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# Social determinants and osteoarthritis outcomes

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# Abstract

Osteoarthritis (OA) is one of the most frequently occurring musculoskeletal diseases, posing a significant public health problem due to its impact on pain and disability. Traditional risk factors fail to account for all of the risk observed for OA outcomes. In recent years, our view of disease causation has broadened to include health risks that are created by an individual's socioeconomic circumstances. Early research into social determinants has focused on social position and explored factors related to the individual such as education, income and occupation. Results from these investigations suggest that low education attainment and nonprofessional occupation are associated with poorer arthritis outcomes. More recently, research has expanded to examine how one's neighborhood socioeconomic environment may be relevant to OA outcomes. This narrative review proposes a framework to help guide our understanding of how social context may interact with pathophysiological processes and individual-level variables to influence health outcomes in those living with OA.

### Keywords

osteoarthritis; sociall determinant; socioeconomic status

Despite the growing capacity of societies to extend life and prevent disease, the disparities in health between and within countries remain ubiquitous and vexing [1–4]. Historically, society has attributed disparities in health and disease to the medical sector. While access to medical care is certainly relevant to one's health, the literature over the past 15–20 years on the social and environmental influences on people's health has invariably shown that socioeconomically disadvantaged populations experience worse health outcomes than more affluent populations [5–7]. The disproportionate burden of illness experienced by these

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populations is a result of structural inequalities reflected in the unequal distribution of power and income, globally and nationally [8,9].

In 1998, the W HO published *Social Determinants of Health-The Solid Facts* [201], a booklet that presented policy makers, researchers and practitioners with a more holistic approach to views on health that acknowledged the impact of the social environment on health, as represented by social inequalities in health. In 2001, the WHO approved the International Classification of Functioning, Disability and Health (ICF), a framework that synthesizes the medical model of disability with the social model of disability. The ICF conceptual model of disability is innovative in that it recognizes disability as a complex phenomenon that includes the interaction of an individual with features of the social and environmental context [202]. With the creation of the WHO Commission on Social Determinants in 2005, views of disease causation have continued to expand to include discourse on health risks that are created and maintained by the social context above and beyond individual risk factors [10–12].

Early research related to social determinants and health disparities focused on social position and explored downstream determinants primarily related to the individual, such as mechanisms of human biology and the clinical issues of how people cope with disease and disability. Increasingly, research findings have focused on the broader view of upstream determinants related to the community, such as place of residence, work environment or wider social and economic policies of society [10–12].

It is important to focus on social determinants and osteoarthritis (OA) outcomes to advance our understanding of how social context affects health outcomes, particularly in older adults who may experience multiple chronic conditions. OA is one of the most frequently occurring musculoskeletal diseases in western nations, posing a significant public health problem due to its impact on pain, disability and older adults [13–16,203]. Despite the rise in literature on the impact of environment on physical and mental health and functioning of individuals [17–19], the majority of studies have not focused on older adults with OA who may be particularly vulnerable to barriers in their surrounding social and physical environments [20,21]. With the aging of the population worldwide, the prevalence of OA is expected to increase and is likely to result in substantial economic costs for health-care systems. As noted in the recent Institute of Medicine report, *Living Well with Chronic Illness: A Call for Public Health Action*, although the health of Americans is better now than any other time in history, the gains are not shared by all members of our society [22].

Studies on risk factors of OA have focused on variables at the individual level (e.g., BMI, older age and female gender); however, even after controlling for these individual factors, variation in arthritis prevalence is not completely explained. Evidence exists linking lower levels of individual socioeconomic status (SES) and poorer health outcomes in many diseases [23–27], including arthritis [28–31]. Recent data suggest that the social context (i.e., factors in society that cannot be directly measured at the individual level), or socioeconomic environment of an individual's neighborhood may be relevant to arthritis prevalence and health outcomes as well. Although studies have focused on how community contexts interact with individual characteristics for individuals with rheumatoid arthritis (R A), systemic lupus erythematosus and self-reported arthritis [32–38], far fewer studies have investigated this relationship with populations of individuals with OA [22,39,40].

Prior research has demonstrated a strong association between SES and numerous health outcomes and several frameworks currently exist that explain the relationship between social determinants and poor health outcomes [41,204]. The most common dimensions used in research as proxies for SES include educational attainment, occupation and income or

wealth. We recognize that SES is a complex phenomenon and that there are numerous dimensions of SES that are not captured by these factors; however, it is not within the scope

dimensions of SES that are not captured by these factors; however, it is not within the scope of this paper to discuss this at length. While these measures are interrelated and often combined because they reflect aspects underlying socioeconomic stratification, it remains critical to decompose these measures individually to better understand the specific mechanisms and pathways through which these factors impact specific health outcomes at different stages of the life course, in various populations. Whereas occupation is widely used as a proxy indicator for occupational exposure, social standing, or social networks, education can capture knowledge-related assets of an individual [42] that may affect an individual's problem-solving abilities or influence values [43]. Within the life course framework, educational attainment can also be a reflection of one's early life circumstances [44]. Income and wealth are likely to influence health primarily through the transformation of money and assets into health enhancing environments, commodities and services [45], whereby increased income provides access to additional, or higher quality health-promoting resources.

In this narrative review, we apply these general concepts and propose an OA-specific framework to help guide our understanding of how social context and social position may interact with pathophysiological processes and individual-level variables to influence health outcomes in those living with OA (Figure 1). Recognizing that social context has not been extensively researched in OA, our proposed schematic is based on previous models for other chronic diseases and what current research findings in other rheumatologic diseases have demonstrated [22,30,31,38,40,46–49]. We summarize the current literature on research about OA and social determinants including discussion of current measures of social context, social position, and potential mediators and moderators between social determinants and OA outcomes. Our review closes by focusing on future directions and identifying conceptual and methodological limitations in our current knowledge about social determinants and OA, which will contribute to the framework and further elucidate our understanding of the mechanisms of the health–environment relationship.

# OA: definitions & outcomes

## Definition of OA

Studies report different measurements of OA. While some studies may use self-reported OA based on questionnaires, radiographic definitions of OA are more common. Studies that measure self-reported arthritis often use the response to a question such as "Have you ever been told by a doctor or nurse that you have any of the following long-term health conditions?", listing OA as one of the conditions to which respondents are asked to indicate 'yes' or 'no' [38]. Radiographic definitions are often based on the Kellgren-Lawrence (K-L) scale, which assesses radiographic features of OA such as joint space narrowing and osteophyte development [50,51]. Typically, individuals who score 2 or above on the K-L scale are considered as having radiographic knee OA [50]. Symptomatic OA is defined as the combination of radiographic evidence of OA and self-reported pain in the same joint. Although OA can in theory exist in any joint in the body, it most commonly affects the weight bearing joints such as the hips and knees, but can also frequently affect the hands, feet and spine [52]. In this review, all studies have reported radiographic or symptomatic OA unless otherwise noted.

#### OA health outcomes

In addition to measuring the prevalence of OA in the general population and assessing severity and progression of OA through radiographs and joint evaluation, recent research has examined physical and psychosocial health outcomes of individuals living with OA (Figure

1). Arthritis is both a leading cause of disability [13] and a frequent source of chronic pain among older adults [53]. Joint change and destruction can often lead to the need for joint replacement surgery [53]. Depressive symptoms are also common among people living with chronic arthritis pain [54,55] and previous studies show that individuals with OA often report an overall lower health-related quality of life [56].

