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Osteoarthritis Cartilage. 2016 September ; 24(9): 1518–1527. doi:10.1016/j.joca.2016.04.012.**Annual incidence rates of hip symptoms and three hip OA outcomes from a U.S. population-based cohort study: the Johnston County Osteoarthritis Project****A.S. Moss[†], L.B. Murphy^{‡,*}, C.G. Helmick[‡], T.A. Schwartz[§], K.E. Barbour[‡], J.B. Renner^{||}, W. Kalsbeek[¶], and J.M. Jordan[#]**[†]Department of Mathematics and Statistics, Georgia State University, Atlanta, GA, USA[‡]Division of Population Health, Centers for Disease Control and Prevention, Atlanta, GA, USA[§]Department of Biostatistics, Gillings School of Global Public Health, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA^{||} Departments of Radiology and Allied Health Sciences, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA[¶]Carolina Survey Research Laboratory, Department of Biostatistics, Gillings School of Global Public Health, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA[#]Thurston Arthritis Research Center, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA**SUMMARY****Objective**—Estimate annual incidence rates (IRs) of hip symptoms and three osteoarthritis (OA) outcomes (radiographic, symptomatic, and severe radiographic) overall and by race, sociodemographic characteristics, and hip OA risk factors.**Design**—Analyze baseline (1991–1997) and first follow-up (1999–2003) data ($n = 1446$) from the Johnston County Osteoarthritis Project, a population-based, prospective study of adults 45 years in North Carolina. Hip symptoms were pain, aching, and/or stiffness on most days, or groin pain. Radiographic and severe radiographic OA were Kellgren–Lawrence (KL) grades 2 and 3, respectively. Symptomatic OA was radiographic OA with symptoms in the same hip. Sociodemographics were age, gender, race, highest attained education, and annual household income. Hip OA risk factors were self-reported body mass index (BMI) at age 18 years, clinically measured BMI at baseline, and history of hip injury.**Results**—Annual IRs (median = 5.5 years follow-up) were 37, 23, 13, and 2.9 per 1000 person-years for hip symptoms, and radiographic, symptomatic, and severe radiographic hip OA, respectively. We found low IRs of radiographic and symptomatic hip OA among African Americans and high IRs of hip symptoms among the obese and the very poor. Across outcomes, IRs were highest for those with hip injury.

* Address correspondence and reprint requests to: L.B. Murphy, Arthritis Program, Division of Population Health, Centers for Disease Control and Prevention, 4770 Buford Highway NE, Mailstop F78, Atlanta, GA 30341, USA. Tel: 1 770-488-5102; Fax: 1 770-488-5486. lmurphy1@cdc.gov (L.B. Murphy).

Conclusion—No prior studies have reported IRs of hip symptoms; IRs of radiographic and severe radiographic hip OA were similar to, and the IR of symptomatic hip OA was higher than, previous estimates. Prevention efforts should target low socioeconomic status (SES) populations and obese adults; interventions for hip OA and hip symptoms are imperative for those with hip injuries.

Keywords

Hip osteoarthritis; Hip symptoms; Race; Socioeconomic status; Incidence rates

Introduction

Hip osteoarthritis (OA) is a common cause of pain and disability among older adults and a predominant reason for total hip replacement (THR) surgery¹, which, in the US, is expected to grow to half a million procedures annually by 2030². Among adults aged ≥45 years, radiographic hip OA affects 27%, and symptomatic hip OA (defined as radiographic OA with hip symptoms) may affect 3–9%^{3,4}. Hip pain and other OA symptoms can substantially impair health and functioning, limit daily activities and reduce quality of life^{5,6}, and can significantly affect individuals financially through lost wages and need for informal care^{7,8}. Additionally, hip OA has been associated with an increased risk of mortality among older women⁹. These costs and the poor health associated with hip OA are part of a global public health problem that is predicted to only worsen¹⁰.

Epidemiological studies of incident hip OA have examined the role of demographics (women, older age, and white/African American race)^{11–15} and modifiable risk factors (hip injury and obesity)^{11,16–25}, but most of these studies have reported ratio measures (e.g., odds ratios). Descriptive studies quantifying the incidence rate (IR) – the rate at which new cases arise in the population – provide data that can be used for projecting prevalence and forecasting health service utilization and costs. Further, hip pain and symptoms can have a substantial health and economic impact, but no studies have reported IRs of hip pain/symptoms independently of radiographic hip OA, which can be poorly correlated with symptoms^{5,6,26}.

Of the few studies that have reported IRs of hip OA, some were in limited populations (e.g., white race or women only) and others reported only cumulative incidences²⁷, which may not adequately account for variation in observation time among participants. Furthermore, some US studies were conducted several decades ago and may not represent IRs in current populations given aging, increasing racial/ethnic diversity, and increasing prevalence of obesity in the US over past decades. Of the previous studies, two in the US estimated IRs for symptomatic hip OA from health records; Wilson *et al.* estimated an age and sex-adjusted rate of 47 per 100,000 person-years for the entire population of Rochester, MN in 1985¹², and Oliveria *et al.* estimated an age and sex-adjusted rate of 88 per 100,000 person-years among HMO members aged 20–89 in Worcester County, MA between 1998–1992¹¹. Three studies reported cumulative incidence of radiographic OA; from the Study of Osteoporotic Fractures, a multicenter US cohort of Caucasian women ≥65 years (mean follow-up 8 years), Lane *et al.* found 33% and 14% developed radiographic and severe radiographic OA (KL

3) respectively of the hip between 1986–1998²⁸; from the Rotterdam Study, a population-based Dutch cohort aged 55 (mean follow-up 6.6 years), Reijman *et al.* estimated 17% developed incident radiographic hip OA as defined by Kellgren–Lawrence (KL) grade 2, between 1990–1999²⁹; from the Johnston County OA Project (JoCo OA), Kopec *et al.* reported that radiographic OA developed in 7% of hips (joint-based analysis)¹³. Grotle *et al.* estimated 10-year cumulative incidence of self-reported hip OA of 6% in a population-based cohort ages 24–76 years in Ullensaker, Norway in 1994–2004.

Given these limited data on IRs of hip OA-related outcomes, we conducted a comprehensive hip OA incidence study using data from the JoCo OA, a large population-based, prospective cohort study in Johnston County, North Carolina. The purpose of this study was to quantify annual IRs of hip symptoms and three types of hip OA (radiographic, symptomatic, and severe radiographic hip OA). For three outcomes (hip symptoms, radiographic, symptomatic hip OA), we estimated overall IRs among African Americans and whites. For these three outcomes, we also estimated IRs by selected sociodemographic variables and hip OA risk factors, overall and for each race. Annual IRs for subgroups can be used to prioritize susceptible populations for further research and prevention efforts.

