Health Related Quality of Life Among Adults Reporting Arthritis: Analysis of Data from the Behavioral Risk Factor Surveillance System, US, 1996–99

FATIMA MILI, CHARLES G. HELMICK, MATTHEW M. ZACK, and DAVID G. MORIARTY

ABSTRACT. Objective. To characterize health related quality of life (HRQOL) among people with and without selfreported arthritis in the general population by selected demographic and behavior characteristics.

> Methods. We analyzed data from a cross sectional random-digit telephone survey [the Behavioral Risk Factor Surveillance System (BRFSS)] of civilian noninstitutionalized adults aged 18 years or older from 15 states and Puerto Rico, all of which used an optional arthritis survey module for one or more years from 1996 through 1999. We compared HRQOL among people with arthritis, defined as chronic joint symptoms (CJS) or doctor-diagnosed arthritis, those within one of 3 arthritis subgroups (i.e., only doctor-diagnosed arthritis, only CJS, and both doctor-diagnosed arthritis and CJS), and those without

> Results. On an age-adjusted basis, respondents with arthritis had significantly worse HRQOL than respondents without arthritis. Members of all 3 arthritis subgroups had significantly worse HRQOL than those without arthritis. Those with both CJS and doctor-diagnosed arthritis had consistently worse HRQOL than those with only CJS, who in turn had worse HRQOL than those with only doctor-diagnosed arthritis. In some of the demographic and behavioral subgroups, HRQOL differences between those with and without arthritis greatly exceeded the differences for the overall study.

> Conclusion. Because many adults report arthritis and because arthritis substantially worsens their HRQOL, HRQOL measures like those in the BRFSS may be useful in monitoring the burden of arthritis and in tracking the success of population interventions for arthritis. (J Rheumatol 2003;30:160-6)

Key Indexing Terms: **ARTHRITIS** QUALITY OF LIFE

CROSS SECTIONAL SURVEY

EPIDEMIOLOGY SOCIOECONOMIC FACTORS

Arthritis and other rheumatic conditions are a large and growing public health problem that affected 43 million people in the United States in 1997¹, and with the aging of the baby boom generation will affect an estimated 60 million Americans by 2020². These conditions constitute the most frequent cause of disability in the US³, cost \$65 billion in direct and indirect costs in 19924, and were associated with 744,000 hospitalizations and 44 million ambulatory-care visits in 1997⁵. Because arthritis and other rheumatic conditions seldom cause death but have a substantial impact on health, health related quality of life (HRQOL) measures indicate their

measures may also help track the success of clinical and public health interventions for arthritis.

influence better than mortality rates. In addition, HRQOL

Although HRQOL among people with arthritis has often been measured in clinical, research, and other special populations⁶⁻¹⁰, it has been measured less frequently in general populations larger than a single community^{11,12}. Standard generic health status measures such as the Medical Outcomes Study Short-Form 36 (SF-36) and the Sickness Impact Profile (SIP) have been shown to be reliable, valid, and responsive to change in various clinical populations including people with arthritis¹³⁻¹⁷. These clinically valid measures have also sometimes been used for population assessment of HRQOL but, because of their overall length and associated respondent burden, have mostly been used for small-scale assessments¹⁸ or in specialized populations¹⁹.

Measuring HRQOL in the larger, general population is important in understanding and monitoring the effects of arthritis on HRQOL because 16% of people with arthritis do not see a doctor for their arthritis²⁰; therefore clinic based studies cannot capture the entire spectrum of disease. Beginning in the early 1990s, the Centers for Disease Control and Prevention (CDC), with advice and help from its academ-

From the Health Care and Aging Studies Branch, Division of Adult and Community Health, National Center for Chronic Disease Prevention and Health Promotion, Centers for Disease Control and Prevention, Atlanta,

F. Mili, MD, PhD, Medical Epidemiologist; C.G. Helmick, MD, Medical Epidemiologist; M.M. Zack, MD, MPH, Medical Epidemiologist; D.G. Moriarty, Program Analyst.

Address reprint requests to Dr. C.G. Helmick, Health Care and Aging Studies Branch, Division of Adult and Community Health, National Center for Chronic Disease Prevention and Health Promotion, Centers for Disease Control and Prevention, 4770 Buford Highway, Mailstop K45, Atlanta, GA 30341.