# Measures of social determinants

There are several dimensions to measuring social position and social context as they relate to populations with OA (Figure 1). Similar concepts are often composite measures that are aggregated differently depending on the population of interest, or unique aspects of the country of research. In addition to measuring social position and social context, our model also includes a number of mediators, moderators and common confounders, through which social position and social context could affect OA health outcomes. While many aspects of social position and social context can be difficult or impossible to change, understanding mediators and moderators in these associations can provide more easily modifiable opportunities for public health interventions to improve health outcomes.

#### Social position, compositional effects & individual level measures of SES

Compositional attributes of SES are measured at the individual level and have commonly included variables such as occupation type (professional/managerial versus nonprofessional/ nonmanagerial) or occupational grade ranking in Europe, level of educational attainment, income (both individual and household), home ownership, and social class (Figure 1) [57]. Education and occupation tend to be two of the most readily attainable variables. Several studies have examined the relationship between formal education level and rheumatic disease outcomes, particularly in RA [58–64] and to a lesser degree in OA [65,66].

**Educational attainment**—Lower levels of educational attainment have frequently been associated with the increased prevalence, morbidity and mortality of many chronic diseases [67]. Although it is often used as a proxy for SES, in comparison to income and occupation, educational attainment is unlikely to be changed by disease. Educational attainment has been measured as a continuous variable by the number of years of schooling in studies [65,68]. However, as relatively few individuals in developed countries have received no formal education, or only elementary school education [69], studies often dichotomize levels of schooling at <12 and 12 years to represent less than high school or completion of high school and postsecondary education, respectively [22,66,69].

**Occupation**—In the USA, several studies have measured occupation type based on US census classifications, with self-reported job titles grouped into one of seven occupational groups [22,70] (farming, forestry or fishing; management or professional; fabrication or manual labor; precision production, crafting and repair; service; technology, sales or administration; and military) [205]. These occupational groups are often then classified as nonmanagerial or managerial. The term managerial is intended to imply occupations that tend to include office work, lower physical demand, more control over the pace of one's work and higher SES, based on census group descriptions.

In other studies measuring occupational classification with arthritis, workers have also been grouped according to the 2000 Standard Occupational Codes in four major occupational groups: white-collar, service, farm and blue-collar [71,72]. Similarly, in Britain, social class quintiles have been used in OA research [73] and are linked to operational categories of the National Statistics Socio-Economic Classification (NS-SEC) including: I: higher managerial, administrative and professional occupations; II: managerial and technical occupations; IIIN: skilled occupation-nonmanual; IIIM: skilled occupations-manual; IV:

partly skilled occupations; and V: unskilled occupations [74]. Low social class is defined by the residents in households with an economically active head of household in social class IV or V, as a proportion of all residents in households [75]. In a study of prevalence of self-report arthritis in Quebec, researchers measured occupation using the Pineo Porter-McRoberts [76] socioeconomic classification of occupation scale with a total of 16 occupational categories distinguished by their shared attributes in the labor force. Researchers trichotomized these categories into skilled, and semi-skilled or lower scores [77].

**Income**—Far fewer studies have included income as a measure of SES; Hawker and colleagues created a dichotomized variable of annual household income of \$20,000 (Canadian) versus >\$20,000 and also created a composite education/income variable to explore the confluence of low educational attainment and low income [69]. The differences in health outcomes by these compositional measures could reflect relatively direct health benefits of having more economic resources (e.g., better nutrition or housing) or unmeasured behavioral psychology variables [78]. The majority of studies examining social position and OA outcomes have focused on an individual's current SES. The income of one's parents in addition to an individual's own income in adulthood can influence health outcomes (Figure 1) [79,80]. To our knowledge, no studies have examined associations of childhood SES, such as parental education, occupation, income or homeownership with the prevalence of OA or OA health outcomes.

#### Contextual effects & multilevel modeling: individual & community level variables

Social context can be defined as the "social and cultural forces that shape people's day-today experiences and that directly and indirectly affect health and behavior" [81]. Aspects of one's environment can be either health-promoting or health-damaging [57]. The socioeconomic context of communities may affect characteristics of the environment of communities to which all residents are exposed, regardless of their social position. The complex nature of social context can be described in terms of both physical and social components, as well as their objective (i.e., actual) or subjective (i.e., perceived qualities) and their scale or immediacy to individuals and groups [82].

**Community SES**—In an Australian study examining the variation in rates of hip and knee joint replacement, SES was derived from the Index of Disadvantage [83]. This index is an area-based measure that represents the average level of disadvantage across a geographical area (i.e., Statistical Local Area) [84] and is a composite of the income, educational attainment, levels of public sector housing, unemployment and jobs in relatively less skilled occupations. These areas are classified into quintiles where 1 indicates an area with significantly disadvantaged households and quintile 5 represents the least disadvantaged households. Another Australian study, examining the association between SES and utilization of primary total hip replacements used postcodes and residential addresses of each subject and matched them to corresponding ABS Census Collection Districts (i.e., 250 households.) [85]. These reference data were then used to determine the Socio Economic Indexes for Areas (SEIFA) value from the census and then the Index of Relative Socioeconomic Disadvantage (IRSD) was applied.

The Townsend Deprivation Score (TDS) is a similar measure of deprivation, which uses census data to enumerate district (i.e., 450 people in 200 households) or ward (i.e., 5500 people in 2400 households) levels and provide a composite score based upon [86]:

- The percentage of homes that are not owner-occupied;
- The percentage of economically active residents who are unemployed;

- The percentage of households that do not have access to a car;
- The percentage of households with more than one person per room.

In the USA, studies have used census data to calculate the household poverty rate at the block-group level (i.e., 600–3000 people) [22,40,87], which is the percentage of households with income below the poverty level, as defined by US Census Bureau poverty thresholds. Participants' physical addresses are geocoded, linked to a block-group identification number, and then used to extract aggregate census information. Poverty level variables can then be categorized, for example, into tertiles: low, medium and high. In studies of self-reported arthritis in Canada, provincial and regional level measures, such as families in a region with low income have been used [46].

**Built environment**—Previous studies that have examined features of the neighborhood environment have used modified versions of the Home and Community Environment Instrument (HACE) [88], a self-report measure that examines how different environmental domains influence levels of participation, an instrument that has both construct validity and reliability [88]. In studies of individuals with OA, features of the neighborhood environment have examined community mobility barriers (i.e., uneven sidewalks or other walking areas; parks and walking areas that are easy to get to and easy to use; safe parks or walking areas; places to sit and rest at bus stops, in parks, or in other places where people walk; curbs with curb cuts) and transportation facilitators (i.e., public transportation close to home; public transportation with adaptations for people who are limited in their daily activities; and adequate handicap parking, able to drive, have a car available to you at your home) [89,90]. Scores are summed and dichotomized into low barriers (0–1 barrier) and high barriers (2–5 barriers) and high facilitators (4–5 facilitators) and low facilitators (0–3 facilitators), respectively.

While many studies have seen moderate associations between certain occupational activities (e.g., kneeling, squatting and climbing) and the increased risk of knee OA as a result of increased biomechanical loads on the knee joint [91–93], far fewer studies have examined associations of OA with organizational characteristics and policies in the work environment. In one such study, Chen and colleagues asked study participants about work accommodations (i.e., opportunities to switch to physically less demanding jobs; availability for part-time work for people needing reduced time), and benefits (i.e., paid sick leave, short-term or long-term disability payment) [70].