Method

We analyzed baseline (1991–1997) and first follow-up (1999–2003) data from the JoCo OA. The study's overall methodological approach is detailed elsewhere³⁰. Enrolled participants represented civilian, non-institutionalized African Americans and whites 45 years who were residents of one of six designated townships in Johnston County for at least 1 year and who were physically and mentally capable of study completion. At both baseline and first follow-up, participants completed two in-home interviews, approximately 2 weeks apart, and a clinical examination. Supine anterior–posterior pelvis radiographs for both hips were read using standard Kellgren–Lawrence (KL) grade (0–4)³¹. Pelvic radiographs were not obtained from women <50 years. The institutional review boards of the Centers for Disease Control and Prevention and the University of North Carolina Schools of Medicine and Public Health approved the study's protocol.

JoCo OA staff employed various methods to minimize attrition between baseline and first follow-up. Staff used annual newsletters, personal networks, local advertising, medical providers, and community inquiries to locate and retain participants. Deaths were identified through multiple sources including local obituaries, word-of-mouth, local and North Carolina death records, and the National Death Index (NDI), the most complete source of mortality data.

Measurement of hip symptoms and hip OA outcomes

Annual IRs were estimated for hip symptoms and three types of hip OA (radiographic, symptomatic, and severe radiographic hip OA). Hip symptoms were defined as either a “yes” to the question “On most days, do you have pain, aching, or stiffness in your (*right, left*) hip?” or reported the presence of (*right, left*) groin pain. Radiographic and severe radiographic OA were defined as KL grades of 2 and 3, respectively. Inflammatory arthritis occurred in 21 participants; joints with radiographic evidence of inflammatory

arthritis were treated as having missing KL grades and thereby excluded from analysis. Symptomatic OA was defined as having radiographic OA and hip symptoms in the same hip. Because symptomatic OA is the underlying cause in the majority of THRs¹, those with THR were classified as affected for all four outcomes, similar to previous studies^{13,32}.

For each outcome, an incident case was a participant who was not affected at baseline who developed the outcome in at least one hip by first follow-up (range 3–13 years). Baseline prevalent cases (i.e., participants having a given outcome in one or both hips at baseline) were excluded from analysis of that outcome, and therefore the number of participants varied for each outcome analysis (Table I). Analyses for hip symptoms included participants aged ≥45 years. Because women of reproductive age (<50 years) ($n = 312$) did not have hip radiographs, we also excluded men <50 years ($n = 175$) (Table I) so that the three hip OA outcome analyses comprised the same age groups for women and men.

Sociodemographic and hip OA risk factors

Sociodemographic characteristics (and their categorized levels) analyzed were age (45–54 or 50–54 [as appropriate], 55–64, 65–74, and ≥75 years), sex (men, women), race (African American, white), highest attained education (<high school [grade 12], some/completed high school [grade 12/GED {general equivalency diploma}], >high school [college/grad school] and annual household income (\$0–<15,000, \$15,000–<35,000, ≥\$35,000). The three hip OA risk factors were self-reported body mass index (BMI) in kg/m² at age 18 (under/healthy weight [<25], overweight/obese [≥25]) and clinically measured BMI at baseline (under/healthy weight [<25], overweight [25–<30], and obese [≥30] {Class I [30–<35] and Class II–IV [≥35]}), and history of hip injury (yes, no). Self-reported BMI at age 18 was based on participants' report of their weight at age 18. The denominator for all BMI calculations was clinically measured height at baseline. Hip injury was ascertained during the baseline clinic examination with “Have you ever injured your (*right, left*) hip?”.

Statistical analysis

Using person as the unit of analysis, we estimated annual IRs using a Poisson model with an offset (a term incorporated into the model to compute rates) taken as the natural logarithm of participant observation time (in years) to accommodate the variation in follow-up time among participants. We computed a crude IR (overall and stratified by sociodemographics and hip OA risk factors described above) with 95% confidence intervals (CI). For overall IRs, we also computed age- and age- and sex-standardized IRs using the 2000 projected US population with the same age categories that we analyzed (45–54 or 50–54 [as appropriate], 55–64, 65–74, and ≥75 years). We estimated race-specific IRs stratified by the same sociodemographics and hip OA risk factors using models that included race as an additional independent variable. We also tested for interactions with race but none was significant, most likely due to small sample sizes and low statistical power to detect these interactions. We did not compute stratified or race-specific IRs of severe radiographic hip OA because these estimates were unreliable due to the small number of unweighted cases ($n = 30$) (Table I).

To account for the complex sampling design, observations were weighted with population-calibrated sampling weights so that results are generalizable to the target Johnston County population. We calculated 95% CIs using jackknife methods that accounted for stratification and clustering in the sampling design with a finite population correction to adjust for sampling without replacement^{33,34}. Jackknifing produces accurate CIs even in the presence of overdispersion (i.e., variance greater than predicted by a Poisson distribution). We tested for differences in IRs using a Wald test with jackknife variance estimates that accounted for the sampling design. For age, education, self-reported BMI at age 18, and clinically measured BMI at baseline, we also tested for a trend through assessing the slope in models that included each of these characteristics as a continuous variable. Statistical significance was determined using a Bonferroni-corrected alpha of 0.0166 (<0.05/3 outcomes) to adjust for multiple comparisons across three outcomes (hip symptoms, and radiographic and symptomatic hip OA, but not severe radiographic hip OA owing to its lack of formal comparisons)³⁵. We performed statistical analyses with SAS version 9.3.

Sensitivity analysis

We performed sensitivity analyses for income imputation and selection bias following procedures detailed in Murphy *et al.*³⁶. We conducted these analyses to determine the extent to which missing income data and selection bias due to attrition may have influenced the IRs.

Income imputation

Based on weighted percentages, income was unknown for 18% of participants due to nonresponse (“refused” or “don’t know”), so we repeated the IR analyses using imputed income values to assess bias due to missing data. We performed multiple imputation with baseline variables for sociodemographics, hip OA risk factors/outcomes, and characteristics associated with income or income nonresponse. We found that analyses based on imputed income data produced similar IRs to those presented here. Thus any bias due to missing income data was likely small.

Selection bias

Because approximately 48% of eligible participants were excluded from IR estimation (Table I), we performed additional analyses to evaluate the extent of selection bias in the overall estimates. We computed approximate IRs for the analytic and eligible baseline populations (Table II)³⁶. For all outcomes we found that these approximate IRs were similar to each other. Thus differences between the analytic and baseline eligible populations appeared to have had little impact on IR estimates, and any potential selection bias was likely small.

Results

Of the 3068 participants in the JoCo OA sample at baseline, 2788 were potentially eligible for analysis after applying baseline exclusions (Table I). Of these 2788, about half ($n = 1446$) had both baseline and follow-up data for at least 1 of the 4 outcomes and were

included in the analytic sample (Table I). The median follow-up time for analyzed participants was 5.5 years (3–13 years).