Submitted January 16, 2002; revision accepted July 4, 2002.

ic and public health practitioner colleagues, began to develop and validate a feasible set of HRQOL measures that could be used in population surveys. The product of this collaboration was a core set of 4 CDC HRQOL survey items that ask participants to rate their general health, physical health, mental health, and activity limitations. This set of survey items has been shown to be reliable, valid, and responsive to naturally occurring changes in populations²¹⁻²³. It has been used to characterize the burden of disease and disability²⁴⁻²⁶; to identify unmet health and social service needs of older adults^{27,28}; to determine the size of known or suspected health disparities by demographic, socioeconomic, and employment status; and to reflect seasonal patterns and time trends^{23,29}.

An earlier study used data from the Behavioral Risk Factor Surveillance System (BRFSS) to characterize details of the prevalence of arthritis in 15 states and Puerto Rico³⁰. We used the same data to characterize HRQOL among people with and without self-reported arthritis in the general population by selected demographic and behavior characteristics.

MATERIALS AND METHODS

Telephone survey. The BRFSS is an ongoing state-based, random-digit dialed telephone survey, conducted by the CDC and state health departments. The BRFSS has been described in detail³¹. Briefly, it collects self-reported health status, demographic, behavioral risk factor, and other information from a representative sample of the civilian, noninstitutionalized population aged ≥ 18 years in each US state and some territories³². The BRFSS is exempt from human subjects review by the CDC Institutional Review Board because it is surveillance and not research. The study population we used for this analysis consisted of 54,154 respondents from 15 states (Alabama, Arizona, Georgia, Hawaii, Kansas, Louisiana, Mississippi, Missouri, Montana, Nebraska, New Jersey, Ohio, Oklahoma, Rhode Island, West Virginia) and Puerto Rico, all of which used a standard 6 item optional BRFSS Arthritis Module in one or more years from 1996 through 1999. HRQOL was measured by participants' responses to the following 4 item set of questions developed and validated by the CDC for use in population surveillance: (1) Self-rated health: Would you say that in general your health is excellent, very good, good, fair, or poor? (2) Recent physical health: Now thinking about your physical health, which includes physical illness and injury, for how many days during the past 30 days was your physical health not good? (3) Recent mental health: Now thinking about your mental health, which includes stress, depression, and problems with emotions, for how many days during the past 30 days was your mental health not good? (4) Recent activity limitation: During the past 30 days, for about how many days did poor physical or mental health keep you from doing your usual activities, such as self-care, work, or recreation?

Analysis. We calculated overall unhealthy days as the sum of physically and mentally unhealthy days, with this sum not to exceed 30 days for an individual. As outcome measures, we calculated the percentage of participants who rated their health as fair or poor, as well as participants' mean number of physically unhealthy days, mentally unhealthy days, recent activity limitation days, and overall unhealthy days.

We classified people as having arthritis if they reported having either chronic joint symptoms (CJS) or doctor-diagnosed arthritis. They were considered to have CJS if they responded yes to 2 questions: (1) During the past 12 months, have you had pain, aching, stiffness, or swelling in or around a joint? (2) Were these symptoms present on most days for at least one month? They were considered to have doctor-diagnosed arthritis if they responded yes to the question, Have you ever been told by a doctor that you have arthritis? All other respondents, including those who responded don't know, refused to answer the questions, or had missing data, were considered not to have arthritis.

Covariates included in the analyses were age, sex, race, Hispanic origin,

marital status, education, employment status, physical activity, body mass index (BMI), health care coverage or insurance, cigarette smoking, and alcohol beverage drinking. We grouped respondents into 3 age categories (18–44, 45–64, and 65+ years) and 4 physical activity categories (inactive; active, irregular and not sustained; active, regular but not intensive; and active, regular and intensive). We calculated respondents' BMI [weight (kg) divided by height (m²)] from their self-reported height and weight and grouped these values into the 4 National Institutes of Health weight classes³3: underweight (BMI < 18.5), normal weight (18.5 \leq BMI < 25.0), overweight (25.0 \leq BMI < 30.0), and obese (BMI \geq 30.0). We grouped respondents by their cigarette smoking into those who smoke daily, those who smoke some days, former smokers, and never smokers. We grouped respondents by their alcohol consumption into nondrinkers, occasional drinkers (\leq 29 drinks/month), and regular drinkers (\geq 29 drinks/month).

