

Original article

Annual consultation incidence of osteoarthritis estimated from population-based health care data in England

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Abstract

Objectives. To estimate the consultation incidence of OA using population-based health care data in England and compare OA incidence figures with those derived in other countries.

Methods. A population-based health care database (Consultations in Primary Care Archive) in England was used to derive the consultation incidence of OA (overall and by joint site) using the maximum available run-in period method. These estimates, and their distribution by age and sex, were compared with those published from population-based health care databases in Canada, the Netherlands and Spain. A novel age-stratified run-in period method was then used to investigate whether the consultation incidence has been increasing over time in younger adults.

Results. The annual consultation incidence of OA (any joint) was 8.6/1000 persons ≥ 15 years of age (95% CI 7.9, 9.3) [6.3 (95% CI 5.5, 7.1) in men and 10.8 (95% CI 9.8, 12.0) in women]. Incidence increased sharply between 45 and 64 years of age, peaking at 75–84 years. The joint-specific incidence was 1.4 (95% CI 1.1, 1.7), 3.5 (95% CI 3.1, 3.9) and 1.3 (95% CI 1.1, 1.6) for hip OA, knee OA and hand OA, respectively. The estimates and their distribution by age and sex were broadly consistent with international estimates. Between 2003 and 2010, incidence in those aged 35–44 years increased from 0.3 to 2.0/1000 persons.

Conclusion. Newly diagnosed cases of OA in England occur in 9 in 1000 at-risk adults each year, similar to other international estimates. Although lower, the consultation incidence proportion in younger adults appears to have increased in the past decade.

Key words: osteoarthritis, incidence, population-based health care data, consultation.

Rheumatology key messages

- Approximately 9 in 1000 persons aged ≥ 15 years is newly diagnosed with OA each year in England.
- An increase in England in newly diagnosed OA observed between 2003 and 2010 in 35- to 44-year-olds warrants further investigation.

Introduction

OA accounts for 3% of all years lived with disability in the high-income countries [1]. An ageing population, increasing prevalence of risk factors such as obesity and increasing strain on health budgets, including costs

of increasing numbers of patients undergoing joint arthroplasty [2, 3], suggest a growing challenge for population health, the National Health Service and the UK economy [4].

A key recommendation of the UK Chief Medical Officer is the need for better information in musculoskeletal disorders [5]. Alongside the National Joint Registry (<http://www.njrcentre.org.uk/>) and bespoke surveys and cohorts, population-based health care databases based on anonymized, routinely collected clinical information recorded in the electronic health record are important sources of such information, whose advantages in terms

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of representativeness, cost, size and continuity over time are well recognized.

Data from general practice—the setting where most cases of OA are assessed and managed—has been used to estimate the consultation prevalence of OA (the proportion of the population with a diagnosed consultation over a defined period of time) in several countries. Fewer studies [6–10], and indeed none from the UK, have estimated consultation incidence (the rate of new cases presenting to general practice). Yet incidence rates are an important measure of occurrence in the population, responding more quickly to changes in risk factors and being less influenced by disease duration [11]. Previous estimates from Canadian provincial health care databases would suggest an annual consultation incidence for OA in men and women of 9.3–12.2/1000 and 11.0–17.4/1000, respectively [7, 9, 10]. Recent estimates from a large regional primary care database in Spain found a 3-fold higher incidence of knee OA than hip or hand OA (6.5 vs 2.4 and 2.1/1000 persons >45 years of age) [8]. However, direct international comparisons are limited by different OA case definitions and population strata. In addition, these previous estimates cannot be assumed to be representative of current rates in the UK since consultation incidence proportions for OA may be influenced by cohort and period effects, some of which may be particular to location (e.g. the effect of the UK Quality and Outcomes Framework on coding of OA since 2004).