Based on weighted percentages, the analytic sample represented a population at baseline that was over half women (58%) and predominantly white (80%) having a median age of 55 years (Table II). Most were married (73%) and had completed at least some high school (88%). About 25% had an annual household income under \$15,000, 28% had an income of \$35,000 or more, and income was unknown for 7% due to “don’t know” and 11% due to “refused” responses. At age 18, only 8% were overweight or obese whereas most were overweight (44%) or obese (25%) at the time of study baseline. Among those who were obese, almost a third were Class II–IV (BMI \geq 35). Only 3% had a history of hip injury.

Crude and standardized annual IRs

Crude and standardized annual IRs are presented in Table III. Crude IRs per 1000 person-years for hip symptoms, and radiographic, symptomatic, and severe radiographic hip OA were 37 (95% CI = 33, 41), 20 (95% CI = 18, 23), 13 (95% CI = 12, 16), and 2.9 (95% CI = 2.3, 3.7), respectively. Age-standardized and age- and sex-standardized IRs were higher than crude estimates (Table III).

Stratified and race-specific IRs

Annual IRs stratified by sociodemographic characteristics and hip OA risk factors are presented in Table IV, and race-specific annual IRs are presented in Table V, for the three outcomes of hip symptoms, radiographic OA, and symptomatic OA. We presented overall and race-specific IRs by age group in Figure 1.

Sociodemographic characteristics

Race—Across the three outcomes, IRs were lower among African Americans with significant differences for radiographic and symptomatic OA. The IR of radiographic OA was 24/1000 person-years for whites and 7/1000 person-years for African Americans, just over a quarter the rate of whites. The IR of symptomatic OA was 15/1000 person-years for whites and 7/1000 person-years for African Americans, less than half the rate of whites (Table IV).

Age—Across the three outcomes, IRs increased with greater baseline age, and trends were significant for radiographic and symptomatic OA. The IR of radiographic OA more than doubled from 12 to 27/1000 person-years between ages 50–54 and 55–64 and then remained steady among older ages. The annual IR of symptomatic hip OA almost tripled from 8 to 23/1000 person-years from youngest to oldest ages (Table IV, Fig. 1). Race-specific IRs for age showed similar trends (Table V, Fig. 1).

Sex—Across the three outcomes, IRs were higher among women than men with significant differences for hip symptoms and symptomatic OA, which were almost 50% higher among women (Table IV). Race-specific IRs for sex showed similar patterns (Table V).

Highest education—For hip symptoms and radiographic OA, IRs decreased moderately with greater educational attainment with a significant trend only for radiographic OA (Table IV). The trend in race-specific IRs of radiographic OA was also significant; compared with the overall sample, the decreasing trend appeared stronger among whites but was attenuated among African Americans (Table V).

Annual household income—The annual IR of 45/1000 person-years of hip symptoms for those with lowest income was among the highest rates observed across all subpopulations that we analyzed (Table IV). Across all three outcomes, race-specific IRs decreased with rising income, but differences were significant only for hip symptoms (Table V).

Hip OA risk factors

Self-reported BMI at age 18: The IR of hip symptoms was moderately higher among those with greater BMI at age 18, but there was no significant trend. Conversely for symptomatic OA, the IR among those who were overweight/obese was less than half the IR among those who were under/normal weight at age 18, and though the difference was significant, there was no significant trend. Across the three outcomes, the IRs of radiographic OA varied little (Table IV). Race-specific IRs of BMI at age 18 showed similar patterns (Table V).

Clinically measured BMI at baseline: For hip symptoms, the IR increased significantly with greater BMI at baseline; the IR of 48/1000 person-years among obese persons (>30) was high at twice the IR of 24/1000 person-years among under/healthy weight (<25) persons (Table IV). The IR of radiographic OA varied significantly across BMI levels but there was no significant trend. The IR of symptomatic OA varied little across BMI levels (Table IV). Race-specific IRs for BMI at baseline showed similar patterns (Table V).

History of hip injury: Across the three outcomes, those with hip injury had the highest IRs of all characteristics that we analyzed. Those with hip injury had higher IRs than those without hip injury, but these differences were not significant in the overall sample (Table IV). After stratifying by race, the difference was significant for symptomatic OA. Within each race, those with hip injury had double the IR of symptomatic OA (Table V).

Discussion

In this study of adults aged 45 years followed for a median of 5.5 years, IRs were 37, 20, 13, and 2.9 per 1000 person-years for hip symptoms, and radiographic, symptomatic, and severe radiographic hip OA, respectively. Overall and across sociodemographic subgroups and hip OA risk factors, the IRs of radiographic and symptomatic hip OA among African Americans were significantly lower than IRs for whites. For both African Americans and whites, the highest IRs of hip symptoms occurred among those aged 75 years, those with an annual household income <\$15,000, those who were obese, and those with history of hip injury. Those with hip injury also had highest IRs of radiographic and symptomatic hip OA, indicating the importance of OA prevention efforts among this subgroup.

Previous studies have assessed hip OA incidence data by race, age, and sex. Our findings of lower IRs of radiographic and symptomatic hip OA among African Americans are consistent with a joint-based (as opposed to person-based) analysis of incident radiographic hip OA in the JoCo OA cohort¹³. Across outcomes, IRs increased with older age with significant trends for radiographic and symptomatic OA. Although this same pattern has been found with radiographic OA^{20,21}, previous studies have shown declining IRs of incident symptomatic hip OA among the oldest ages^{11,12}. Differences in study populations and small numbers of oldest participants¹² likely contribute to variation in trends in oldest ages¹². Also reported previously^{11,12,37}, women had significantly higher IRs of hip symptoms and symptomatic hip OA than men.

This is among the first studies of socioeconomic status (SES) and hip OA, and it is the first study to quantify IRs of hip outcomes by household income and educational attainment. Lower education was associated with a higher IR of radiographic OA, and lowest income (< \$15,000) was associated with a higher IR of hip symptoms after adjusting for race. Two previous studies, one in the JoCo OA cohort, have also reported higher risk of hip OA among low SES populations^{38,39}, but our IR data further showed that those with lowest income had among the highest rates of hip symptoms across the large number of subpopulations that we analyzed. Regardless of cause, high IRs among low-SES populations are a concern given the substantial economic costs due to OA-associated hip symptoms⁷ and the disproportionate impact those costs may have on low-income households. Therefore, public health initiatives that prioritize low-SES populations and promote low-cost, evidence-based strategies (e.g., physical activity, weight loss, and self-management) are needed. Furthermore, conducting interventions and addressing barriers to uptake in low SES populations³⁸ may be an effective means of lowering incidence of hip symptoms.

Adults with greater BMI at baseline had a significantly higher IR of hip symptoms with a strong positive trend. We found wide variation in IRs of radiographic OA, with lowest rates among overweight persons, but no significant trend. The IRs of symptomatic hip OA varied little across levels of BMI. Similarly discordant patterns have been reported previously^{19,40}, and our findings for hip OA are consistent with studies reporting a lack of association with incident hip OA^{20–22,25,41}. Given the strong positive association between BMI and hip symptoms and the very high IR of hip symptoms among obese, focusing prevention among overweight/obese adults and emphasizing strategies to maintain healthy weight may reduce incident hip symptoms.