To account for the complex sample survey design of BRFSS, we used sample weights and SUDAAN statistical software³⁴ in estimating the 95% confidence intervals (CI) for the 5 HRQOL measures among respondents without arthritis, those with arthritis, and those in each of the 3 arthritis subgroups (i.e., those with only doctor-diagnosed arthritis, those with only CJS, and those with both doctor-diagnosed arthritis and CJS). To assess HRQOL among participants in each of these groups, we treated the HRQOL measures as continuous variables and used linear regression procedures. We also computed differences in HRQOL between those with arthritis and those without arthritis as well as differences in HRQOL among the 3 arthritis subgroups. Because both the prevalence of arthritis and the HRQOL measures among those with arthritis vary with age³⁵, we stratified by age group or adjusted all the analyses for age (as a continuous variable) using the age distribution of the sample population. Mean group HRQOL measures in which 95% CI did not overlap were considered to be significantly different.

We used a mosaic plot³⁶ to depict the sample population stratified by age group, sex, and arthritis status; to illustrate the relative sizes and proportions of affected populations and the mean overall unhealthy days for these groups; and to identify those populations most severely affected by arthritis.

RESULTS

Of the 54,154 respondents to the BRFSS Arthritis Module, 17,556 (32%) reported having arthritis. Of those with arthritis, 36% reported having only doctor-diagnosed arthritis, 25% only CJS, and 38% both doctor-diagnosed arthritis and CJS.

Respondents with arthritis had significantly worse HRQOL than respondents without arthritis (Figure 1). Respondents with arthritis reported having fair or poor health 3 times as often as those without arthritis (23.8 vs 7.3%), and they averaged 4.0 more physically unhealthy days, 2.3 more mentally unhealthy days, 4.9 more overall unhealthy days, and 2.3 more recent activity limitation days. All 3 arthritis subgroups had significantly worse mean HRQOL measures than the nonarthritis group (Figure 1). Those with both CJS and doctor-diagnosed arthritis had consistently worse HRQOL than those with only CJS, who in turn had worse HRQOL than those with only doctor-diagnosed arthritis.

Trends in HRQOL by demographic and behavioral variables were generally similar for those with and without arthritis (Table 1). In some of these demographic and behavioral-risk subgroups, however, HRQOL differences between those with arthritis and those without arthritis exceeded (by at least 8.0% or 0.5 days) differences for the overall study population (Table 1). Groups whose HRQOL was thus especially affected by arthritis included: those 45–64 years old, blacks and other races, Hispanics, those separated from their spouses,

Personal, non-commercial use only. The Journal of Rheumatology Copyright © 2003. All rights reserved.

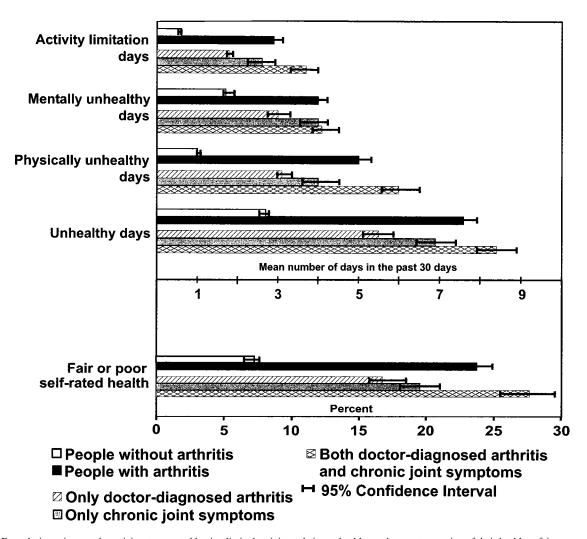


Figure 1. Days during prior month participants reported having limited activity or being unhealthy, and percentage rating of their health as fair or poor, by arthritis status and arthritis subgroup.

those with less than a high school education, those unemployed more than one year, homemakers, those unable to work, the physically inactive, those who were underweight, those who lacked health care coverage or insurance, those who currently smoke, and those who did not drink alcohol. In other complementary subgroups, the HRQOL differences between those with arthritis and those without arthritis were correspondingly much less (at least 0.5 fewer days) than expected. These groups included: whites and Asian/Pacific Islanders, those other than Hispanics, those currently married, widowed or never married, college graduates, the employed, students, the retired, those reporting regular physical activity, and those with health care coverage or insurance.

