

vention and early intervention that could save the pool money in future expenses. An expanded income-based subsidy, analogous to the earned income tax credit, combined with a requirement that each person have evidence of ability to pay for minor and preventive care, would increase efficiency and equity. Some funds for such coverage are already included in Medicaid program expenditures.

This fundamental restructuring of the payment system would achieve both universal coverage and improved efficiency. Focusing attention on patient outcomes would free clinicians and hospitals to creatively explore ways to deliver care and eliminate payers' focus on fee constraints and micromanagement of clinical decisions. The government's role in the operation of the system would shift to ensuring information availability and transparency in payment. Government would maintain current employment-based subsidies (if a payroll tax is not substituted) and implement income-based redistribution for individuals with low incomes.

A collective risk pool would reallocate funds so all can access appropriate care regardless of their individual health status. The simplified system would eliminate unnecessary administration. Market-disciplined carriers would facilitate payment, provide information, and respond to patient preferences. Appropriate incentives would help ensure that health care expenditures are driven by informed patient and clinician decisions about the care needed to achieve high-quality outcomes.

Financial Disclosures: None reported.

Funding/Support: This work is supported by a grant from the Robert Wood Johnson Foundation Investigators in Health Policy Program. Ingenix Inc provided the data set and software used for the estimation of episodes and costs for various types of episodes but provided no funding for the project.

Role of the Sponsors: Neither the Robert Wood Johnson Foundation nor Ingenix Inc had any role in the preparation, review, or approval of the manuscript.

Acknowledgment: I thank Laura Eaton, MD, MPH, Department of Family and Community Medicine and Institute for Health Policy Studies, University of California, San Francisco (UCSF), for data analysis support and Amy Markowitz, JD, and Beth Newell, BA, Institute for Health Policy Studies, UCSF, for editing. Mss Markowitz and Newell received compensation for their contributions from project funds of the Robert Wood Johnson Foundation Investigators in Health Policy Program; Dr Eaton received no compensation.

REFERENCES

1. DeNavas-Walt C, Proctor BD, Lee CH. *Income, Poverty, and Health Insurance Coverage in the United States: 2005*. Washington, DC: US Census Bureau; 2006. Current Population Reports P60-231.
2. World Health Organization. *World Health Report 2000*. Geneva, Switzerland: World Health Organization; 2000.
3. Balla SJ. Administrative procedures and political control of the bureaucracy. *Am Polit Sci Rev*. 1998;92:663-673.
4. Cotter D, Thamer M, Narasimhan K, Zhang Y, Bullock K. Translating epoetin research into practice: the role of government and the use of scientific evidence. *Health Aff (Millwood)*. 2006;25:1249-1259.
5. Foote SB. Why Medicare cannot promulgate a national coverage rule: a case of regula mortis. *J Health Polit Policy Law*. 2002;27:707-730.
6. Medicare Payment Advisory Commission (MedPAC). *A Data Book: Health-care Spending and the Medicare Program*. Washington, DC: MedPAC; June 2006.
7. Selden TM, Gray BM. Tax subsidies for employment-related health insurance: estimates for 2006. *Health Aff (Millwood)*. 2006;25:1568-1579.
8. Chernew M, Cutler DM, Keenan PS. Increasing health insurance costs and the decline in insurance coverage. *Health Serv Res*. 2005;40:1021-1039.
9. Farber HS, Levy H. Recent trends in employer-sponsored health insurance coverage: are bad jobs getting worse? *J Health Econ*. 2000;19:93-119.
10. Gabel JR, Pickreign JD, Whitmore HH, Schoen C. Embraceable you: how employers influence health plan enrollment. *Health Aff (Millwood)*. 2001;20:196-208.
11. Polsky D, Stein R, Nicholson S, Bundorf MK. Employer health insurance offerings and employee enrollment decisions. *Health Serv Res*. 2005;40:1259-1278.
12. Shen Y-C, Long SK. What's driving the downward trend in employer-sponsored health insurance? *Health Serv Res*. 2006;41:2074-2096.
13. Cooper PF, Schone BS. More offers, fewer takers for employment-based health insurance: 1987 and 1996. *Health Aff (Millwood)*. 1997;16:142-149.
14. Conwell L, Cohen J. *Characteristics of Persons With High Medical Expenditures in the U.S. Civilian Noninstitutionalized Population*, 2002. Rockville, Md: Agency for Healthcare Research and Quality; March 2005. Statistical brief No. 73.
15. Machlin SR, Zodet MW. *Out-of-Pocket Health Care Expenses by Age and Insurance Coverage, 2003*. Rockville, Md: Agency for Health Care Research and Quality; May 2006. Statistical brief No. 126.
16. Wennberg JE, Fisher ES, Stukel TA, Skinner JS, Sharp SM, Bronner KK. Use of hospitals, physician visits, and hospice care during last six months of life among cohorts loyal to highly respected hospitals in the United States. *BMJ*. 2004;328:607-610.
17. Wennberg JE, Fisher ES, Stukel TA, Sharp SM. Use of Medicare claims data to monitor provider-specific performance among patients with severe chronic illness [Web exclusive]. *Health Aff (Millwood)*. October 7, 2004. <http://content.healthaffairs.org/cgi/reprint/hlthaff.var.5v1>. Accessed January 26, 2007.
18. Wennberg JE, Fisher ES, Skinner JS. Geography and the debate over Medicare reform [Web exclusive]. *Health Aff (Millwood)*. February 13, 2002. <http://content.healthaffairs.org/cgi/content/full/hlthaff.w2.96v1/DC1>. Accessed January 26, 2007.
19. Goodman DC, Stukel TA, Chang CH, Wennberg JE. End-of-life care at academic medical centers: implications for future workforce requirements. *Health Aff (Millwood)*. 2006;25:521-531.
20. Fuchs VR. Economics, values, and health care reform. *Am Econ Rev*. 1996;86:124.