One of the methodological challenges in estimating incidence is how to ensure prevalent cases (those with previous OA consultations) are excluded from the calculation. This is achieved in many databases by using a run-in period (disease-free observation period [12], look-back period [13] or clearance period [14]), which for OA can be at least 9 years [15] (e.g. one needs to look back to 2001 to be confident that a patient consulting with OA in 2010 is receiving that diagnosis for the first time). One disadvantage of this is that databases must be established with satisfactory data quality for many years before trends in incidence can begin to be described [16]. However, we propose that prevalent cases can be effectively excluded using shorter run-in periods in younger age groups since they are less likely to have a long prior history of OA consultations. Therefore our study had two main objectives: first, to estimate the consultation incidence of OA (overall and separately for hip, knee and hand) by applying a maximum available (10 year) run-in period to data from a local primary care database in North Staffordshire, England and compare OA incidence figures with those derived in other countries; second, in the same database, to develop a novel age-stratified run-in period method and demonstrate its application by testing the hypothesis that there has been an increase in the consultation incidence proportion of diagnosed OA among younger adults (ages 35–44 years; i.e. the 10 year age band below the typical age stated in guidelines for clinical diagnosis of OA [17]). Although the incidence of OA in this age group is lower than in older age groups, increasing

incidence of OA over time in this age group could be of particular concern in the context of increasing childhood obesity.

Methods

Data source

The Consultations in Primary Care Archive (CiPCA), which contains all recorded consultation data by general practitioners (GPs) and practice nurses from 11 general practices in North Staffordshire, England between 2000 and 2010 (total practice population consisted of 94 955 people in 2010) [16]. North Staffordshire is more deprived than England as a whole, although the CiPCA practices cover both deprived and less deprived areas. In England, general practice provides the first point of access to the National Health Service for most non-emergency care and also provides continuing care for many chronic diseases. The vast majority of the population are registered at a general practice [18, 19]. Within CiPCA practices, 97% of contacts with a GP are assigned a morbidity code and practices undergo an annual cycle of assessment, feedback and training in morbidity coding [20]. A similar annual primary care consultation prevalence figure for OA from CiPCA practices has been shown compared with those derived from national UK [21] and international (Swedish [16]) databases. Moreover, in CiPCA, secondary care information (e.g. hospital letters) is obtained and coded at the discretion of the practices, thus completeness varies by practice.

Ethical approval for the CiPCA database was given by the North Staffordshire Research Ethics Committee to download, store and analyse anonymized medical records information for research use from participating general practices in CiPCA (reference 03/04). Patients are informed by a poster at their GP's practice and by leaflet that the practice is a Keele research practice and that their anonymized records (with identifiable information removed) may be used for research, and that they can opt out if they wish by informing the practice staff. Therefore no separate ethical approval was required for our study.

Case definition

An OA consultation was defined as a Read code starting with N05 (OA and allied disorders), equivalent to ICD9 codes beginning with 715. Knee, hip and hand OA were each defined by Read code lists drawn up through a consensus process involving local GPs (code lists are available upon request from the authors [22]). A single medical contact with an OA diagnosis is a standard definition used in most previous studies [6, 23] and is preferred in previous validation studies to case definitions based on multiple contacts, referrals and prescription records [24, 25]. Prieto-Alhambra *et al.* [8] report that only 1.3% of cases defined in this way are subsequently given an alternative diagnosis.

Estimating consultation incidence using maximum available run-in period

The overall annual consultation incidence of OA and incidence rates by age (defined as age on 31 December 2010) and sex were estimated for the calendar year 1 January 2010 to 31 December 2010. The numerator for the incidence calculation was defined as those with a recorded OA code in 2010 with no prior OA code and complete registration in the previous 10 years. Hence prevalent cases who were diagnosed with OA during the 10 year run-in period from 1 January 2000–31 December 2009 were not eligible to be included in the numerator. Each patient continuously registered in 2010 and for the run-in period (1 January 2000–31 December 2009) was included in the denominator, with exclusion of those who were prevalent cases in the run-in period. The incidence proportions were then corrected for the loss of patients with <10 years registration (supplementary Tables S1–S24, available at *Rheumatology* Online). A previous study of consultation incidence of OA using health administrative data in British Columbia, Canada concluded that a run-in period of 9 years [10] was generally sufficient to remove prevalent cases. We verified that the 10 year run-in period (the maximum available within CiPCA) was sufficient to observe a stable estimate of consultation incidence in CiPCA by rerunning the analyses varying the run-in period from 0 to 9 years (see supplementary Fig. S1, available at *Rheumatology* Online).