Although the proportions of respondents with only doctordiagnosed arthritis or with both CJS and doctor-diagnosed arthritis increased substantially with age, the proportions with only CJS remained constant from 18 through 64 years old before declining at older ages (Figure 2). At all ages, women reported more doctor-diagnosed arthritis than men. Among those without arthritis and among all those with arthritis (except those 45 years old or older with CJS only), women consistently reported more overall unhealthy days than men. Mean unhealthy days ranged from 2.6 days for men aged 45–64 years without arthritis to 12.7 days for women aged 18–44 years with doctor-diagnosed arthritis and CJS.

DISCUSSION

In our age adjusted analysis, respondents with arthritis had significantly worse HRQOL than respondents without arthritis. All 3 arthritis subgroups had significantly worse HRQOL than those without arthritis. Those with both CJS and doctor-diagnosed arthritis had consistently worse HRQOL than those with only CJS, who in turn had worse HRQOL than those with only doctor-diagnosed arthritis. In some of the demographic and behavioral risk subgroups, HRQOL differences between those with arthritis and those without arthritis greatly exceeded the differences for the overall study.

This study has several strengths. First, because we ana-

Table 1. Percentage of those reporting arthritis* and health related quality of life measures among people reporting arthritis (n = 17,556) and those not reporting arthritis (n = 36,598), age adjusted**, by selected characteristics — Behavioral Risk Factor Surveillance System, US, 1996–99.

		HRQOL Self-Rated Mean Number of Days During the Past 30 Days					
	337141- A41 ***		oor Health, %	Mean Number of Days During the Past 30 Days Unhealthy Days Recent Activity Limitation Day			
Characteristic	With Arthritis, %	Arthritis	No Arthritis	Arthritis	No Arthritis	Arthritis	No Arthritis
Age group, yrs							
18–44	16	19.7	2.4	8.9	4.1	3.2	0.8
45–64	39	31.8	7.0	9.0	3.4	3.8	0.9
≥ 65	55	40.3	18.2	7.5	3.7	3.5	1.3
Sex							
Women	26	31.3	9.3	9.2	4.4	3.5	1.1
Men	33	29.7	9.1	7.7	3.1	3.5	0.9
Race/ethnicity							
White	31	22.2	4.1	7.3	3.7	3.1	0.8
Black	28	36.8	12.3	9.5	4.2	4.6	1.1
Asian/Pacific Islander	14	21.9	5.3	6.3	2.6	2.6	0.6
American Indian/Alaska Nat	ive 29	34.3	13.0	10.2	3.9	3.4^{\dagger}	1.6^{\dagger}
Other	18	37.3	11.1	9.0	4.4	3.8	1.0
Hispanic ethnicity							
Hispanic origin	22	39.4	13.4	8.9	3.8	3.8	1.2
Non-Hispanic origin	31	21.6	5.0	8.0	3.7	3.2	0.8
Marital status							
Married	31	22.5	4.1	6.3	2.4	2.2	0.5
Divorced	35	29.1	8.7	8.8	4.1	3.9	1.2
Widowed	56	29.0	10.4	7.5	4.1	2.9	1.2
Separated	32	41.9	11.6	11.7	4.6	4.7	1.2
Never married	14	27.0	9.6	7.1	3.1	2.7	0.9
Unmarried couple	21	33.6	10.6	9.4	4.1	4.6	1.0
Education level							
Less than high school	43	50.9	19.4	11.5	4.7	5.4	1.5
High school graduate or							
some college	30	26.4	6.5	8.0	3.7	3.2	0.9
College graduate	22	14.3	1.5	5.9	2.9	1.9	0.6
Employment status							
Employed for wages	22	14.2	0.7	3.7	1.5	0.4	0.0
Self-employed	25	10.2	0.0	4.3	1.4	0.7^{\dagger}	0.0^{\dagger}
Out of work > 1 year	34	41.4	14.2	13.6	4.5	6.2	1.6
Out of work < 1 year	23	29.2	8.9	11.2	4.7	4.3	1.7
Homemaker	33	32.3	5.9	6.5	2.4	2.3	0.7
Student	9	16.7	2.2	3.5^{\dagger}	1.3^{\dagger}	0.0^{\dagger}	0.2^{\dagger}
Retired	53	28.6	10.6	5.9	2.2	2.5	0.6
Unable to work	69	73.7	50.7	19.1	11.9	13.6	8.2
Physical activity							
Inactive	35	42.2	13.3	11.2	4.3	5.6	1.4
Irregular, not sustained	27	28.9	8.3	7.9	3.6	3.2	0.9
Regular, not intensive	25	23.0	6.6	7.0	3.6	2.4	0.9
Regular, intensive	29	19.3	4.7	6.3	3.0	2.0	0.6
Body mass index							
Underweight	22	40.8	10.9	10.6	4.4	5.3	1.6
Normal	24	21.7	5.9	7.0	3.7	2.4	0.7
Overweight	30	23.8	6.7	7.0	3.0	2.6	0.6
Obese	43	35.5	12.7	9.2	4.2	3.6	1.1
Health care coverage or insuranc	e						
Yes	31	23.5	5.9	6.7	3.2	2.9	0.8
No	25	37.6	12.5	10.2	4.3	4.1	1.2
Smoking status							
Current, every day	32	37.7	11.9	10.4	4.3	4.3	1.1
Current, some days	25	31.0	9.9	9.4	4.7	4.7	1.3
Former	40	27.7	7.8	7.1	3.3	2.6	0.9
Never	25	25.9	7.1	7.0	2.8	2.4	0.7
Alcohol drinker							
Non-drinker	34	43.4	13.1	10.8	3.7	5.2	1.2
≤ 29 Drinks/mo	25	25.2	7.3	7.6	3.8	2.8	1.0
> 29 Drinks/mo	25	27.6	9.9	8.0	4.3	3.1	1.1