Health Disparities and Access to Health

Nicole Lurie, MD, MSPH

Tamara Dubowitz, MSc, SM, ScD

RACIAL AND ETHNIC MINORITIES—POPULATIONS WHO are more likely to be poor, have lower educational levels, or both—are fundamentally at greater risk of ill health than their nonminority, nonpoor, better educated peers.¹ Multiple factors, both within and outside the health care delivery system, probably explain these disparities. Health care and social factors associated with such disparities relate directly to access to care, and

access to care is important because it is believed to lead to better health.

The ideals related to universal access to care might precisely be termed universal access to health, which, by definition, includes the elimination of health disparities. However, erasing disparities in health cannot be accomplished simply by achieving universal access to care; policies that affect public health and the nonmedical determinants of health are also necessary.

Author Affiliations: RAND Center for Population Health and Health Disparities, Arlington, Va.

Corresponding Author: Nicole Lurie, MD, MSPH, RAND Corp, 1200 S Hayes St, Arlington, VA 22302 (lurie@rand.org).

Access to and Quality of Care

Differential access to care is one key contributor to disparities in health. Rates of uninsurance are substantially higher among Hispanics (34%) and blacks (21%) than among whites (13%).² To eliminate disparities in care, having health insurance is necessary but not sufficient. The availability of health insurance does not guarantee access to care—and certainly does not guarantee access to high quality of care. Eisenberg and Power³ likened this phenomenon to an electrical system in which a current passes through a series of resistors, encountering voltage drops as it travels along the circuit. In the health care system circuit, individuals must enroll in available insurance plans that cover needed services, must be able to choose a primary care clinician whom they see regularly and consistently, and must be able to receive appropriate specialty services and high quality of care. Even then, communication challenges such as language differences between patient and clinician, or low health literacy, can impair the effectiveness of that care.^{4,5}

Racial and ethnic minorities and individuals of lower socioeconomic status are more likely to experience these voltage drops, in that even once insured, they are less likely to enter the health care system, establish a regular source of care, or receive care of similar quality to their more advantaged and nonminority peers.³ A report from the Institute of Medicine on disparities in health care concluded that racial disparities in the amount and quality of care exist even for similarly insured patients.⁶ One implication of this conclusion is that actions to eliminate health disparities must go well beyond equalizing insurance coverage.

Thus far, much of health disparities research and efforts to address health disparities have focused largely on factors that are actionable within the context of the health care system. Disparities in care are increasingly viewed as a problem with quality, implying that methods to improve quality will narrow the disparities in care, and by extension, the disparities in health. Unfortunately, many of the uninsured are left out of efforts to address disparities through improving quality because they do not access the health care system.

Despite the limitations of focusing solely on the health care system, encouraging evidence suggests that some disparities in care are narrowing. For example, Trivedi et al⁷ have shown that differences in receipt of a low-density lipoprotein cholesterol or a hemoglobin A_{1c} test between blacks and whites enrolled in Medicare have narrowed substantially, a sign of what can happen when quality is measured and reported. However, these improvements were not accompanied by similar reductions in either lipid or glucose control for those with heart disease or diabetes, suggesting that narrowing disparities in long-term outcomes—or disparities in health—cannot be achieved by simply ordering appropriate tests or prescribing appropriate medication. Although quality measures may be based on incorrect metrics (eg, process of care vs outcomes achieved), these seem-

ingly paradoxical findings also suggest that the solution to reducing disparities in outcomes entails far more than what health care services can provide and that there is a complex interplay of health care, public health, and social factors at work.