**Environmental exposures**—Regional differences in OA prevalence may be an indication of physical environmental factors contributing to health outcomes. Early studies suggest that whole blood lead levels are associated with radiographic knee OA [94].

#### Life course

The life course approach is particularly relevant in understanding the long-term effects of both social position and social context on chronic diseases. Life course measures encompass the aforementioned factors while explicitly including the variable of time, which allows researchers to study disease outcomes across generations and at the individual level beginning from gestation and through childhood, adolescence, young adulthood, middle-age and senescence [95]. Several proposed theoretical models that explain the influence of life course on disease risk may include critical or sensitive periods, which state that exposures during a specific window of time may have long-lasting effects that increase disease risk [96].

While life course is difficult to assess in practice, with most studies measuring life course SES at only two or three time points, there is suggestion that the impact of childhood SES on health decreases over time in adults [97]. Other life course models propose that effects accumulate over the life course, such that increased duration or number of exposures are compounded over time, leading to increased health risk in later years [10]. Although methodological challenges exist in creating summary measures, combinations of the aforementioned indicators (e.g., educational attainment, environmental exposures, occupation type among others) in OA studies would allow researchers to better understand the trajectory and development of OA for an individual over time.

# Potential confounders, mediators & moderators between social determinants & OA outcomes

Our model also includes commonly related personal risk factors associated with OA (Figure 1), including smoking status [98], older age [99], female gender [100], occupation-tasks [101–103], BMI (i.e., obesity) [104,105], injury [100], genetics [106,107], race [105] and physical activity [108]. Many of these personal risk factors are also associated with one's current social position [109]. Although race is not considered a social determinant of health, we have included it in the personal factors in our framework and discussion as it is highly correlated with social position, and a large body of research in OA has explored race as an effect modifier [110–113]. In addition to the more commonly associated risk factors for OA, our figure also includes more proximal mediator and moderators, such as community perceptions and several dimensions of psychological influences (i.e., perceived helplessness, social support/coping resources, psychological disposition, perceived discrimination and/or catastrophizing), that previous research in other rheumatologic conditions or emerging literature in OA have demonstrated to be associated with health outcomes. Including these sets of factors provides for a more transparent relationship to be made linking social determinants to health outcomes, but also allows us to identify areas for public health or behavioral interventions.

While we acknowledge that it is important to recognize individual risk factors associated with OA prevalence and health outcomes, the WHO ICF model also recognizes that attributes beyond an individual affect outcomes. Thus our model of OA health outcomes includes both external, environmental factors, which we refer to as the social context, and internal, personal factors, which are included in our measures of social position and mediators and moderators (Figure 1). This integrated biopsychosocial approach allows us to understand the mechanisms through which distal factors can affect more modifiable, proximal determinants of OA health outcomes.

# Studies of social determinants & OA outcomes

#### Social position & outcomes in knee & hip OA

Significant associations between radiographic and symptomatic knee OA and low educational attainment in the National Health and Nutrition Examination Survey has been reported by Hannan and colleagues [65]. Symptomatic knee pain and self-reported OA remained significantly associated with educational attainment (Table 1), even after adjustment for the risk factors of age, race, BMI, knee injury and occupation. The association between educational attainment and radiographic OA no longer remained significant among women after adjustment for risk factors. A later study of the Johnston County Osteoarthritis Project [66], revealed gender differences as well, such that for men, associations of low educational attainment with radiographic NA and low educational attainment persisted after adjusting for smoking status, alcohol use, occupation, knee injury

and obesity. Similar to Hannan and colleagues, this study found that symptomatic knee OA was more prevalent in both men and women with limited educational attainment. Significant associations between education and OA of the hip were also noted in the Johnston County Osteoarthritis Study [114].

Few studies have examined associations between educational attainment and other OA outcomes, such as disability and pain, in hip OA. In a Finnish study conducted in a clinical setting, educational level was identified as a significant factor for self-reported pain and physical function in participants with hip OA [68]. The EUROHIP consortium [115] reported that greater severity of pain, stiffness and physical function symptoms, as measured by the Western Ontario and McMaster Universities (WOMAC) Arthritis Index, was associated with lower levels of educational attainment in a population of patients with hip radiographic OA who were seeking hip replacements. In a study conducted with patients of the Dresden Hip Surgery Registry [116], significant differences in mean gain in WOMAC scores indicated better scores with increasing levels of educational attainment when comparing those who finished school after 12 years to those who ended their education after 8 years; however, beyond 12 years of educational attainment, no significant differences in WOMAC scores were seen. Similar findings were seen in a community-based sample, whereby WOMAC scores were significantly associated with low educational attainment and nonmanagerial occupation in radiographic and symptomatic hip OA, and disability was significantly associated with low educational attainment [22]. Although associations remained even when all SES variables (e.g., educational attainment, occupation and community poverty) were in the model, they were significantly diminished.

A community-based prospective observational study of hip and knee OA conducted in the UK confirmed that individuals in lower social classes were more likely to experience greater deterioration with more severe knee and hip pain and disability at follow-up, while another study conducted in Canada found that individuals with lower levels of educational attainment or lower income were more likely to have potential unmet need for arthroplasty [69].

International studies examining SES and total joint replacement (TJR) have been mixed, but some have demonstrated an association. TJR is a cost-effective elective procedure that is used in severe cases of OA [117]. SES, particularly less education and lower income have been shown to be independently associated with a greater likelihood of having the potential need for TJR [69]. However, those with low SES are less likely to receive a consultation for their OA or TJR [118,119]. Despite having more severe arthritis [69], those with lower SES are often recommended for TJR less frequently [120]. It has been suggested that this may be due to physicians' perceptions of patients with low SES, which has been shown to be associated with beliefs about abilities, behavioral tendencies and role demands [121]. Further affecting the rates of TJR among those with lower SES is their willingness to consider receiving the procedure. Lower SES, particularly less education, may affect perceived risks of those with lower SES [122] and subsequently influence willingness to receive TJR [123]. Thus, rates of receiving TJR have been consistently associated with SES worldwide, with low rates of TJR amongst those with lower SES in Australia [85], Europe [118,124,125], Canada [119,126] and the USA [127-129]. Furthermore, SES has been associated with higher risk of acute adverse medical events and worse outcomes [124,130,131]. Although SES may play some role in joint replacement, research has shown that the procedure is a complex relationship that involves patient-related factors and issues surrounding equitable access to healthcare service [127].

#### Social context & outcomes in knee & hip OA

Community SES-To our knowledge, only one study has examined associations of individual level SES and community level SES on the prevalence of radiographic and symptomatic knee OA [40]. This study confirmed previous findings in rheumatology research that neighborhoods have an independent effect on outcomes in arthritis [30,31,46]; independent associations were noted between neighborhoods with higher rates of household poverty and outcomes of knee radiographic OA, bilateral knee radiographic OA and symptomatic knee OA in the Johnston County Osteoarthritis Project (Table 2) [40]. A recent multilevel Australian study using self-report measures of arthritis also found independent associations of neighborhood disadvantage with prevalence of arthritis such that residents of the most disadvantaged neighborhoods were 42% more likely than those in the least disadvantaged neighborhoods to self-report arthritis [38]. Although individual characteristics, such as BMI, educational attainment, and occupation type, have been linked to the risk and prevalence of OA, they do not explain all the variance seen across different geographic regions. Thus, it will be of continued importance to understand how community characteristics (e.g., access to physical activity resources, environmental exposures), beyond poverty, may be linked to the prevalence of disease.