^{*} Persons with arthritis were defined as those having either chronic joint symptoms or doctor-diagnosed arthritis.

Personal, non-commercial use only. The Journal of Rheumatology Copyright © 2003. All rights reserved.

^{**} Using age as a continuous variable, except for age-group analyses.

[†] In these comparisons, those with arthritis did not have significantly worse HRQOL than those without arthritis. In all other comparisons, those with arthritis did have significantly worse HRQOL.

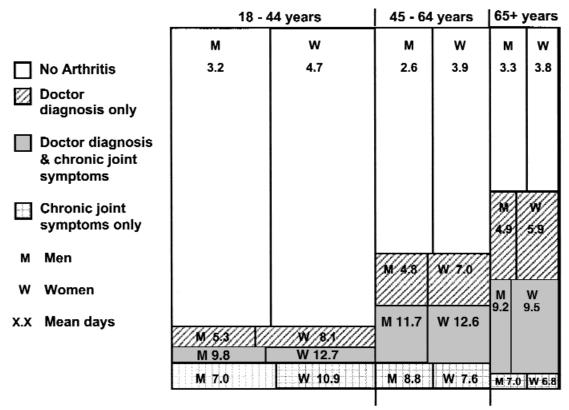


Figure 2. Proportions of adults by age group, sex, and arthritis status and the mean number of unhealthy days for each subgroup, 1996–99 BRFSS (15 states and Puerto Rico). Areas of the mosaic correspond to the proportions these strata represent in the BRFSS states included in this study.

lyzed state based population survey data, our results are generalizable to the populations surveyed. Second, because the CDC HRQOL measures used here have been validated by CDC and others for use in population surveillance^{21-23,37,38}, they can be confidently used to describe the nature and extent of the perceived health and activity limitation burden associated with different subgroups of arthritis in different populations. This standard CDC HRQOL 4 item set correlated well with related SF-36 scales both in general community populations³⁷ and among adults with known disabilities³⁹. It has also been acceptably validated in cognitive studies conducted among elders by the National Center for Health Statistics^{40,41}. In addition, it has been found to predict rates of mortality, hospital utilization, and outpatient visits among low income elderly people⁴²⁻⁴⁴. Third, because the BRFSS combines data from identical state based surveys to create a large sample, the estimates of HRQOL among people with arthritis and among subgroups of people with arthritis derived from it are more precise than those based on smaller samples. Fourth, by examining HRQOL for the 3 subgroups of people with arthritis and for various demographic groups, we were able to identify high risk groups. Fifth, the large sample size allowed us to identify age and sex-specific HRQOL differences for better targeting of interventions.