We found that a high BMI at age 18 did not predict the development of incident hip OA among those aged 50 years. In fact, those who were overweight/obese at age 18 had a lower IR of symptomatic hip OA, and further we found no significant trend between BMI at age 18 and onset of any hip outcomes. Variation in estimates and lack of significance was also found for self-reported hip OA with clinically measured and self-reported BMI in a previous study among young men¹⁶. Characteristics that we did not analyze such as life-course BMI^{42–45} and other population characteristics (e.g., sex, age, and BMI at baseline) may explain the patterns in IRs that we observed.

Consistent with previous studies, hip injury was associated with a higher IR of symptomatic hip OA^{17,18,22,24} after adjusting for the effect of race. For the three hip outcomes, those with hip injury had highest IRs across all characteristics that we analyzed. Therefore targeted interventions for hip OA prevention and self-management may greatly benefit those with hip injuries.

Compared with previous studies, the overall IRs of radiographic and severe radiographic hip OA are consistent with annual rates reported previously²⁷⁻²⁹. We found no other studies reporting IRs of hip pain/symptoms to compare with IRs of hip symptoms. Our overall IR of symptomatic hip OA was substantially higher than previous IR estimates based on health records of inpatient and outpatient visits for clinical care^{11,12}, which may be conservative since a majority of adults with joint pain do not seek care for their symptoms^{46,47}. Differences in study population characteristics and OA definitions also likely contribute to differences in estimates.

This study has several limitations. First, similar to most longitudinal studies, the results are in a small geographic area and may have limited generalizability to other populations. Second, death and loss to follow-up are common for longitudinal studies in older populations, particularly with longer follow-up intervals as in this study, and attrition in the JoCo OA was not trivial. Although we determined that selection bias resulting from attrition was likely small, there remains the possibility that we may not have accounted for other types of bias potentially associated with attrition. Third, because we estimated incident hip symptoms and hip OA outcomes, relatively rare conditions, we had insufficient numbers of cases and low power to detect some effects (e.g., hip injury) and assess differential patterns of association (i.e., interactions) by race.

This study also has several important strengths. First, we conducted a comprehensive, descriptive analysis of incident hip outcomes from which we have presented IRs of four hip OA-related outcomes for a large population-based prospective study. We estimated IRs for African Americans and whites, and we identified dramatically lower IRs of hip OA for African Americans, overall and across all subpopulations that we analyzed. These population-based IRs provide relatively more recent data to improve projections of costs, prevalence, and demand for health services in increasingly diverse populations. Secondly, we estimated IRs of hip symptoms which can substantially impair health even in the absence of radiographic signs of hip OA⁵. Knowing these IRs can help clinicians and public health professionals recognize additional high risk groups for intervention opportunities to reduce and prevent hip symptoms. Thirdly, we quantified IRs overall and by numerous sociodemographic characteristics and hip OA risk factors. Quantifying IRs can identify subpopulations in which the prevalence and consequent burden of hip OA is growing most rapidly in addition to determining those at increased risk of hip OA-related outcomes.

The IRs we present provide valuable insights to clinicians and public health practitioners in their ongoing efforts to reduce the future impact of hip OA. In the JoCo OA cohort, the IR of hip symptoms was high, and, compared with earlier estimates, we found substantially higher IRs of symptomatic hip OA. The IRs of radiographic and severe radiographic hip OA were similar to previous estimates. African Americans had low IRs of radiographic and

symptomatic hip OA compared with whites. In addition to higher IRs among women (hip symptoms and symptomatic OA), oldest ages (radiographic and symptomatic OA), and those with hip injury (symptomatic OA), those with lower education had a higher IR of radiographic OA, and several of the highest IRs of hip symptoms occurred among those with lowest income and greatest BMI. Across outcomes, IRs were highest among those with a hip injury. Public health efforts should prioritize low-SES populations, obese adults, and those with a hip injury for further action and prevention of hip symptoms and hip OA.

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Disclaimer

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Appendix

Contributions

A Susan Moss: conception and design; analysis and interpretation of the data; drafting of the article; critical revision of the article for important intellectual content; final approval of the article; statistical expertise; responsibility for the integrity of the work as a whole, from inception to finished article zbr3@cdc.gov.

Louise B Murphy: conception and design; analysis and interpretation of the data; critical revision of the article for important intellectual content; final approval of the article; responsibility for the integrity of the work as a whole, from inception to finished article lmurphy1@cdc.gov.

Charles G Helmick: conception and design; analysis and interpretation of the data; critical revision of the article for important intellectual content; final approval of the article; administrative, technical, or logistic support.

Todd A Schwartz: conception and design; analysis and interpretation of the data; critical revision of the article for important intellectual content; final approval of the article; statistical expertise; administrative, technical, or logistic support.

Kamil E Barbour: conception and design; analysis and interpretation of the data; critical revision of the article for important intellectual content; final approval of the article.

Jordan B Renner: critical revision of the article for important intellectual content; final approval of the article; collection and assembly of the data.

William Kalsbeek: conception and design; analysis and interpretation of the data; critical revision of the article for important intellectual content; final approval of the article; statistical expertise.

Joanne M Jordan: conception and design; analysis and interpretation of the data; critical revision of the article for important intellectual content; final approval of the article; provision of study materials or patients; obtaining of funding; administrative, technical, or logistic support; collection and assembly of the data.

Ethics

The procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the Helsinki Declaration of 1975, as revised in 2000. The study was approved by the Institutional Review Boards of the University of North Carolina Schools of Medicine and Public Health, and the Centers for Disease Control and Prevention. All participants gave written informed consent at recruitment and follow-up.

Competing interests

Ms. Moss has nothing to disclose.

Dr. Murphy has nothing to disclose.

Dr. Helmick has nothing to disclose.

Dr. Schwartz has nothing to disclose.

Dr. Barbour has nothing to disclose.

Dr. Renner has nothing to disclose.

Dr. Kalsbeek has nothing to disclose.