The relationship between community-level SES and several OA outcomes has also been examined in the Johnston County Osteoarthritis Project. One study investigating SES factors associated with hip OA found that living in a community with high household poverty rate (>25%) was associated with radiographic hip OA [114]. In another investigation, although Knight and colleagues did not find any association between community poverty and disability among those with radiographic hip OA [22] after adjusting for education and occupation, they did find that individuals with radiographic OA of the knee who lived in areas with poverty rates higher than 25% had higher total WOMAC scores when compared with those living in areas with poverty rates lower than 12%. This association remained strong when other sociodemographic factors were considered [132].

In addition to studying associations with prevalence and physical health outcomes, several studies suggest that social context may contribute to the variations in the utilization of knee and hip replacements (Table 2). Although a 2007 study by Cookson and colleagues in England found that socioeconomic small area inequality in total hip replacement decreased from 1991 and 2001 [133], a recent Australian study reported by Dixon and colleagues noted significant variations in rates of hip and knee joint replacements by community SES [83]. People living in the most disadvantaged and remote areas had more knee, but less hip replacements compared with those who lived in the least advantaged areas. Researchers posit that this variation in knee as compared with hip replacement procedures may result from a greater proportion of individuals from disadvantaged areas being engaged in occupations that involve kneeling, heavy lifting and combined kneeling/squatting and lifting, all of which are known risk factors for knee OA [134].

**Built environment**—While census and administrative data can be used to capture objective aspects of communities (e.g., poverty rates), descriptive, qualitative data describing an older adult's community environment (e.g., safe walking areas and even sidewalks) may provide more insight on how neighborhood features may promote mobility or allow adults to remain engaged in daily activities. Older adults often report mobility barriers resulting from features in their community environment [135], however, fewer studies have examined the association of the community environment with disability (Table 2) [136]. Understanding qualitative aspects of the built environment as health enhancing or as perceived barriers to older adults provides information to policy makers that can allow them to more effectively allocate resources at the community level to promote quality of

life, enhance mobility and reduce disability in the lives of older adults, particularly those living with OA. A study conducted with a knee pain and disability subcohort of the Multicenter Osteoarthritis Study revealed that walking areas, handicapped parking and public transportation were important features of the neighborhood that affected health outcomes of older adults with functional limitations such that the absence of parks was associated with more reports of disability with social and recreational activities and the presence of handicapped parking was associated with less disability for social, leisure and work role activities [90].

In addition to considerations of neighborhood-level features that may affect health outcomes of those living with OA, the prevalence of knee OA has been examined in conjunction with workplace policies. It was found that individuals employed in workplaces offering better policies (e.g., work accommodations and benefits such as paid sick leave and long-term disability payment) had significantly less knee symptoms compared with those with fewer workplace policies, taking into account age, race, sex, SES, BMI, smoking habit, prior knee injury, frequencies of occupational physical activities, or occupational group [70].

Access to affordable medical services and interventions and access to quality healthcare can be an important determinant of health outcomes in OA [137]. Previous research has shown that among individuals with hip and knee OA, those with lower levels of educational attainment were less likely to receive advice on exercise [138]. A systematic review of evidence on the effectiveness of interventions to improve healthcare quality for disadvantaged populations with OA noted that while several arthritis self-management interventions target disadvantaged populations with OA, there is a dearth of literature on how well these programs work compared with more advantaged population to reduce disparities in care [139].

**Environmental exposures**—There is strong, consistent evidence in the USA that shows that there is a disproportionate burden of risk to suboptimal unhealthy environmental conditions for those with low SES [140]. Early studies on environmental exposure and OA have suggested that increasing levels of whole blood lead exposure is associated with small but significant increases in the prevalence and severity of radiographic knee OA [94]. It is currently unclear whether or not this association is explained by a direct toxic effect of lead on joint tissues or through an indirect mechanism, however this preliminary research suggests that features of the environment, such as environmental exposure to this toxicant may play some role in mobility and disability in knee OA.

#### Life course

Although there are no current studies that have examined composition and context over the life course, there is evidence to suggest that BMI, an important risk factor for OA, may have unique effects across the life course. Researchers in the UK examined the relationship between lifetime BMI and midlife knee OA in a population-based British cohort of 3035 men and women with knee OA [141], comparing outcomes of prolonged exposure to high BMI to high BMI at a particular period. Wills and colleagues found that prolonged exposure to high BMI throughout adulthood had a greater influence on the risk of knee OA [141]. Analyses supported an accumulation model of risk indicating that having a high BMI in earlier stages of life was not independently associated with knee OA above and beyond BMI in adulthood. Earlier findings by Macfarlane and colleagues on the association between BMI throughout the life course and adult knee pain found that while association with adult obesity is strongest, associations of high BMI at the age of 11 years was associated with increased risk of knee pain at the age of 45 years [142].

Recent research on life course SES and RA found that patients with R A reported lower childhood SES compared with noncases, with cumulative effects across multiple measures, such as household education, food insecurity and young maternal age [143]. Another study conducted in southern Sweden with patients in an outpatient clinic and private rheumatology clinics assessed perinatal characteristics in relation to the risk of adult RA. High birth weight, initiation of breast feeding during inpatient care after delivery, and paternal occupation were associated with OA indicating that characteristics of the perinatal period may be relevant to the pathogenesis of RA [144]. These findings on the impact of life course BMI and OA and previous research on life course SES and RA suggest that it will also be important to examine social position and social context across the life course in OA health outcomes.

# Potential mediators & moderators

Social position and social context most likely affect health outcomes indirectly through other factors such as personal lifestyle, demographic characteristics as well as community perceptions and psychological influences (Figure 1). As social position and aspects of the community are more difficult to intervene upon or impossible to change in adulthood, it is critical to identify and better understand modifiable risk factors, which may be more amenable to intervention in order to eliminate disparities in the prevalence of OA and improve health outcomes of those with OA. As we have previously noted common risk factors associated with OA prevalence, this section will highlight other potential mediators and moderators (Table 3).

#### Personal factors

**Occupational tasks**—Specific occupational tasks related to radiographic OA and symptomatic OA have not been consistent across studies [145,146]. However, research has consistently shown a strong, positive relationship between work involving knee bending and knee OA in men [147]. Specific occupations however, such as farming, have shown moderate associations with radiographic and symptomatic OA [148], particularly for those who have ten or more years in agricultural work [149]. Far fewer studies have examined the associations of specific occupational tasks with OA of the hip. However, research suggests that prolonged standing at work and heavy lifting increase the risk of degenerative hip disease for farmers [149]. Recent research by Allen and colleagues demonstrated an association between occupational tasks and knee OA (Table 3) [102]. Four specific tasks were associated with symptomatic hip OA: lifting >10 lbs, bending/twisting/reaching, crawling, doing heavy work while standing, as well as the cumulative number of tasks. Additionally, there was an association of lifetime exposure to jobs involving specific occupational tasks with radiographic knee OA, but not radiographic hip OA. However, a Danish study using register data of the whole Danish working population confirmed that occupations with heavy physical workload were associated with a high risk for both hip and knee OA in men and women separately, with risks increasing with cumulative years in occupation (Table 3) [150].