This study also has several limitations. First, the BRFSS case definition for self-reported arthritis has not yet been val-

idated, although validation studies are under way. The more comprehensive definition of arthritis in this report (people with CJS or doctor-diagnosed arthritis), however, may better identify those with arthritis and other rheumatic conditions than a previous case definition that included only those with CJS^{45,46}. Second, because the BRFSS does not ask about many other common chronic health conditions that affect people's HRQOL, we could not adjust for these conditions. Third, because the BRFSS excludes people without telephones, those in the military, those in institutions (for example, nursing homes), and those younger than 18 years, the data do not represent the entire population in these states and thus underestimate the total number affected by arthritis. Fourth, because the time and functional capacity required to complete the BRFSS may limit participation by people with arthritis who have poor health and functional limitations, our data may underestimate the proportion of people with poor HRQOL. Fifth, unhealthy days may be overestimated for people who report both physical and mental unhealthy days when these days overlap. Sixth, no published studies have examined how these HRQOL measures change over time or how these measures among people with arthritis are affected by clinical and public health interventions. Finally, the states using the BRFSS arthritis module may not be representative of other states.

The substantial detrimental effect that arthritis has on

HRQOL, especially after age 45 years, suggests the importance of identifying potential public health interventions. Older people and those in other groups whose HRQOL was disproportionately affected by arthritis may need interventions to reduce barriers to effective arthritis treatment such as lack of insurance, cost of treatment, distance from treatment, and lack of transport. Because most of the burden from only CJS occurs among those 18-44 years old, CJS may represent subacute effects of injury associated with occupation or life style. Identifying specific causes of only CJS might lead to interventions to prevent future arthritis or reduce its effect or severity. For example, 16% of people with arthritis in the National Health Interview Survey had not seen a doctor for their arthritis²⁰. Compared with those who saw a doctor for their arthritis, these people were less likely to be overweight or not have health insurance, and more likely to be male, younger, and to have better self-perceived health and fewer activity or work limitations due to arthritis²⁰.

Because arthritis substantially affects people's health, using HRQOL measures like those in the BRFSS may be useful in monitoring the effect and the burden of arthritis and in identifying high risk groups for targeted interventions such as the Arthritis Self-Help Course⁴⁷, which helps people with arthritis to decrease their pain and number of physician visits. The Arthritis Foundation estimates, however, that these interventions reach less than 1% of the target population⁴⁸. Other interventions may include water and land exercise, social support, and improved access to health care services.

Further research is needed to determine the specific effect of arthritis and disability on HRQOL after controlling for other factors, such as common chronic health conditions or low socioeconomic status⁴⁹. Longitudinal studies are also needed to examine how these HRQOL measures of people with arthritis change over time and respond to clinical and public health interventions. Such research could improve current clinical and public health interventions developed for people with arthritis, and guide efforts in reaching the Healthy People 2010⁵⁰ goal of increasing the quality and years of healthy life for people with arthritis.

ACKNOWLEDGMENT

We thank the Behavioral Surveillance Branch at CDC and the state BRFSS coordinators who helped conduct the surveys. We thank Barbara Dougherty for doing the graphics.

REFERENCES

- Centers for Disease Control and Prevention. Prevalence of arthritis

 United States, 1997. MMWR Morb Mortal Wkly Rep 2001;50:334-6.
- CDC. Arthritis prevalence and activity limitations United States, 1990. MMWR Morb Mortal Wkly Rep 1994;43:433-8.
- CDC. Prevalence of disabilities and associated health conditions United States, 1999. MMWR Morb Mortal Wkly Rep 2001; 50:120.5
- Yelin E, Callahan LF. The economic cost and social and psychological impact of musculoskeletal conditions. Arthritis Rheum 1995;38:1351-62.