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References

1. Katz JN, Losina E, Barrett J, Phillips CB, Mahomed NN, Lew RA, et al. Association between hospital and surgeon procedure volume and outcomes of total hip replacement in the United States medicare population. *J Bone Joint Surg Am*. 2001; 83-A:1622–9. [PubMed: 11701783]
2. Kurtz S, Ong K, Lau E, Mowat F, Halpern M. Projections of primary and revision hip and knee arthroplasty in the United States from 2005 to 2030. *J Bone Joint Surg Am*. 2007; 89:780–5. [PubMed: 17403800]
3. Lawrence RC, Felson DT, Helmick CG, Arnold LM, Choi H, Deyo RA, et al. Estimates of the prevalence of arthritis and other rheumatic conditions in the United States. Part II Arthritis Rheum. 2008; 58:26–35. [PubMed: 18163497]
4. Jordan JM, Helmick CG, Renner JB, Luta G, Dragomir AD, Woodard J, et al. Prevalence of hip symptoms and radiographic and symptomatic hip osteoarthritis in African Americans and Caucasians: the Johnston County Osteoarthritis Project. *J Rheumatol*. 2009; 36:809–15. [PubMed: 19286855]
5. Birrell F, Croft P, Cooper C, Hosie G, Macfarlane G, Silman A. Health impact of pain in the hip region with and without radiographic evidence of osteoarthritis: a study of new attenders to primary care. The PCR Hip Study Group *Ann Rheum Dis*. 2000; 59:857–63.
6. Odding E, Valkenburg HA, Algra D, Vandenouweland FA, Grobbee DE, Hofman A. Associations of radiological osteoarthritis of the hip and knee with locomotor disability in the Rotterdam Study. *Ann Rheum Dis*. 1998; 57:203–8. [PubMed: 9709175]
7. Gupta S, Hawker GA, Laporte A, Croxford R, Coyte PC. The economic burden of disabling hip and knee osteoarthritis (OA) from the perspective of individuals living with this condition. *Rheumatology (Oxford)*. 2005; 44:1531–7. [PubMed: 16091394]
8. Gabriel SE, Crowson CS, Champion ME, O’Fallon WM. Indirect and nonmedical costs among people with rheumatoid arthritis and osteoarthritis compared with nonarthritic controls. *J Rheumatol*. 1997; 24:43–8. [PubMed: 9002009]
9. Barbour KE, Lui LY, Nevitt MC, Murphy LB, Helmick CG, Theis KA, et al. Hip osteoarthritis and the risk of all-cause and disease-specific mortality in older women: population-based cohort study. *Arthritis Rheumatol*. 2015; 73:1798–805.
10. Cross M, Smith E, Hoy D, Nolte S, Ackerman I, Fransen M, et al. The global burden of hip and knee osteoarthritis: estimates from the global burden of disease 2010 study. *Ann Rheum Dis*. 2014; 73:1323–30. [PubMed: 24553908]
11. Oliveria SA, Felson DT, Reed JI, Cirillo PA, Walker AM. Incidence of symptomatic hand, hip, and knee osteoarthritis among patients in a health maintenance organization. *Arthritis Rheum*. 1995; 38:1134–41. [PubMed: 7639811]
12. Wilson MG, Michet CJ Jr, Ilstrup DM, Melton LJ 3rd. Idiopathic symptomatic osteoarthritis of the hip and knee: a population-based incidence study. *Mayo Clin Proc*. 1990; 65:1214–21. [PubMed: 2402161]
13. Kopec JA, Sayre EC, Schwartz TA, Renner JB, Helmick CG, Badley EM, et al. Occurrence of radiographic osteoarthritis of the knee and hip among African Americans and whites: a population-based prospective cohort study. *Arthritis Care Res (Hoboken)*. 2013; 65:928–35. [PubMed: 23281251]
14. Scher DL, Belmont PJ Jr, Mountcastle S, Owens BD. The incidence of primary hip osteoarthritis in active duty US military servicemembers. *Arthritis Rheum*. 2009; 61:468–75. [PubMed: 19333991]
15. Prieto-Alhambra D, Judge A, Javaid MK, Cooper C, Diez-Perez A, Arden NK. Incidence and risk factors for clinically diagnosed knee, hip and hand osteoarthritis: influences of age, gender and osteoarthritis affecting other joints. *Ann Rheum Dis*. 2014; 73:1659–64. [PubMed: 23744977]
16. Gelber AC, Hochberg MC, Mead LA, Wang NY, Wigley FM, Klag MJ. Body mass index in young men and the risk of subsequent knee and hip osteoarthritis. *Am J Med*. 1999; 107:542–8. [PubMed: 10625021]

17. Gelber AC, Hochberg MC, Mead LA, Wang NY, Wigley FM, Klag MJ. Joint injury in young adults and risk for subsequent knee and hip osteoarthritis. *Ann Intern Med.* 2000; 133:321–8. [PubMed: 10979876]
18. Felson DT, Lawrence RC, Dieppe PA, Hirsch R, Helmick CG, Jordan JM, et al. Osteoarthritis: new insights. Part 1: the disease and its risk factors. *Ann Intern Med.* 2000; 133:635–46. [PubMed: 11033593]
19. Lieveense AM, Bierma-Zeinstra SM, Verhagen AP, van Baar ME, Verhaar JA, Koes BW. Influence of obesity on the development of osteoarthritis of the hip: a systematic review. *Rheumatology (Oxford).* 2002; 41:1155–62. [PubMed: 12364636]
20. Reijman M, Pols HA, Bergink AP, Hazes JM, Belo JN, Lieveense AM, et al. Body mass index associated with onset and progression of osteoarthritis of the knee but not of the hip: the Rotterdam Study. *Ann Rheum Dis.* 2007; 66:158–62. [PubMed: 16837490]
21. Grotle M, Hagen KB, Natvig B, Dahl FA, Kvien TK. Obesity and osteoarthritis in knee, hip and/or hand: an epidemiological study in the general population with 10 years follow-up. *BMC Musculoskelet Disord.* 2008; 9:132. [PubMed: 18831740]
22. Juhakoski R, Heliövaara M, Impivaara O, Kroger H, Knekt P, Lauren H, et al. Risk factors for the development of hip osteoarthritis: a population-based prospective study. *Rheumatology (Oxford).* 2009; 48:83–7. [PubMed: 19056801]
23. Lohmander LS, Gerhardsson de Verdier M, Roloff J, Nilsson PM, Engstrom G. Incidence of severe knee and hip osteoarthritis in relation to different measures of body mass: a population-based prospective cohort study. *Ann Rheum Dis.* 2009; 68:490–6. [PubMed: 18467514]
24. Richmond SA, Fukuchi RK, Ezzat A, Schneider K, Schneider G, Emery CA. Are joint injury, sport activity, physical activity, obesity, or occupational activities predictors for osteoarthritis? A systematic review *J Orthop Sports Phys Ther.* 2013; 43:515–B519. [PubMed: 23756344]
25. Mork PJ, Holtermann A, Nilsen TI. Effect of body mass index and physical exercise on risk of knee and hip osteoarthritis: longitudinal data from the Norwegian HUNT Study. *J Epidemiol Community Health.* 2012; 66:678–83. [PubMed: 22511797]
26. Birrell F, Lunt M, Macfarlane G, Silman A. Association between pain in the hip region and radiographic changes of osteoarthritis: results from a population-based study. *Rheumatology (Oxford).* 2005; 44:337–41. [PubMed: 15536064]
27. Pereira D, Peleteiro B, Araujo J, Branco J, Santos RA, Ramos E. The effect of osteoarthritis definition on prevalence and incidence estimates: a systematic review. *Osteoarthritis Cartilage.* 2011; 19:1270–85. [PubMed: 21907813]
28. Lane NE, Lin P, Christiansen L, Gore LR, Williams EN, Hochberg MC, et al. Association of mild acetabular dysplasia with an increased risk of incident hip osteoarthritis in elderly white women: the study of osteoporotic fractures. *Arthritis Rheum.* 2000; 43:400–4. [PubMed: 10693881]
29. Reijman M, Hazes JM, Pols HA, Koes BW, Bierma-Zeinstra SM. Acetabular dysplasia predicts incident osteoarthritis of the hip: the Rotterdam Study. *Arthritis Rheum.* 2005; 52:787–93. [PubMed: 15751071]
30. Jordan JM, Helmick CG, Renner JB, Luta G, Dragomir AD, Woodard J, et al. Prevalence of knee symptoms and radiographic and symptomatic knee osteoarthritis in African Americans and Caucasians: the Johnston County Osteoarthritis Project. *J Rheumatol.* 2007; 34:172–80. [PubMed: 17216685]
31. Kellgren JH, Lawrence JS. Radiological assessment of osteoarthrosis. *Ann Rheum Dis.* 1957; 16:494–502. [PubMed: 13498604]
32. Murphy LB, Helmick CG, Schwartz TA, Renner JB, Tudor G, Koch GG, et al. One in four people may develop symptomatic hip osteoarthritis in his or her lifetime. *Osteoarthritis Cartilage.* 2010; 18:1372–9. [PubMed: 20713163]
33. Shao, J., Tu, D. *The Jackknife and Bootstrap.* New York, NY, USA: Springer Verlag; 1995.
34. Rust KF, Rao JN. Variance estimation for complex surveys using replication techniques. *Stat Methods Med Res.* 1996; 5:283–310. [PubMed: 8931197]
35. Miller, RG. *Simultaneous Statistical Inference.* 2nd. New York: Springer-Verlag; 1981.

