug-susceptibility rates among clinical isolates from respiratory and sinus samples, we believe that it is less likely to explain variability in susceptibility rates among blood isolates since thresholds for sending blood cultures are more standardized across clinical practices, particularly in the management of patients hospitalized with community-acquired pneumonia.

Third, hospitals may differ in their methodologic approach and quality assurance for pneumococcal susceptibility testing. However, a study based on a nationwide College of American Pathologists proficiency survey of hospital laboratories demonstrated that, while the use of specific susceptibility tests varies substantially, few major interpretive errors occurred in assigning the susceptibility to standardized pneumococcal isolates (17). In addition, prior studies have found that the results of susceptibility data generated at local hospital facilities provide a reasonable surrogate for susceptibility data generated at centralized facilities (18).

Finally, geographic proximity of hospitals does not ensure that the patient populations served by those hospitals come from the same communities. Indeed, substantial literature has established important variation in hospital referral patterns that creates diversity in the source of patient populations among geographically proximate hospitals. Future research will need to consider the role of referral patterns in explaining some of the hospital-level variability observed in this and other studies.

One limitation of this study is our focus on a single geographic region in eastern Pennsylvania. Patterns and determinants of hospital-level variation in pneumococcal drug

susceptibility may produce different results in other regions. In addition, because we focused on hospitals providing adult care, we cannot generalize our results to pediatric hospitals.

Regardless of whether individual hospitals provide valid information on local levels of pneumococcal susceptibility, variability in these levels is only clinically meaningful if the range of susceptibility crosses some threshold for the empirical choice of specific antimicrobial drugs. Currently in the United States, empirical use of penicillins for the treatment of community-acquired pneumonia is uncommon relative to the use of fluoroquinolones and macrolides (19). However, as resistance to these classes of antimicrobial drugs continues to grow, physicians will need to determine appropriate thresholds for switching to yet newer antimicrobial agents. In these settings, valid information on local rates of pneumococcal drug susceptibility will become increasingly important. Ideally, data should be derived from large samples reflective of the region for which the empirical recommendations apply. Continued research is needed to determine whether susceptibility data provided by local hospital microbiology laboratories can ever serve this purpose.

Acknowledgments

The authors acknowledge the valuable contributions of Emily Cohen, Linda Crossette, M. Annette Hill, and Tonya Mincer for their help in the collection of data and preparation of this manuscript; the following laboratory directors and their staff members whose participation and continued support made this research possible: Herbert Auerbach, Ierachmiel Daskal, Stephen B. Chasko, Albert Keshgegian, Olarae Giger, David Wright, Ru Lin Ko Tung, Jim Heald, Harvey B. Spector, Lawrence M. Matthews, Rose M. Kenny, Xiaoli Chen, Ila Mirchandani, Howard Elefant, Fernando Garcia, Irwin Hollander, Christopher Emory, Pantaleon Fagel, Paul Edelstein, D.W. Anderson, Harvey J. Bellin, Paul Belser, Gary Stopyra, Leonas Bekeris, Dante DiMarzio, Susan Yaron, Pradeep Bhagat, James Monihan, John McCormick, David Steinberg, Zenon Gibas, Helen Kroh, Allan Truant, Donald Junkind, Donald Stieritz, and Manjula Balasubramanian.

The following hospitals participated in this study: Abington Memorial Hospital, Albert Einstein Medical Center, Brandywine Hospital, Bryn Mawr Hospital, Lankenau Hospital, Paoli Hospital, Central Montgomery Medical Center, Chestnut Hill Hospital, Chester County Hospital, Crozer Chester Medical Center, Taylor Hospital, Springfield Hospital, Delaware County Memorial Hospital, Doylestown Hospital, Elkins Park Hospital, Jeanes Hospital, Fox Chase Cancer Center, Frankford Hospital Torresdale Division, Frankford Hospital Bucks County Campus, Frankford Hospital Frankford Division, Graduate Hospital, Parkview Hospital, Grand View Hospital, Hahnemann University

Hospital, Medical College of Pennsylvania, Holy Redeemer Hospital and Medical Center, Hospital of the University of Pennsylvania, Presbyterian Hospital, Mercy Fitzgerald, Mercy Suburban, Mercy Hospital of Philadelphia, Methodist Hospital, Montgomery Hospital, Pennsylvania Hospital, Phoenixville Hospital, Pottstown Memorial Hospital, Riddle Memorial Hospital, Roxborough Memorial Hospital, Jennersville Regional Medical Center, St. Agnes Medical Center, St. Luke's Bethlehem, St. Mary Medical Center, Temple University Hospital, Northeastern Hospital, Episcopal Hospital, Thomas Jefferson University Hospital, Philadelphia VA Medical Center, and Warminster Hospital.

J.P.M. is supported by an Advanced Research Career Development Award from the Health Services Research and Development Service of the Department of Veterans Affairs. This research was partly supported by grant R01-AI46645 from the National Institutes of Health.