**Health literacy**—Although health literacy has not yet been explored as a mediator or effect modifier in OA health outcomes, it has been associated with poorer health outcomes in other common chronic diseases [151], such as diabetes [152] and RA [153], as well as in the functional health status among older adults [154]. Swearingen and colleagues analyzed the literacy of 194 consecutive patients at an academic rheumatology clinic using the Rapid Estimate of Adult Literacy in Medicine (REALM) and an 'Arthritis-Adapted' REALM (A-REALM) [155]. This study found that more than 10% of patients living with R A for an average of 11 years could not read 'cartilage', 'diagnosis', 'rheumatologist', or 'symptom',

highlighting the potential for literacy-associated barriers in socioeconomic disparities in rheumatologic health.

#### **Community perceptions**

Understanding qualitative aspects of the built environment as health enhancing or as perceived barriers to older adults provides information to policymakers that can allow them to more effectively allocate resources at the community level to promote the quality of life, enhance mobility and reduce disability in the lives of older adults, particularly those living with OA.

Despite the vast body of research that links features of the built environment to physical activity [156–159], research on environmental features as they relate to disabled individuals living with OA, is relatively recent. Keysor and colleagues confirmed the hypothesis that people who reported more mobility barriers and fewer facilitators in the community perceived more limitation in their daily activities [89]. A recent qualitative study with six focus groups in Johnston County (NC, USA), identified community resources and environmental characteristics that are facilitators or barriers to OA disease management [39]. Using the Corbin and Strauss chronic illness trajectory framework, the study divided emergent community resources into three self-management tasks: medical and behavioral management, role management and emotional management. Local area senior and civic centers, a local health facility and aquatics center, community parks and religious organizations were viewed as community resources that promoted physical activity. Social support networks were also used for self-management and to facilitate the completion of daily tasks. The positive benefits of socializing contributed to emotional management of OA. Focus group participants also discussed environmental features in their neighborhood as barriers and facilitators for OA management. Although short distances between residences and community resources promoted physical activity, the availability and quality of lighting, sidewalks and level surfaces could limit one's ability to engage in physical activity [39]. However, although Martin and colleagues' research revealed older adults' recognition of existing neighborhood barriers, Keysor and colleagues found no association with the way people perceived how often daily activities were performed; while features in the neighborhood may make involvement in daily activities difficult, older adults with functional limitations are able to adapt (Table 3) [89].

#### **Psychological influences**

**Perceived helplessness**—Previous research in other rheumatologic conditions has demonstrated an association with helplessness and outcomes [59,160,161]. Early research by Callahan and colleagues confirmed that helplessness mediated a component of the association between formal education level and mortality over 5 years in patients with RA [59]. A recent investigation of patients in the Norfolk Arthritis Registry examined the cross-sectional relationship between SES, learned helplessness, and disease outcome in patients with inflammatory polyarthritis [160]. Camacho and colleagues found that patients with the lowest SES had significantly worse health outcomes compared with patients with the highest SES, a relationship that appeared to be mediated by learned helplessness [160].

**Social support/coping resources**—An extensive body of research has demonstrated the positive health benefits of social support acting as an appropriate buffer against psychological distress caused by disease [162], particularly for older adult populations [163,164] and in other chronic diseases [165]. In OA, social support has been associated with higher physical functioning, general health, mental health, social functioning and vitality [166–168]. In combination with hardiness and pain, social support accounted for 53

and 54% of the variance in psychological status and physical function, respectively, in older adults with OA [166].

Additional psychosocial factors may explain the variation in self-reports of and disability of those who have OA. Catastrophizing has been conceptualized as both a coping response [169] and a cognitive process in which individuals perceive catastrophic threat, harm or consequences from a stressor (e.g., pain or disability), including negative self-statements and negative predictions about the future. In combination with learned helplessness, Keefe and colleagues demonstrated that levels of catastrophizing were inversely associated with reports of knee pain, disability and psychological distress [170], extending previous research on pain, coping and self-reported disability in older adults [171]. Recent research also found that gender differences in reporting of pain, pain behavior and physical disability were mediated by catastrophizing, even after controlling for depression [172]. A separate study showed that at the end of a 10-week skills intervention focused on providing pain coping skills among older OA patients, the skills training group reported significantly lower levels of pain and psychological disability than the control group [173]. Even at a 6-month follow-up, low catastrophizing scores were associated with better health outcomes.

**Perceived discrimination**—There is little research on perceived discrimination as it affects OA outcomes. In a recent study by Hausman and associates that examined the impact of perceived discrimination on patient-provider communication, African–Americans reported greater levels of perceived racism and classism [174]. Although Hausman *et al.* studied patient–provider communication as the outcome, there is reason to believe that the mechanisms of discrimination operate through complex pathways that could ultimately affect both the prevalence of OA in populations, and other OA health outcomes. Additional research is needed to identify the causal pathway through which perceived discrimination can produce negative mental health and induce physiological stress responses.

# Conclusion

This article reviews the impact of social determinants on OA outcomes and proposes an OAspecific framework that synthesizes a medical model with a social model of disability. Our narrative review highlights several important components of this topic, including discussion on the current methods of measurement of social determinants at both the individual (i.e., occupation type, educational attainment and/or income) and contextual level (i.e., community-level measures of SES, the built environment and/or environmental exposure), while also identifying potential mediators and moderators that may affect OA health outcomes.

Educational attainment is one of the most investigated measures of social determinants. It has been found to be associated with prevalence of knee and hip OA, as well as markers of disease severity and progression among those with knee and hip OA, including disability, pain and physical function. Other dimensions of social position can be captured through occupation type or measures of income, but vary from study to study, and country to country.

Research on contextual measures of social determinants has demonstrated that living in areas with high community poverty rates is associated with higher rates of self-reported arthritis and higher prevalence of radiographic knee and hip OA and increased disability among those with knee OA. More qualitative, descriptive approaches to capturing contextual environment have included studies on features of the built environment, community perceptions and workplace policies that can be viewed as facilitators or barriers to OA self-management.

We recognize that the mediating, moderating and confounding factors included in our schematic are not an exhaustive list. However, in our review of the literature, we created three categories to reflect conceptual distinctions, including personal factors, community perceptions and psychological influences. As the body of research on social determinants in OA continues to expand, and the measurement of concepts becomes more precise, we expect that additional factors will be added to the model and the mechanisms explaining the relationship between social determinants and OA health outcomes to be better explicated.

# Future perspective

In reviewing the existing literature on social determinants of OA, it is important to identify gaps in the literature and areas in need of future research in order to reach out to decision-makers and guide the development of health promoting policies that reduce the disparities seen in OA and other chronic diseases. Our paper has identified several factors of interest and provided rationale as to how they may potentially affect OA health outcomes, most of which have largely been unexplored. These include: social cohesion/social capital, environmental exposure, neighborhood environment (perceived and actual), health literacy, perceived discrimination; and life course. We encourage researchers to consider our proposed schematic in their future studies and develop their research around these factors to help build and strengthen our understanding of the complex causal linkages that underlie disparities in OA health outcomes.