36. Murphy LB, Moss S, Do BT, Helmick CG, Schwartz TA, Barbour KE, et al. Annual incidence of knee symptoms and four knee osteoarthritis outcomes in the Johnston County Osteoarthritis Project. *Arthritis Care Res (Hoboken)*. 2016; 68:55–65. [PubMed: 26097226]
37. Thiem U, Lamsfuss R, Gunther S, Schumacher J, Baker C, Endres HG, et al. Prevalence of self-reported pain, joint complaints and knee or hip complaints in adults aged \geq 40 years: a cross-sectional survey in Herne, Germany. *PLoS One*. 2013; 8:e60753. [PubMed: 23646102]
38. Cleveland RJ, Schwartz TA, Prizer LP, Randolph R, Schoster B, Renner JB, et al. Associations of educational attainment, occupation, and community poverty with hip osteoarthritis. *Arthritis Care Res (Hoboken)*. 2013; 65:954–61. [PubMed: 23225374]
39. Reyes C, Garcia-Gil M, Elorza JM, Mendez-Boo L, Hermosilla E, Javaid MK, et al. Socio-economic status and the risk of developing hand, hip or knee osteoarthritis: a region-wide ecological study. *Osteoarthritis Cartilage*. 2015; 23:1323–9. [PubMed: 25819582]
40. Lubbeke A, Duc S, Garavaglia G, Finckh A, Hoffmeyer P. BMI and severity of clinical and radiographic signs of hip osteoarthritis. *Obesity (Silver Spring)*. 2009; 17:1414–9. [PubMed: 19197252]
41. Oliveria SA, Felson DT, Cirillo PA, Reed JI, Walker AM. Body weight, body mass index, and incident symptomatic osteoarthritis of the hand, hip, and knee. *Epidemiology*. 1999; 10:161–6. [PubMed: 10069252]
42. Apold H, Meyer HE, Espehaug B, Nordsletten L, Havelin LI, Flugsrud GB. Weight gain and the risk of total hip replacement a population-based prospective cohort study of 265,725 individuals. *Osteoarthritis Cartilage*. 2011; 19:809–15. [PubMed: 21524707]
43. Holliday KL, McWilliams DF, Maciewicz RA, Muir KR, Zhang W, Doherty M. Lifetime body mass index, other anthropometric measures of obesity and risk of knee or hip osteoarthritis in the GOAL case-control study. *Osteoarthritis Cartilage*. 2011; 19:37–43. [PubMed: 21044695]
44. Wang Y, Wluka AE, Simpson JA, Giles GG, Graves SE, de Steiger RN, et al. Body weight at early and middle adulthood, weight gain and persistent overweight from early adulthood are predictors of the risk of total knee and hip replacement for osteoarthritis. *Rheumatology (Oxford)*. 2013; 52:1033–41. [PubMed: 23362222]
45. Abbate LM, Jordan JM. Weight change in osteoarthritis. *Osteoarthritis Cartilage*. 2012; 20:268–70. [PubMed: 22178466]
46. Thorstensson CA, Gooberman-Hill R, Adamson J, Williams S, Dieppe P. Help-seeking behaviour among people living with chronic hip or knee pain in the community. *BMC Musculoskelet Disord*. 2009; 10:153. [PubMed: 19968876]
47. Paskins Z, Sanders T, Hassell AB. What influences patients with osteoarthritis to consult their GP about their symptoms? A narrative review *BMC Fam Pract*. 2013; 14:195. [PubMed: 24359101]

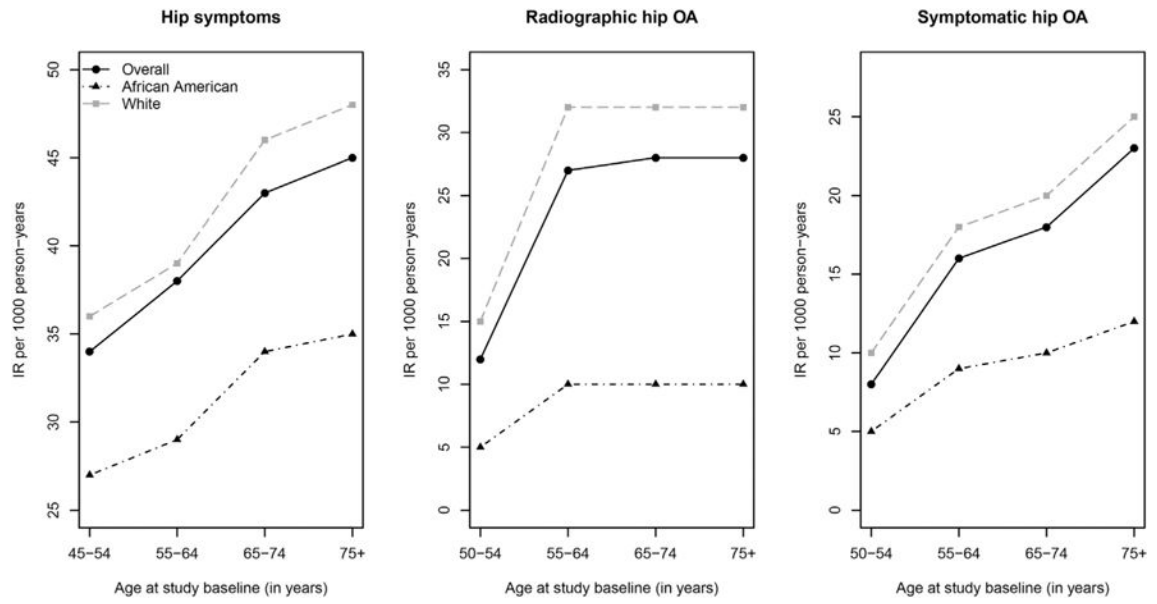


Fig. 1. Incidence rate (IR) per 1000 person-years for hip symptoms and OA outcomes by age group, overall and within each race.