Using the maximum run-in period method, the consultation incidence of OA was estimated for the total population, among people ≥ 15 years of age and among people ≥ 45 years of age, as was consultation incidence of OA by gender and specific age. All analyses were repeated for OA in three selected body regions. We then compared the consultation incidence of OA in women with that in men using overall (age-adjusted) and age-specific female:male incidence rate ratios (IRRs) with 95% CIs using Poisson regression.

Comparison with international estimates

An earlier rapid literature review by us identified six original English-language articles that reported potentially comparable OA incidence estimates from 10 databases in Canada, the Netherlands and Spain [6, 8–10, 26] (supplementary Table S25, available at *Rheumatology* Online). Using tables and plots we compared these international estimates with those obtained in the present study using the maximum available run-in period method in CiPCA. To facilitate the comparisons, direct standardization was used, in which the published age-stratified incidence rates from other international studies were applied to the age-sex distribution of the CiPCA population.

Trend in consultation incidence of OA in younger adults

A novel age-stratified run-in period method (supplementary data, the time series regression model for deriving age-stratified run-in periods section, and supplementary

Fig. S2, available at *Rheumatology* Online) was used to investigate the recent trend in annual consultation incidence of OA among adults aged 35–44 years. Since this age group had the lowest minimum run-in period of all age groups, more data points were available to evaluate trends. The annual incidences in 2003–10 were estimated and the trend was observed by fitting a log-linear model with Wald test to test for linear trend. To increase the stability of the rates with a minimum loss of information, 3 year moving average incidence proportions for 2005–10 were estimated by using the mean annual number of incident cases and mean denominator population over the 3 years. Statistical analyses were carried out using Stata 12.0 (StataCorp, College Station, TX, USA).

Results

On 1 January 2010, 94 955 people (48 237 women, 46 718 men) including 1953 people with a recorded OA diagnosis (1251 women, 702 men) were registered in CiPCA.

Annual consultation incidence of OA

Based on the maximum available run-in period method, the annual consultation incidence of OA at any joint for persons aged ≥ 15 years was 8.6/1000 persons (95% CI 7.9, 9.3) [6.3 (95% CI 5.5, 7.1) for men and 10.8 (95% CI 9.8, 12.0) for women]. In those aged ≥ 45 years, the corresponding estimates were 16.1 (95% CI 14.8, 17.5), 12.0 (95% CI 10.5, 13.7) and 20.1 (95% CI 18.1, 22.2), respectively (Table 1). The consultation incidence proportions, expressed per 1000 persons aged ≥ 15 years, were 3.5 (95% CI 3.1, 3.9) for knee OA, 1.4 (95% CI 1.1, 1.7) for hip OA and 1.3 (95% CI 1.1, 1.6) for hand OA.

Age- and gender-specific consultation incidence estimates are shown in Fig. 1 (see also supplementary Table S26, available at *Rheumatology* Online). Similar patterns of age-specific consultation incidence were found for OA at any joint, hip, knee and hand OA: a progressive increase from age 25 to 34 years, with the steepest change in slope for age group 55–64 years, peaking at 75–84 years, with a slight decrease in the age group ≥ 85 years. The pattern of age-specific consultation incidence was similar between men and women at any joint, knee and hand OA except hip OA, which showed an earlier peak in men than in women. The consultation incidence of OA was significantly higher in women, with the exception of knee OA: age-adjusted IRR 1.6 (95% CI 1.3, 1.9) for OA at any joint, 1.6 (95% CI 1.0, 2.5) for hip OA, 1.0 (95% CI 0.7, 1.2) for knee OA and 3.3 (95% CI 2.0, 5.4) for hand OA. There was no strong evidence that the female:male IRRs differed across age strata. At age 55–64 years, a higher consultation incidence of knee OA was noted among men compared with women (10.2/1000 persons vs 5.9/1000 persons), although all such age- and sex-stratified joint-specific estimates were based on small numbers (see supplementary Table S27, available at *Rheumatology* Online).