While there is strong evidence for individual SES/compositional effects on prevalence and outcomes of OA, most studies have focused primarily on educational attainment and occupation type. The inclusion of more studies examining the association of income or wealth might provide more understanding on the conversion of assets and money into health-enhancing commodities, environments and services. Measurement of social phenomena and other aggregate constructs in social determinants at both the individual level and particularly at the group level remain underdeveloped [175]. Definitions of occupation type, community poverty and deprivation often vary from study to study and country to country. Thus, we encourage continued conceptualization in the development of valid measures that would allow for the comparison of data across countries and improvement in methodological approaches and study designs that can better capture dimensions of social position and social context. In particular, additional research is needed to determine if aggregate-level measures of SES are providing accurate estimations of individual SES as well as determining if there are specific populations where SES measures should be measured contextually or individually.

More research, particularly multilevel modeling, is needed. Although there is evidence linking social context and individual SES in other chronic conditions, there is little research regarding how community contexts can interact with individual-level characteristics in OA. Using the ICF conceptual model of disability as a framework for research in OA requires adequate consideration of the complex relationship of individuals within their social and environmental context [202]. In particular, greater use of multilevel models in OA and determinants research will better allow us to understand how populations interact with aggregate phenomena (i.e., community poverty) and change over time.

With a growing aging population, and one that lives longer, the life course perspective will allow researchers to understand trends within chronic diseases and understand trajectories of illness over time. Currently, no studies have explored the role of social determinants across the life course and OA.

Although radiographic assessment of OA is considered the gold standard, some studies on OA still use self-report as their measure of disease. A movement towards radiographic,

clinical assessment as compared with self-report is needed to improve accuracy and comparability between studies. Additionally, there has been a large focus on OA of the knee and hip. More research is needed to assess the effects of social determinants on other types of OA, including the hands, spine and feet.

With the exception of the Johnston County Osteoarthritis Study, which has enrolled a sizeable number of African–Americans, most research investigating the effect of social determinants on OA outcomes has focused primarily on Caucasian populations. Future research needs to be carried out to determine whether OA outcomes have similar associations with SES among different racial and ethnic groups/settings, different cultures, developing countries and older adults. In closing, it is of continued importance to focus on social determinants and OA outcomes in order to advance our understanding of how social context affects health outcomes. Identifying potential mediators, such as social support or psychological disposition, which are proximal to OA health outcomes, may provide public health practitioners with more accessible opportunities to develop health interventions to improve the quality of life of those living with OA. However, we emphasize that only through understanding the mechanisms of distal, social determinants can we expect to truly affect the structural inequalities that underlie the disproportionate burden of illness.

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#### **Executive summary**

#### Overview

- Early research related to social determinants and health disparities focused on social position and explored downstream determinants primarily related to the individual, such as mechanisms of human biology and the clinical issues of how people cope with disease and disability. Over the past 15–20 years, our view of disease causation has broadened to include discourse on the health risks that are created and maintained by the social context above and beyond individual risk factors.
- This review proposes a framework to help guide our understanding of how social context may interact with pathophysiological processes and individuallevel variables to influence health outcomes in those living with osteoarthritis (OA). Our schematic is based on previous models for chronic disease and what current research findings in other rheumatologic diseases have demonstrated.
- We summarize the current literature on research on OA and social determinants, focus on future directions for research, and identify conceptual and methodological limitations in our current knowledge about social determinants and OA.

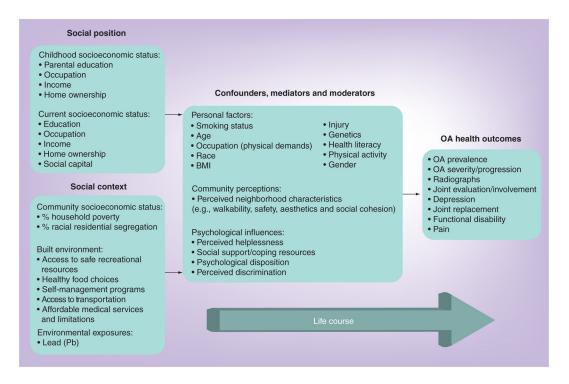
#### Social determinants & OA outcomes

- Education and occupation are the compositional measures of socioeconomic status (SES) that have been examined most extensively in OA.
- Low educational attainment and/or having a nonmanagerial/nonprofessional occupation have been demonstrated to be associated with increased OA prevalence and morbidity (poor health outcomes), after adjusting for other known risk factors (gender, BMI, injury, and age).
- In the limited studies to date, social context (percentage of the neighborhood living in poverty) has been shown to be associated with OA outcomes, independent of individual measures of SES.

#### **Conclusion & future perspective**

- Due to the complex nature of community variables often referred to as social context, better and more consistent definitions of this measure need to be developed.
- Research on the effects of social determinants on OA outcomes has focused primarily on education and occupation type. Definitions of SES need to be expanded to include other measures that will address methodological issues in defining social determinants, including:
  - Social cohesion/social capital
  - Environmental exposure
  - Neighborhood factors (actual and perceived)
  - Health literacy
  - Perceived discrimination
  - Life course
  - Income

• Most research investigating the effect of social determinants on OA outcomes has focused primarily on Caucasian populations. Future research needs to be carried out to determine if OA outcomes have similar associations with SES among different racial and ethnic groups/settings, different cultures, developing countries and older adults.



**Figure 1.** The relationships between social position, social context and osteoarthritis outcomes OA: Osteoarthritis.

| ► : | Type of OA  | Social position or social context   | Data source/population  | Health outcomes   | Covariates<br>measured<br>(personal factors)  | Outcome  | Country | Ref. |
|-----|---|---|---|---|---|--|---------|------|
|     | Knee OA<br>radiographs,<br>symptomatic<br>knee pain, self-<br>report arthritis<br>at any site   | Educational attainment  | National Health and<br>Nutrition Examination<br>Survey  | Prevalence of OA  | Age, knee injury,<br>race, obesity,<br>occupation   | Low educational<br>attainment associated with<br>a high prevalence of knee<br>OA in both men and<br>women, according to all<br>three definitions, known<br>risk factors explain<br>association between<br>educational attainment,<br>adthough risk factor<br>contributions differ by<br>gender | USA     | [65] |
|     | Self-reported<br>hip and knee<br>OA followed by<br>clinical and<br>radiographic<br>examinations | Educational attainment, income  | Mail, telephone survey of persons aged 55 years and older   | Potential need for<br>and willingness to<br>consider hip and<br>knee arthroplasty | Age, sex, region of<br>residence  | Less education and lower<br>income independently<br>associated with greater<br>likelihood of having<br>potential need for<br>arthroplasty, individuals<br>with less education and/or<br>lower income more likely<br>to have potential unmet<br>need for arthroplasty                           | Canada  | [69] |
|     | Self-reported<br>hip and knee<br>OA, followed<br>by clinical<br>examination                     | Social class, employment status<br>(self and partner)                                       | Somerset and Avon Survey<br>of Health, community-based,<br>age-sex stratified survey<br>registered in 40 general<br>practices | Change in severity<br>of pain and<br>disability                                   | Sex, age group,<br>BMI, marital<br>status,<br>comorbidities,<br>healthcare<br>utilization   | Those reporting greater<br>knee or hip pain had more<br>pain and worse function at<br>follower social class,<br>hypertension and higher<br>BMI was associated with<br>greater deterioration  | UK      | [73] |
|     | Hip OA  | Educational level   | Rehabilitation clinic in a<br>Finnish Hospital  | Self-reported pain<br>and self-reported<br>disease-specific<br>physical function  | Age, sex,<br>depression, life<br>satisfaction,<br>sporting activities,<br>radiologic score,<br>BMI,<br>comorbidities,<br>duration of knee<br>pain | Educational level,<br>comorbidities, and BMI<br>significant factors for self-<br>reported pain and physical<br>functioning, no direct<br>relationship with pain and<br>psychological factors<br>detected   | Finland | [68] |
|     | Unilateral and<br>bilateral<br>radiographic<br>and  | Educational attainment: number<br>of years of formal education,<br>dichotomized at 12 years | Johnston County<br>Osteoarthritis Project, (NC,<br>USA), African–American<br>and Caucasian men and                            | Presence of<br>radiographic and<br>symptomatic knee<br>OA                         | Age, race, gender,<br>smoking status,<br>alcohol use, history<br>of previous knee<br>injury, BMI,   | Association of low<br>educational attainment<br>with radiographic knee OA<br>explained by age and race,<br>but association with  | USA     | [66] |