Unweighted sample sizes in the entire baseline, eligible baseline, and analytic samples with incident cases, by hip symptoms and three hip OA outcomes

Table 1

	All analyses*	Hip OA outcomes		
		Hip symptoms	Radiographic	Severe radiographic
Entire baseline sample (1991–1997)	3068	3068	3068	3068
Outcome at baseline		1228	803	72
Participants at risk at baseline		1840	2265	2996
Other baseline exclusions				
Hip replacement		24	24	24
Women <50 years		0	312	312
Men <50 years		0	175	175
Missing baseline		72	58	60
Eligible baseline sample	2788	1744	1696	2425
Participant exclusions at 1995–2003 follow-up [†]				
Deceased		200	223	338
Mentally/physically unable		98	123	181
Moved out of study area		129	119	176
Declined participation		276	228	296
No clinic exam		75	83	107
Unable to locate		30	38	55
Eligible at follow-up		936	865	1242
Missing outcome at follow-up		0	17	30
Analytic sample	1446	936	864	1212
Incident cases at follow-up		200	107	30

* 3068 were enrolled at baseline; 2788 were eligible for analysis of at least 1 of the 4 outcomes; 1446 were analyzed for at least 1 of the 4 outcomes.

[†] Participants who were excluded were those who died (deceased), were mentally or physically unable to participate, moved out of the study area, declined to participate, participated in the in-home questionnaire but not the clinic exam (no clinic exam), or were lost to follow up for other reasons (unable to locate).

Table II

Weighted* distribution (%)† of baseline sociodemographic characteristics and hip OA risk factors for the analytic and eligible baseline samples

	Analytic sample <i>n</i> = 1446	Eligible baseline sample <i>n</i> = 2778
Sociodemographic characteristics		
Age (years)		
45–54	52	30
55–64	26	29
65–74	18	28
75	5	13
Median	55	62
Sex		
Men	42	43
Women	58	57
Race		
African American	20	18
White	80	82
Marital status		
Never married	4	3
Married	73	66
Separated/divorced	9	10
Widowed	13	22
Highest education		
<High school	11	21
Some/completed high school	54	52
>High school	34	28
Annual household income		
\$0–<\$15,000	25	35
\$15,000–<\$35,000	29	27
\$35,000	28	17
Don't know	7	9
Refused	11	12
Hip OA risk factors		
Self-reported BMI at age 18 (kg/m ²)		
Under or healthy weight (<25)	92	90
Overweight or obese (≥ 25)	8	10
Clinically measured BMI at study baseline (kg/m ²)		
Under or healthy weight (<25)	31	34
Overweight (25–<30)	44	42
Obese (≥ 30)	25	24
Obese Class I (30–<35)	17	17
Obese Class II–IV (≥ 35)	7	6

	Analytic sample $n = 1446$	Eligible baseline sample $n = 2778$
History of hip injury		
No	97	95
Yes	3	5

* The analytic and eligible samples were weighted to the Johnston County population in 2000 and 1990, respectively. Differences in weighted numbers are due to the use of different sampling weights applied to the baseline and follow-up samples.

† Percentages may not sum to 100% due to rounding.

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Overall crude, age-standardized, and age- and sex-standardized IRs per 1000 person-years and their 95% CIs for hip symptoms and three hip OA outcomes

Table III

	Hip symptoms		OA outcomes			
	IR (95% CI)		Radiographic IR (95% CI)	Symptomatic IR (95% CI)	Severe radiographic IR (95% CI)	
Crude	37 (33, 41)		20 (18, 23)	13 (12, 16)	2.9 (2.3, 3.7)	
Age-standardized*	39 (34, 44)		24 (20, 30)	17 (14, 20)	3.2 (2.5, 4.1)	
Age- and sex-standardized*	38 (34, 43)		24 (20, 30)	17 (14, 19)	3.2 (2.5, 4.1)	

* Standardized to the 2000 projected US population.

IRs per 1000 person-years and their 95% CIs for hip symptoms and OA outcomes by sociodemographic characteristics and hip OA risk factors

Table IV

	Hip symptoms			OA		
				Symptomatic		
	IR (95% CI)	P value* (P_{Trend}^{\ddagger})	P value* (P_{Trend}^{\ddagger})	IR (95% CI)	IR (95% CI)	P value* (P_{Trend}^{\ddagger})
Sociodemographic characteristics						
Race		0.082	<0.001 [‡]			<0.001 [‡]
African American	28 (20, 39)			7 (5, 10)	7 (5, 11)	
White	39 (34, 44)			24 (21, 27)	15 (13, 18)	
Age (years)		0.31	<0.001 [‡]			<0.001 [‡]
45 – 54/50 – 54 [§]	34 (28, 41)	(0.018)	(<0.001 [‡])	12 (10, 16)	8 (6, 12)	(<0.001 [‡])
55–64	38 (32, 44)			27 (22, 34)	16 (13, 20)	
65–74	43 (36, 52)			28 (22, 36)	18 (14, 24)	
75	45 (28, 72)			28 (14, 52)	23 (16, 33)	
Sex		0.008 [‡]	0.23			0.013 [‡]
Men	30 (24, 37)			19 (15, 23)	11 (8, 14)	
Women	42 (36, 48)			22 (19, 27)	16 (14, 19)	
Highest education		0.48	0.25			0.81
<High school (HS)	40 (31, 50)	(0.030)	(0.016 [‡])	22 (16, 30)	14 (10, 20)	(0.98)
Some/completed HS	38 (32, 45)			22 (18, 26)	13 (10, 16)	
>HS	33 (26, 41)			17 (13, 22)	14 (11, 19)	
Annual household income		0.022	0.57			0.76
\$0–<\$15,000	45 (37, 53)			20 (17, 24)	14 (12, 17)	
\$15,000–<\$35,000	34 (27, 42)			22 (18, 28)	13 (10, 18)	
\$35,000	28 (21, 37)			18 (13, 25)	12 (8, 18)	
Hip OA risk factors						
Self-reported BMI(kg/m ²) at age 18		0.40	0.67			0.005 [‡]
Underweight/Normal(<25)	35 (31, 40)	(0.19)	(0.21)	21 (18, 24)	14 (12, 17)	(0.10)
Overweight/Obese (≥ 25)	43 (28, 66)			19 (14, 28)	5 (3, 10)	
Clinically measured BMI (kg/m ²) at baseline		<0.001 [‡]	<0.001 [‡]			0.73