Public Health and Nonmedical Determinants

Health is the result of an individual's genetic makeup, income and educational status, health behaviors, communities in which the individual lives, and environments to which he or she is exposed. Indeed, the contribution of health care to health status is modest, estimated to be approximately 15%.⁸ Although a person's genetic composition is established, other factors—environment, health behaviors, community resources, and even income—can be influenced by a combination of a robust public health system and changes in social and economic policy. Public health efforts have been critical for eliminating disparities in exposure to environmental toxins (eg, lead and asbestos), promoting healthful behaviors (eg, smoking cessation and physical activity), and improving community resources (eg, parks, lighting, and sidewalks in disadvantaged neighborhoods). These actions can help prevent disparities in the incidence and prevalence of chronic disease. Surveillance and disease-control efforts contribute not only to the health of the population overall but to the health of low-income and minority populations who are more likely to experience higher incidence, morbidity, and mortality from infectious diseases (eg, human immunodeficiency virus or tuberculosis).⁹

In addition to the role of individual socioeconomic and psychosocial determinants of health, characteristics of the neighborhood in which one lives also have an independent effect on health.¹⁰ Researchers are even beginning to identify biological pathways through which individual and neighborhood socioeconomic status, for example, “get under the skin.”¹¹ Some of these pathways involve excess cortisol, inflammation, oxidative stress, and gene methylation and are associated with increased risk of chronic diseases such as coronary heart disease, diabetes, and certain cancers.¹²

How might the transformation from environment to poor health occur? One proposed mechanism is residential racial segregation which, over a century in the making, leads to racial differences in socioeconomic status.¹³ Individuals who are members of racial/ethnic minority groups are likely to have lower individual socioeconomic status and are more likely to live in racially and economically segregated and stressful environments that lack resources, such as employment opportunities; high-quality, affordable food; and safe places in which to play and be physically active. These neighborhoods are also more likely to contain environmental toxins and have higher rates of crime. Neighborhood disadvantage also links back to the health care delivery system; access to care in such neighborhoods is also poor. In addition to having higher rates of uninsurance and sicker populations, health care services in such neighborhoods may also

operate at a disadvantage. For example, Bach et al¹⁴ reported that approximately 20% of physicians care for 80% of the black population in the United States. Those physicians disproportionately report that they are less able to access resources for their patients, including specialty consultation and diagnostic tests.

Several health plans that are part of the National Health Plan Disparities Collaborative, a group of health insurance policy makers addressing racial/ethnic disparities in care, have found that in sociodemographically similar neighborhoods, members living in one neighborhood may receive elements of high-quality care while those residing in another do not.¹⁵ This suggests that the causes of disparities in care are not as simple as either insurance or socioeconomic status but that other factors are likely to be operating. Thus, insurance coverage, quality of care, public health measures, and community resources all appear to be important in addressing disparities in care and in health.

Policies to Reduce Disparities in Health

Is the goal of narrowing disparities in health achievable? There is cause for optimism in the finding that some countries achieve both universal access to care and have better health outcomes, including fewer health disparities. Some national governments have enumerated the kinds of policies likely to improve health.^{16,17} In Britain, the Acheson Commission provided a set of 39 evidence-based recommendations for social policies that could contribute to such a goal. Notably, only 3 of these recommendations pertained directly to health care. In the United States, the Centers for Disease Control and Prevention publishes the Community Guide, an evidence-based analysis of interventions that would improve health,¹⁸ including interventions targeted at the health care system (eg, reducing financial barriers to vaccination and disease management), the public health system (eg, community-wide campaigns to promote physical activity and immunization programs), and the social environment (eg, comprehensive early childhood development programs and tenant-based rental voucher or housing mobility programs).

Some social conditions that contribute to poor health may be amenable to federal policy changes, such as improving income by increasing the minimum wage or expanding the earned income tax credit. Improving environmental air quality could improve health and reduce Medicare expenditures.¹⁹ The law offers a panoply of tools for improving health; for example, legal approaches could be used to facilitate more healthful lifestyles and help address obesity.²⁰

Fortunately, neither improving access nor solely focusing on conditions that promote health is exclusively dependent on federal policy. State and local policies may also play a role. From an access perspective, in California, parents of children newly enrolled in the State Children's Health Insurance Program reported that their children performed better in school, felt better physically, and were able to get along

better with their peers than they did before they had insurance.²¹ Furthermore, ethnic disparities in children's access to health care were largely reduced. Several other states including Massachusetts, Vermont, California, and Maine have recently initiated plans for universal coverage.²²

States are enacting health-promoting legislation. Following the lead of California, some are moving to set limits on automobile and factory emissions. Local incentives have helped locate supermarkets in low-income areas. Citizens of Los Angeles passed a bond initiative to renovate city parks. Local zoning ordinances that limit urban sprawl and promote walkable communities are becoming more widespread. Governments at all levels could develop creative approaches if their administrative entities, whether cabinet-level departments or city governmental units, convened regularly to identify opportunities to enact health-promoting policies.