- CDC. Impact of arthritis and other rheumatic conditions on the health-care system – United States, 1997. MMWR Morb Mortal Wkly Rep 1999;48:349-53.
- Reynolds D, Torrance GW, Badley EM, et al. Modeling the population health impact of musculoskeletal diseases: arthritis. J Rheumatol 1993;20:1037-47.
- Carr AJ. Beyond disability: measuring the social and personal consequences of osteoarthritis. Osteoarthritis Cartilage 1999; 7:230-8.
- 8. Jordan JM. Effect of race and ethnicity on outcomes in arthritis and rheumatic conditions. Curr Opin Rheumatol 1999;11:98-103.
- Currey SS, Rao JK, Winfield JB, Callahan LF. Variations in health status among rheumatic disease patients using the BRFSS healthrelated quality of life measure [abstract]. Arthritis Rheum 2000;43 Suppl:177.
- Scott DL, Garrood T. Quality of life measures: use and abuse. Bailliere's Clin Rheumatol 2000;14:663-87.
- Raina P, Dukeshire S, Lindsay J, Chambers LW. Chronic conditions and disabilities among seniors: an analysis of population-based health and activity limitation surveys. Ann Epidemiol 1998; 8:402-9
- 12. Hill CL, Parsons J, Taylor A, Leach G. Health-related quality of life in a population sample of arthritis. J Rheumatol 1999;26:2029-35.
- McHorney CA. Health status assessment methods for adults: past accomplishments and future challenges. Annu Rev Public Health 1999;20:309-35.
- Gill TM, Feinstein AR. A critical appraisal of the quality of life measurements. JAMA 1994;272:619-26.
- McDowell I, Newell C. Measuring health: A guide to rating scales and questionnaires. 2nd ed. New York: Oxford University Press; 1996
- Andresen EM, Meyers AR. Health-related quality of life outcomes measures. Arch Phys Med Rehabil 2000;81 Suppl 2:30-45.
- 17. Guillemin F. Functional disability and quality of life assessment in clinical practice. Rheumatology 2000;39:17-23.
- Burdine JN, Felix ME, Abel AL, Wiltraut CJ, Musselman YJ. The SF-12 as a population health measure: an exploratory examination of potential for application. Health Serv Res 2000;35:885-904.
- Cooper JK, Kohlmann T, Michael JA, Haffer SC, Stevic M. Health outcomes. New quality measure for medicare. Int J Qual Health Care 2001;13:9-16.
- Rao JK, Callahan LF, Helmick CG. Characteristics of persons with self-reported arthritis and other rheumatic conditions who do not see a doctor. J Rheumatol 1997;24:169-73.
- Hennessy CH, Moriarty DG, Zack MM, Scherr PA, Brackbill R. Measuring health-related quality of life for public health surveillance. Public Health Rep 1994;109:665-72.
- Ounpuu S, Kreuger P, Vermeulen M, Chambers L. Using the U.S. Behavioral Risk Factor Surveillance System's health-related quality of life survey tool in a Canadian city. Can J Public Health 2000;91:67-72.
- Centers for Disease Control and Prevention. Measuring healthy days: population assessment of health-related quality of life.
 Atlanta: Centers for Disease Control and Prevention; 2000.
 [Cited November 12, 2002] Available from http://www.cdc.gov/hrqol/monograph.htm
- CDC. Health-related quality of life among persons with epilepsy Texas, 1998. MMWR Morb Mortal Wkly Rep 2000;50:24-6.
- Ford ES, Moriarty DG, Zack MM, Mokdad AH, Chapman DP. Selfreported body mass index and health-related quality of life: findings from the Behavioral Risk Factor Surveillance System. Obesity Res 2001;9:21-31.
- Valdmanis V, Smith DW, Page MR. Productivity and economic burden associated with diabetes. Am J Public Health 2001; 91:129-30.

Personal, non-commercial use only. The Journal of Rheumatology Copyright © 2003. All rights reserved.