Dr. Metlay is a research associate and staff physician at the VA Medical Center in Philadelphia. He is an assistant professor of medicine and epidemiology at the University of Pennsylvania School of Medicine. His research focuses on the epidemiology of drug-resistant pneumococcal infections, particularly among adults with community-acquired pneumonia.

References

- Niederman MS, Mandell LA, Anzueto A, Bass JB, Broughton WA, Campbell GD, et al. Guidelines for the management of adults with community-acquired pneumonia. Diagnosis, assessment of severity, antimicrobial therapy, and prevention. *Am J Respir Crit Care Med* 2001;163:1730–54.
- Bartlett JG, Dowell SF, Mandell LA, File TM Jr, Musher DM, Fine MJ. Practice guidelines for the management of community-acquired pneumonia in adults. *Infectious Diseases Society of America. Clin Infect Dis* 2000;31:347–82.
- Mandell LA, Marrie TJ, Grossman RF, Chow AW, Hyland RH. Canadian guidelines for the initial management of community-acquired pneumonia: an evidence-based update by the Canadian Infectious Diseases Society and the Canadian Thoracic Society. The Canadian Community-Acquired Pneumonia Working Group. *Clin Infect Dis* 2000;31:383–421.
- Metlay JP, Singer DE. Outcomes in lower respiratory tract infections and the impact of antimicrobial drug resistance. *Clin Microbiol Infect* 2002;8(Suppl 2):1–11.
- Whitney CG, Farley MM, Hadler J, Harrison LH, Levau C, Reingold A, et al. Increasing prevalence of multidrug-resistant *Streptococcus pneumoniae* in the United States. *N Engl J Med* 2000;343:1917–24.
- Dowell SF, Schwartz B. Resistant pneumococci: protecting patients through judicious use of antibiotics. *Am Fam Physician* 1997;55:1647–54, 1657–8.
- Fine MJ, Stone RA, Singer DE, Coley CM, Marrie TJ, Lave JR, et al. Processes and outcomes of care for patients with community-acquired pneumonia: results from the Pneumonia Patient Outcomes Research Team (PORT) cohort study. *Arch Intern Med* 1999;159:970–80.
- National Committee for Clinical Laboratory Standards. Analysis and presentation of cumulative antimicrobial susceptibility test data; Approved guidelines. 2002. Report no.: NCCLS document M39-A. Philadelphia: The Committee; 2002.
- National Committee for Clinical Laboratory Standards. Performance standards for antimicrobial susceptibility testing: 12th information supplement. 2002. Report no.: NCCLS document M100-S12. Philadelphia: The Committee; 2002.
- Schrag SJ, Zell ER, Schuchat A, Whitney CG. Sentinel surveillance: a reliable way to track antibiotic resistance in communities? *Emerg Infect Dis* 2002;8:496–502.
- Geographic variation in penicillin resistance in *Streptococcus pneumoniae*—selected sites, United States, 1997. *MMWR Morb Mortal Wkly Rep* 1999;48:656–61.
- Fleiss J. The design and analysis of clinical experiments. New York: John Wiley and Sons; 1986.
- Fotheringham A, Brunsdon C, Charlton M. Statistical inference for spatial data. In: Quantitative geography: perspectives on spatial data analysis. London: Sage Publications; 2000. p. 201–11.
- Cliff A, Ord J. Spatial autocorrelation: monographs in spatial and environmental systems analysis. London: Pion; 1973.
- Quale J, Landman D, Ravishankar J, Flores C, Bratu S. *Streptococcus pneumoniae*, Brooklyn, New York: fluoroquinolone resistance at our doorstep. *Emerg Infect Dis* 2002;8:594–7.
- Lave JR, Fine MJ, Sankey SS, Hanusa BH, Weissfeld LA, Kapoor WN. Hospitalized pneumonia. Outcomes, treatment patterns, and costs in urban and rural areas. *J Gen Intern Med* 1996;11:415–21.
- Doern GV, Brueggemann AB, Pfaller MA, Jones RN. Assessment of laboratory performance with *Streptococcus pneumoniae* antimicrobial susceptibility testing in the United States: a report from the College of American Pathologists Microbiology Proficiency Survey Program. *Arch Pathol Lab Med* 1999;123:285–9.
- Chin AE, Hedberg K, Cieslak PR, Cassidy M, Stefonek KR, Fleming DW. Tracking drug-resistant *Streptococcus pneumoniae* in Oregon: an alternative surveillance method. *Emerg Infect Dis* 1999;5:688–93.
- Metlay JP, Shea JA, Crossette LB, Asch DA. Tensions in antibiotic prescribing: pitting social concerns against the interests of individual patients. *J Gen Intern Med* 2002;17:87–94.

Address for correspondence: Joshua P. Metlay, 712 Blockley Hall, 423 Guardian Drive, University of Pennsylvania, Philadelphia PA 19104, USA; fax: 215-573-0198; email: jmetlay@cceb.med.upenn.edu

The opinions expressed by authors contributing to this journal do not necessarily reflect the opinions of the Centers for Disease Control and Prevention or the institutions with which the authors are affiliated.

Search past issues of EID at www.cdc.gov/eid