Table 1

Studies measuring social position and osteoarthritis health outcomes.

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| Country Ref.   |  |
|--|--|
| Outcome Co   | symptomatic knee OA<br>remains after adjustment<br>for covariates; with<br>women, association of low<br>educational attainment<br>with all four knee OA<br>outcomes remain after<br>adjustment; association not<br>explained by current HRT<br>use |
| Covariates<br>measured<br>(personal factors)             | occupational<br>characteristics,<br>hormone<br>replacement<br>therapy  |
| Health outcomes  |  |
| Data source/population                                   | women aged 45 years in<br>rural NC (USA)   |
| Social position or social context Data source/population |  |
| Study (year) Type of OA                                  | symptomatic<br>knee OA   |
| Study (year)   |  |

Luong et al.

HRT: Hormone replacement therapy; OA: Osteoarthritis.

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Luong et al.

Table 2

Studies measuring social context and osteoarthritis health outcomes.

| Study (year)                    | Type of OA                              | Social position or social context  | Data source/population   | Health outcomes                             | Covariates<br>measured<br>(personal factors)   | Outcome  | Country   | Ref.  |
|---------------------------------|---|--|--|---|--|--|-----------|-------|
| Cookson <i>et</i><br>al. (2007) | Total hip replacement                   | Townsend deprivation score   | Hospital episode statistics,<br>English National Health<br>Service   | Elective total hip<br>replacement           | Not available  | Socioeconomic<br>small area inequality<br>in use of total hip<br>replacement fell<br>between 1991 and<br>2001  | England   | [133] |
| Dixon <i>et al.</i><br>(2011)   | Hip and knee joint                      | Community socioeconomic status from the Index of Disadvantage, geographic locality, birthplace | National hospital morbidity<br>database  | Primary hip or<br>knee joint<br>replacement | Indigenous status,<br>age, gender  | Rates of joint<br>replacement were<br>significantly lower<br>in people living in<br>the most<br>disadvantaged and<br>remote areas, people<br>born outside<br>Australia, and<br>indigenous peoples,<br>however knee<br>replacement rates<br>were higher in the<br>most advantaged<br>than in the least<br>disadvantaged areas;<br>thip and knee<br>replacements higher<br>in regional areas<br>than in major cities | Australia | [83]  |
| White <i>et al.</i><br>(2010)   | Incident and<br>symptomatic knee OA     | Individual environment features  | Multicenter osteoarthritis<br>study-knee pain and<br>disability subcohort  | Disability in daily<br>activities           | Knee pain,<br>functional<br>limitation,<br>comorbidities, BMI,<br>age, gender,<br>educational<br>attainment, ethnicity<br>and study site | Older adults whose<br>neighborhoods did<br>not have parks and<br>walking areas less<br>frequently engaged<br>in a regular fitness<br>program, and in<br>social activities  | USA       | [06]  |
| Chen <i>et al.</i><br>(2007)    | Knee OA                                 | Work-related organizational  | Johnston County<br>Osteoarthritis Project, (NC,<br>USA) Affrican-American<br>and Caucasian men and<br>women aged 45 years in<br>rural NC (USA) | Prevalence of knee<br>OA                    | Sociodemographic<br>features, lifestyle<br>factors, knee<br>injuries, BMI, other<br>workplace<br>characteristics                         | Individuals<br>employed in<br>workplaces offering<br>better policies had<br>significantly less<br>knee symptoms  | USA       | [70]  |
| Nelson <i>et al.</i><br>(2011)  | Radiographic and<br>symptomatic knee OA | Environmental exposure   | Johnston County<br>Osteoarthritis Project, (NC,<br>USA), African-American<br>and Caucasian men and   | Prevalence and<br>severity of knee<br>OA    | Age, race, sex, BMI,<br>smoking, alcohol<br>drinking   | Increasing levels of<br>blood lead are<br>statistically<br>significantly<br>associated with  | USA       | [94]  |

| Study (year)                            | Type of OA                             | Social position or social context                                | Data source/population  | Health outcomes   | Covariates<br>measured<br>(personal factors)   | Outcome   | Country   | Ref.  |
|---|--|--|---|---|--|---|-----------|-------|
|   |  |  | women aged 45 years in<br>rural NC (USA)  |   |  | radiographic knee<br>OA   |           |       |
| callahan <i>et</i><br><i>al.</i> (2011) | Tibiofemoral knee OA                   | Educational attainment,<br>occupation, community poverty         | Johnston County<br>Osteoarthritis Project, (NC,<br>USA), African-American<br>and Caucasian men and<br>women aged 45 years in<br>rural NC (USA)                        | Presence of<br>radiographic and<br>symptomatic knee<br>OA | Age, gender, race,<br>BMI, current<br>smoking status,<br>knee injury and<br>occupational<br>activity score | Low educational<br>attainment was<br>significantly<br>associated with<br>radiographic OA,<br>bradiographic OA,<br>symptomatic OA;<br>occupation had no<br>significant<br>independent<br>association beyond<br>educational<br>attainment and<br>attainment and | USA       | [40]  |
| (2011)<br>(2011)                        | Radiographic and<br>symptomatic hip OA | Educational attainment,<br>occupation type, community<br>poverty | Johnston County<br>Osteoarthritis Project, (NC,<br>USA), African-American<br>and Caucasian men and<br>women aged 45 years in<br>rural NC (USA)                        | Pain and disability                                       | Age, gender, race,<br>BMI, presence of<br>knee symptoms  | Pain and disability<br>significantly<br>associated with low<br>accational<br>attainment in<br>radiographic and<br>symptomatic OA;<br>disability<br>associated with and<br>nonmangerial<br>occupation  | USA       | [22]  |
| Brennan <i>et al.</i><br>(2012)         | Hip OA                                 | Relative Index of<br>Socioeconomic Disadvantage                  | Australian Orthopaedic<br>Association National Joint<br>Replacement Registry data<br>among residents of the<br>Barwon Statistical Division<br>of Victoria (Australia) | Primary total hip<br>replacement                          | Age, sex   | Rates of total hip<br>replacement are<br>greater for both the<br>most disadvantaged<br>and least<br>disadvantaged<br>groups   | Australia | [85]  |
| Wills <i>et al.</i><br>(2012)           | Knee OA                                | BMI across life  | British bitth cohort of 3035<br>men and women   | Risk of midlife<br>knee OA                                | Occupation type,<br>activity level at the<br>age of 36 years   | Prolonged exposure<br>to high BMI<br>throughout<br>adulthood associated<br>with the highest risk<br>for knee OA with no<br>additional risk<br>conferred from  | UK        | [141] |
|   |  |  |   |   |  |   |           |       |

OA: Osteoarthritis.