- Diwan S. Unmet health needs and the quality of life of the elderly: development of a survey for states and communities [technical report]. Atlanta: CDC; 1994.
- Calsyn RJ, Winter JP. Predicting four types of service needs in older adults. Eval Prog Planning 2001;24:157-66.
- Broyles RW, McAuley WJ, Baird-Holmes D. The medically vulnerable: their health risks, health status, and use of physician care. J Health Care Poor Underserved 1999;10:186-200.
- Mili F, Helmick CG, Zack MM. Prevalence of arthritis: Analysis of data from the Behavioral Risk Factor Surveillance System, US, 1996–99. J Rheumatol 2002;29:1981-8.
- Mariolis P. Data accuracy: how good are our usual indicators? In: Proceedings of 18th International Symposium on Methodological Issues, October 16–19, 2001; Ottawa. Ottawa: Statistics Canada; 2001
- CDC. Health risks in America: gaining insight from the Behavioral Risk Factor Surveillance System. Revised ed. Atlanta: CDC; 1997.
- World Health Organization. Physical status: the use and interpretation of anthropometry: report of a WHO expert committee. WHO technical report series; 854. Geneva: World Health Organization; 1995.
- Shah BV, Barnell BG, Bieler GS. SUDAAN: software for the statistical analysis of correlated data. User's manual, release 6.40 [software documentation]. Research Triangle Park: Research Triangle Institute; 1995.
- CDC. Health-related quality of life among adults with arthritis Behavioral Risk Factor Surveillance System, 11 states, 1996-1998. MMWR Morb Mortal Wkly Rep 2000;49:366-9.
- ViSta: The Visual Statistics System. [cited October 1, 2002]
 Available from: http://forrest.psych.unc.edu/research/vista-frames/help/lecturenotes/lecture02/repvis4a.html
- Newschaffer CJ. Validation of Behavioral Risk Factor Surveillance System (BRFSS) HRQOL measures in a statewide sample. Atlanta: CDC, National Center for Chronic Disease Prevention and Health Promotion; 1998.
- Ounpuu S, Chambers LW, Patterson C, Chan D, Yusuf S. Validity
 of the US Behavioral Risk Factor Surveillance System's health
 related quality of life survey tool in a group of older Canadians.
 Chron Dis Can 2001;22:93-101.
- Andresen EM, Fouts BS, Romeis JC, Brownson CA. Performance of health-related quality of life instruments in a spinal cord injured population. Arch Phys Med Rehabil 1999;80:877-84.

- Beatty P, Schechter S, Whitaker K. Evaluating subjective health questions: cognitive and methodological investigations. In: Proceedings of the Section on Survey Research Methods, 51st Annual Conference of the American Statistical Association, August 4-8, 1996; Chicago. Alexandria, VA: American Statistical Association, 1996:956-61. [Cited November 12, 2002] Available from http://www.amstat.org/sections/srms/proceedings/papers/ 1996_166.pdf
- Schechter S, Beatty P, Willis GB. Asking survey respondents about health status: Judgment and response issues. In: Schwarz N, Park DC, Knauper B, editors. Cognition, aging, and self-reports. Philadelphia: Psychology Press; 1999.
- 42. Dominick K, Gold C, Ahern F, Heller D. Cardiac drug use and health-related quality of life among older men and women [abstract]. Gerontologist 1999;39:244.
- Ahern F. Quality of life, medications, and health among the elderly (TS-213 Final report – Part I). State College, PA: Pennsylvania State University; 1999.
- Dominick K, Ahern F, Gold C, Heller D. Relationship of selfreported health conditions to health-related quality of life among the elderly [abstract]. Gerontologist 2000;40:93.
- CDC. Prevalence of arthritis Arizona, Missouri, and Ohio, 1991-1992. MMWR Morb Mortal Wkly Rep 1994;43:305-9.
- CDC. Prevalence and impact of chronic joint symptoms seven states, 1996. MMWR Morb Mortal Wkly Rep 1998;47:345-51.
- Lorig KR, Mazonson PD, Holman HR. Evidence suggesting that health education for self-management in patients with chronic arthritis has sustained health benefits while reducing health care cost. Arthritis Rheum 1993;36:439-46.
- Arthritis Foundation, Association of State and Territorial Health Officials, CDC. National Arthritis Action Plan: a public health strategy. Atlanta: Arthritis Foundation; 1999.
- CDC. Health-related quality of life and activity limitation eight states, 1995. MMWR Morb Mortal Wkly Rep 1998;47:134-40.
- US Department of Health and Human Services. Healthy People 2010. 2nd ed. Washington, DC: US Government Printing Office; 2000. [Cited October 1, 2002] Available from: http://www.health.gov/healthypeople/