A substantial proportion of US physicians view issues related to access to care, public health influences on health, and nonmedical determinants of health as important areas for their public responsibilities.²³ However, far fewer physicians reported being engaged in community participation, political action, or collective advocacy regarding these topics in the past 3 years. Increased engagement of health professionals of all types may also be necessary—although not sufficient—to move the nation toward universal access to health care.

Medicine and public health have been likened to trains running on parallel tracks, never to meet.²⁴ Social and economic policies that affect health are often viewed as the third rail. However, bringing together medical care, public health, and other policy reforms that address the nonmedical determinants of health will be essential for making progress on health disparities. Such approaches are likely to improve access to care, access to health, and ultimately reduce health disparities.

Financial Disclosure: None reported.

Funding/Support: Supported in part by National Institute of Environmental Health Sciences grant 5 P50 ES012383-04.

Role of the Sponsor: The funding agency had no role in the preparation, review, or approval of the manuscript.

REFERENCES

1. Lynch JW, Kaplan GA. Socioeconomic position. In: Kawachi I, Berkman L, eds. *Social Epidemiology*. Oxford, England: Oxford University Press; 2001:13-35.
2. Finegold K, Wherry L. Race, Ethnicity, and Health. Snap Shots of American Families 3. http://www.urban.org/uploadedpdf/310969_snapshots3_no20.pdf. Accessed January 15, 2007.
3. Eisenberg JM, Power EJ. Transforming insurance coverage into quality health care: voltage drops from potential to delivered quality. *JAMA*. 2000;284:2100-2107.
4. Dewalt DA, Berkman ND, Sheridan SL, Lohr KN, Pignone M. Literacy and health outcomes: a systematic review of the literature. *J Gen Intern Med*. 2004;19:1228-1239.
5. Aboul-Enein FH, Ahmed F. How language barriers impact patient care: a commentary. *J Cult Divers*. 2006;13:168-169.
6. Institute of Medicine of the National Academies. *Unequal Treatment: Confronting Racial and Ethnic Disparities in Healthcare*. Washington, DC: National Academies Press; 2003.
7. Trivedi AN, Zaslavsky AM, Schneider EC, Ayanian JZ. Trends in the quality of care and racial disparities in Medicare managed care. *N Engl J Med*. 2005;353:692-700.
8. Mokdad AH, Marks JS, Stroup DF, Gerberding JL. Actual causes of death in the United States, 2000. *JAMA*. 2004;291:1238-1245.