Aging health. Author manuscript; available in PMC 2013 June 01.

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| Study (year)                     | Type of OA                             | Independent variables   | Data source/population   | Health outcomes  | Covariates measured  | Outcome  | Country | Ref.  |
|----------------------------------|--|---|--|--|--|--|---------|-------|
| Sandmark <i>et</i><br>al. (2000) | Knee OA                                | Job titles and exposure to<br>physical load in<br>occupational work,<br>housework and leisure-<br>time activities                   | Men and women who had<br>prosthetic surgery due to<br>primary tibiofemoral OA  | Development of<br>knee OA leading to<br>prosthetic surgery | Age, BMI, sports, for<br>women: hormone<br>substitution  | Among men there was an<br>association between<br>lifting at work, squatting<br>or knee bending, kneeling<br>and jumping with knee<br>OA  | Sweden  | [146] |
| (2010)<br>(2010)                 | Knee and hip<br>OA                     | Ten occupational tasks and<br>lifetime exposure to jobs<br>that require spending<br>>50% of their time doing<br>five specific tasks | Johnston County<br>Osteoarthritis Project (NC,<br>USA), African-American and<br>Caucasian men and women<br>aged 45 years in rural NC<br>(USA)  | Presence of knee<br>and hip OA                             | Age, race, gender, BMI,<br>prior knee or hip injury,<br>smoking  | Radiographic hip and<br>knee OA not significantly<br>associated with any<br>occupational tasks;<br>specific activities<br>associated with<br>associated with<br>asymptomatic knee and hip<br>OA; exposure to a greater<br>number of physically<br>demanding occupational<br>tasks at the longest job<br>tasks at the longest job | USA     | [102] |
| Andersen <i>et</i><br>al. (2012) | Knee and hip<br>OA                     | Occupations with heavy<br>workload (i.e., farming,<br>construction, healthcare<br>work)   | National patient register data<br>on the Danish working<br>population from 1981 to 2006<br>followed-up for hip and knee<br>OA during 1996–2006 | Increased risk of<br>OA, particularly for<br>women         | Age, personal income,<br>personal unemployment,<br>previous knee injury  | Male floor layers and<br>bricklayers and male and<br>female healthcare<br>assistants had the highest<br>risk of knee OA; farmers<br>had the highest risk of hip<br>OA; generally, risk<br>increased with cumulative<br>years in the occupation   | Denmark | [150] |
| Keysor <i>et al.</i><br>(2010)   | Incident and<br>symptomatic<br>knee OA | Community mobility<br>barriers, transportation<br>facilitators  | Multicenter Osteoarthritis<br>Study Knee Pain and<br>Disability subcohort  | Disability: DAL<br>and DAF,<br>prevalence of<br>barriers   | Age, gender, race,<br>educational attainment,<br>comorbidity, BMI, knee<br>pain, functional limitation<br>and study site | People who reported more<br>transportation facilitators<br>reported less disability,<br>whereas a greater<br>presence of transportation<br>facilitators was not<br>associated with DAF   | USA     | [89]  |
| (2011)<br>(2011)                 | Hip and knee<br>OA                     | Participant identified<br>community factors   | Johnston County<br>Osteoarthritis Project (NC,<br>USA), African-American and<br>Caucasian men and women<br>aged 45 years in rural NC<br>(USA)  | Facilitators/barriers<br>to OA self-<br>management         | Not applicable   | Individuals use local<br>recreational facilities,<br>senior centers, shopping<br>centers, religious<br>organizations, medical<br>providers, pharmacies,<br>and their social network<br>for OA management; built<br>environmental<br>characteristics both   | USA     | [39]  |

Table 3

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| Type of OA Indepen  | Indepen        | Independent variables  | Data source/population                                  | Health outcomes                                    | Covariates measured  | Outcome   | Country | Ref.  |
|---|----------------|--|---|--|--|---|---------|-------|
|   |                |  |   |  |  | facilitate and hinder use of<br>community resources for<br>OA management  |         |       |
| Knee and hip Social support Patients with consecutive hip<br>OA and knee OA attending an<br>outpatient physical<br>rehabilitation and<br>rheumatology clinic of<br>University Hospital of Liege<br>in Belgium | Social support | Patients with com-<br>and knee OA atter<br>outpatient physics<br>rehabilitation and<br>rheumatology cliri<br>University Hospit<br>in Belgium | secutive hip<br>nding an<br>ul<br>nic of<br>al of Liege | Health-related<br>quality of life                  | Age, sex, BMI, number of<br>comorbid conditions, SES,<br>site of survey completion,<br>severity of OA  | Greater social<br>companionship<br>transactions associated<br>with higher physical<br>functioning, general<br>health, mental health,<br>social functioning and<br>vitality  | Belgium | [166] |
| OA Sociodemographic Older adults<br>characteristics (age, race,<br>education, income),<br>psychosocial factors,<br>health promotion practices<br>and social support, health<br>status and pain                |                | Older adults   |   | Psychological<br>equilibrium,<br>physical function | Not available  | Hardiness, pain and social<br>support were significant<br>contributors to<br>psychological status and<br>physical function  | USA     | [168] |
| Knee OA Pain coping skills Older adults   |                | Older adults   |   | Self-reported<br>disability, walking<br>disability | Demographic variables,<br>BMI, medical<br>comorbidities, history of<br>fractures, vision and<br>hearing impairment   | Pain coping was<br>significantly associated<br>with disability and<br>distance walked. Less<br>catastrophic thinking and<br>prayer was associated<br>with less disability and<br>better physical function   | USA     | [171] |
| Knee OA Gender, catastrophizing Patients with OA  |                | Patients with O  | A   | Pain, pain behavior,<br>physical disability        | Age, ethnic background,<br>income, education level,<br>number of years since OA<br>diagnosis, generalized or<br>localized OA, disability<br>and financial<br>compensation or not | Significant difference in<br>pain, pain behavior, and<br>physical disability in men<br>and women, mediated by<br>catastrophizing  | USA     | [172] |
| OA Past perceived African-American and white<br>discrimination, classism patients treated for OA by<br>orthopedic surgeons in two<br>Veteran Affairs facilities   |                | African-Ameri<br>patients treated<br>orthopedic surg<br>Veteran Affairs  | an and white<br>for OA by<br>eons in two<br>facilities  | Affective tone of patient-provider communication   | Race, age, annual income,<br>highest educational<br>attainment, whether patient<br>had been seen in the<br>orthopedic clinic<br>previously, quality of life                      | Perceived racism and<br>classism were reported by<br>more African–American<br>patients than by white<br>patients. High levels of<br>perceived racism among<br>African–American<br>patients was associated<br>with less positive<br>nonverbal affect | USA     | [174] |
|   |                |  |   |  |  |   |         |       |

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DAF: Daily activity frequency; DAL: Daily activity limitation; OA: Osteoarthritis; SES: Socioeconomic status.