	OA					
	Hip symptoms			Symptomatic		
	IR	(95% CI)	P value* (P _{Trend} [†])	IR	(95% CI)	P value* (P _{Trend} [†])
Under/healthy (<25)	24	(19, 29)	(0,013) [‡]	29	(23, 36)	(0,19)
Overweight (25–<30)	41	(34, 48)		16	(13, 20)	
Obese (30)	48	(37, 61)		20	(16, 26)	
Obese Class I (30–<35)	49	(37, 65)		17	(12, 24)	
Obese Class II–IV (35)	44	(25, 77)		27	(19, 38)	
History of hip injury			0.41			0.26
No	36	(3.2, 4.0)		20	(18, 23)	
Yes	51	(2.2, 11.8)		28	(16, 52)	

* P-value for differences in IRs.

[†] P-value for trend in IRs; performed through assessment of the slope for continuous variables for age, highest education, self-reported BMI at age 18, and clinically measured BMI at baseline.

[‡] Statistically significant at Bonferroni-corrected $\alpha = 0.0166$.

[§] Hip symptoms: 45–54 years; radiographic and symptomatic OA: 50–54 years.

Table V

Race-specific IRs per 1000 person-years and their 95% CIs for hip symptoms and OA outcomes by sociodemographic characteristics and hip OA risk factors

	Hip symptoms				OA outcomes				
	African American		White		African American		White		
	IR (95% CI)	P value† (P _{Trend} ‡)	IR (95% CI)	P value† (P _{Trend} ‡)	IR (95% CI)	P value† (P _{Trend} ‡)	IR (95% CI)	P value† (P _{Trend} ‡)	
Overall									
Crude	28 (20, 39)		39 (34, 44)		7 (5, 10)		24 (21, 27)		15 (13, 18)
Age-standardized*	30 (22, 42)		41 (36, 47)		9 (7, 13)		28 (23, 34)		19 (16, 22)
Age- and sex-standardized*	30 (22, 41)		41 (36, 47)		9 (7, 13)		29 (24, 36)		19 (16, 22)
Sociodemographic characteristics									
Age (years)		0.32							
45–54/50–54 //	27 (18, 38)	(0.025)	36 (30, 43)		5 (3, 7)		15 (11, 19)		5 (3, 8)
55–64	29 (20, 41)		39 (33, 46)		10 (7, 14)		32 (26, 39)		9 (6, 13)
65–74	34 (24, 47)		46 (38, 55)		10 (7, 15)		32 (25, 42)		10 (6, 15)
75	35 (21, 60)		48 (30, 76)		10 (5, 20)		32 (17, 59)		12 (7, 21)
Sex		0.008\$							0.014\$
Men	23 (16, 34)		32 (26, 39)		7 (5, 10)		22 (18, 27)		6 (4, 9)
Women	32 (23, 45)		44 (38, 51)		8 (6, 11)		26 (22, 31)		9 (6, 13)
Highest education		0.36							0.12
<High school (HS)	31 (22, 45)	(0.022)	44 (34, 57)		8 (5, 13)		28 (20, 38)		8 (5, 12)
Some/completed HS	29 (20, 42)		41 (34, 49)		8 (6, 11)		26 (22, 31)		6 (4, 10)
>HS	25 (16, 37)		35 (28, 43)		6 (4, 8)		19 (14, 25)		7 (4, 11)
Annual household income		0.005\$							0.39
\$0–<\$15,000	32 (23, 46)		50 (42, 60)		8 (6, 10)		25 (21, 31)		7 (5, 11)
\$15,000–<\$35,000	23 (16, 34)		36 (28, 45)		8 (5, 11)		25 (20, 32)		6 (4, 11)
\$35,000	19 (12, 29)		29 (22, 39)		6 (4, 9)		20 (14, 27)		6 (3, 10)
Hip OA risk factors									

	Hip symptoms						OA outcomes					
	African American			White			African American			White		
	IR (95% CI)	IR (95% CI)	P value† (P _{Trend} ‡)	IR (95% CI)	IR (95% CI)	P value† (P _{Trend} ‡)	IR (95% CI)	IR (95% CI)	P value† (P _{Trend} ‡)	IR (95% CI)	IR (95% CI)	P value† (P _{Trend} ‡)
Self-reported BMI at age 18 (kg/m ²)			0.37			0.84						0.008§
Under/healthy (<25)	25 (18, 34)	38 (33, 43)	(0.11)	8 (6, 11)	24 (21, 28)	(0.45)	8 (5, 12)	16 (13, 19)				(0.19)
Overweight/obese (≥ 25)	30 (16, 59)	47 (30, 72)	<0.001§	8 (5, 13)	23 (16, 33)		3 (1, 6)	6 (3, 12)				0.94
Clinically measured BMI at baseline (kg/m ²)												
Under/healthy weight (<25)	16 (11, 23)	25 (20, 31)	(<0.004§)	11 (7, 16)	32 (25, 39)	(0.50)	7 (4, 12)	15 (12, 19)				(0.71)
Overweight (25–<30)	29 (20, 41)	44 (37, 53)		6 (4, 9)	19 (15, 23)		7 (4, 11)	15 (12, 20)				
Obese (≥ 30)	34 (23, 51)	53 (41, 68)		8 (6, 11)	24 (19, 32)		7 (4, 10)	14 (11, 19)				
Obese Class I (30–<35)	35 (24, 51)	54 (40, 71)		7 (5, 10)	21 (15, 30)		7 (4, 11)	15 (11, 21)				
Obese Class II–IV (≥ 35)	33 (16, 67)	51 (29, 90)		11 (7, 17)	34 (23, 50)		6 (3, 11)	12 (7, 21)				
History of hip injury			0.28			0.14						0.011§
No	24 (18, 34)	38 (34, 43)		7 (5, 10)	24 (21, 27)		7 (4, 10)	15 (13, 18)				
Yes	38 (17, 86)	60 (27, 134)		12 (6, 21)	36 (20, 65)		15 (7, 29)	33 (19, 59)				

* Standardized to the 2000 projected US population.

† P-value for differences in IRs.

‡ P-value for trend in IRs; performed through assessment of the slope for continuous variables for age, highest education, self-reported BMI at age 18, and clinically measured BMI at baseline.

§ Statistically significant at Bonferroni-corrected $\alpha = 0.0166$.

// Hip symptoms: 45–54 years; radiographic and symptomatic OA: 50–54 years.