9. Evans GW. Psychological costs of chronic exposure to ambient air pollution. In: Isaacson RL, Jensen KF, eds. *The Vulnerable Brain and Environmental Risks*. Vol 3. New York, NY: Plenum; 1994:167-182.
10. Kawachi I, Berkman LF, eds. *Neighborhoods and Health*. New York, NY: Oxford University Press; 2003.
11. Taylor SE, Repetti RL, Seeman T. Health psychology: what is an unhealthy environment and how does it get under the skin? *Annu Rev Psychol*. 1997;48:411-447.
12. Seeman TE, Singer BH, Rowe JW, Horwitz RI, McEwan BS. Price of adaptation—allostatic load and its health consequences: MacArthur studies of successful aging. *Arch Intern Med*. 1997;157:2259-2268.
13. Williams DR, Collins C. Racial residential segregation: a fundamental cause of racial disparities in health. *Public Health Rep*. 2001;116:404-416.
14. Bach PB, Pham HH, Schrag D, Tate RC, Hargraves JL. Primary care physicians who treat blacks and whites. *N Engl J Med*. 2004;351:575-584.
15. National Health Plan Collaborative Phase One Summary Report: Reducing Racial and Ethnic Disparities & Improving Quality of Health Care AHIP Medical Leadership Forum; November 2006. http://www.chcs.org/NationalHealthPlanCollaborative/images/641_104_NHCP_summary_V3.pdf. Accessed January 15, 2007.
16. Acheson D, Barker D, Chambers J, Graham H, Marmot M, Whitehead M *The Report of the Independent Inquiry Into Health Inequalities*. London, England: the Stationary Office; 1998. <http://www.archive.official-documents.co.uk/document/doh/ih/contents.htm>. Accessed January 15, 2007.
17. Health Canada. Achieving health for all: a framework for health promotion. 1986. http://www.hc-sc.gc.ca/english/care/achieving_health.html. Accessed January 15, 2007.
18. Guide to community preventive services: motor vehicle occupant injury. Centers for Disease Control and Prevention Web site. <http://www.thecommunityguide.org/mvoi>. Last updated June 23, 2006. Accessed January 29, 2007.
19. Fuchs VR, Frank SR. Air pollution and medical care use by older Americans: a cross-area analysis. *Health Aff (Millwood)*. 2002;21:207-214.
20. Gostin LO. Law as a tool to facilitate healthier lifestyles and prevent obesity. *JAMA*. 2007;297:87-90.
21. Seid M, Varni JW, Cummings L, Schonlau M. The impact of realized access to care on health-related quality of life: a two-year prospective cohort study of children in the California State Children's Health Insurance Program. *J Pediatr*. 2006;149:354-361.
22. The push for universal healthcare. *State Net Capital J*. December 18, 2006; XIV.
23. Gruen RL, Campbell EG, Blumenthal D. Public roles of US physicians: community participation, political involvement, and collective advocacy. *JAMA*. 2006;296:2467-2475.
24. Remarks by Donna E. Shalala: Secretary of Health and Human Services. Presentation at: the National Congress of the Medicine/Public Health Initiative; March 2, 1996; Chicago, Ill. <http://www.mphi.net/content.php?section=interviews&article=74>. Accessed January 15, 2007.

Structural Impairments That Limit Access to Health Care for Patients With Disabilities

Kristi L. Kirschner, MD

Mary Lou Breslin, MA

Lisa I. Iezzoni, MD, MSc

THE FOLLOWING 3 CASES REPRESENT SUBSTANDARD CARE for patients with disabilities, yet they occurred recently at US tertiary care medical centers with the latest technologies and well-qualified physicians. These failures resulted from basic, "low-tech" structural deficiencies—lack of accessible call systems, diagnostic equipment, and examination tables.

Joe is paralyzed, dependent on a ventilator, and unable to speak. His hospital room was at the end of the corridor and had no accessible call system to summon assistance. When his ventilator became disconnected and then was not promptly recognized, Joe became extremely anxious about being in a hospital.

Susan, who uses a wheelchair, had trouble breathing. She needed an echocardiogram, which was performed while she sat in her wheelchair. The echocardiogram was of poor technical quality and yielded little information.

Chuck has paraplegia and new rectal bleeding. The gastroenterologist refused to perform a diagnostic flexible sigmoidoscopy because the office did not have wheelchair-accessible examination tables or lifting provisions. He sent Chuck home with 3 hemocult cards.

Despite passage of the Americans with Disabilities Act (ADA) in 1990, inaccessible facilities, equipment, and communication systems still compromise health care experi-

ences for individuals with disabilities in the United States.¹⁻³ Although no direct evidence currently exists about the population prevalence of these problems nationwide, increasing numbers of legal cases, small studies, and circumstantial evidence point to widespread access barriers for patients with disabilities within US health care settings.

This commentary reviews the legal and policy contexts for ensuring physical accessibility to health care facilities and also considers potential effects of environmental barriers on patient safety, quality of care, and health care worker safety. These contexts involve 3 fundamental concepts. First, environmental barriers contribute significantly to disability.³ Second, the architectural notion that form follows function holds important implications for health care, where creation of therapeutic environments is a core value. The barriers that disabled patients confront represent quality problems and also heighten patients' sense of stigmatization, disenfranchisement, and demoralization.⁴ And third, the concept of universal design, human-centered design that keeps all potential users in mind,³ recognizes the diversity of patient populations, health care professionals, other workers, and all individuals using these environments. Just as health care has worked toward achieving racial, ethnic, and cultural diversity, the time has come to embrace bodily diversity. Instead of sorting into binary categories of able vs

Author Affiliations: Rehabilitation Institute of Chicago, Northwestern University Feinberg School of Medicine, Chicago, Ill (Dr Kirschner); Disability Rights Education and Defense Fund, Berkeley, Calif (Ms Breslin); and Harvard Medical School, Institute for Health Policy, Massachusetts General Hospital, Boston (Dr Iezzoni).
Corresponding Author: Kristi L. Kirschner, MD, Rehabilitation Institute of Chicago, 345 E Superior St, Room 1136, Chicago, IL 60611 (kkirschner@ric.org).