TABLE 1 Annual consultation incidence (per 1000 persons) of OA (any joint and by selected body region): CiPCA 2010^a

	Total population			Population ≥ 15 years of age			Population ≥ 45 years of age		
	Numerator	Denominator	IR (95% CI)	Numerator	Denominator	IR (95% CI)	Numerator	Denominator	IR (95% CI)
OA	Total	627	87 458	627	73 207	8.6 (7.9, 9.3)	597	37 067	16.1 (14.8, 17.5)
	Men	230	43 907	230	36 637	6.3 (5.5, 7.1)	221	18 358	12.0 (10.5, 13.7)
Hip OA	Women	396	43 548	396	36 567	10.8 (9.8, 12.0)	376	18 707	20.1 (18.1, 22.2)
	Total	109	92 587	109	78 336	1.4 (1.1, 1.7)	102	42 099	2.4 (2.0, 2.9)
Men	Men	40	45 916	40	38 646	1.0 (0.7, 1.4)	40	20 306	2.0 (1.4, 2.7)
	Women	69	46 671	69	39 690	1.7 (1.4, 2.2)	62	21 793	2.8 (2.2, 3.6)
Knee OA	Total	267	90 695	267	76 444	3.5 (3.1, 3.9)	260	40 229	6.5 (5.7, 7.3)
	Men	149	45 138	149	37 868	3.9 (3.3, 4.6)	145	19 543	7.4 (6.3, 8.7)
Women	Men	156	45 557	156	38 576	4.0 (3.4, 4.7)	153	20 684	7.4 (6.3, 8.7)
	Total	103	92 712	103	78 461	1.3 (1.1, 1.6)	100	42 219	2.4 (1.9, 2.9)
Hand OA	Men	24	46 125	24	38 855	0.6 (0.4, 0.9)	22	20 510	1.1 (0.7, 1.6)
	Women	80	46 588	80	39 607	2.0 (1.6, 2.5)	79	21 710	3.6 (2.9, 4.5)

^aCalculated using the maximum available (10 year) run-in period method. CiPCA: Consultations in Primary Care Archive; IR: incidence rate (annual consultation).

Comparison with international estimates

The estimates from CiPCA were broadly comparable to previously published international studies (Table 2), but with some exceptions. While the overall estimate for the total population fell within the range reported recently for Dutch general practice research networks [26], in the population ≥45 years of age the estimates from CiPCA were substantially lower than those reported from the Medical Service Plan health administrative database in British Columbia [10]. Incidence continued to rise sharply with age >50 years in the British Columbia study, in contrast to the slowing or levelling off seen in CiPCA and another provincial health administrative database from Alberta, Canada (supplementary Fig. S3A, available at *Rheumatology* Online) [9]. Estimates for hip and knee OA were noticeably higher in CiPCA than from the second Dutch National Survey of General Practice in 2001. In contrast, there was generally fairly close agreement between CiPCA and a large Catalan general practice database [8] for joint-specific consultation incidence estimates for the population aged ≥40–45 years as well as similar age and gender patterns (supplementary Fig. 3B, available at *Rheumatology* Online). An exception was the relatively high consultation incidence in CiPCA for knee OA among men.

Age-stratified run-in period method

The age-stratified run-in period method resulted in estimates very similar to those obtained from the maximum run-in period method when restricted to the population ≥15 years or ≥45 years (supplementary Table S28, available at *Rheumatology* Online). For example, the consultation incidence of OA estimated by the age-stratified run-in period method was 9.2 (95% CI 8.5, 9.9) for OA at any joint, 1.2 (95% CI 1.0, 1.4) for hip OA, 3.6 (95% CI 3.2, 4.0) for knee OA and 1.1 (95% CI 0.9, 1.4) for hand OA among those ≥15 years of age.

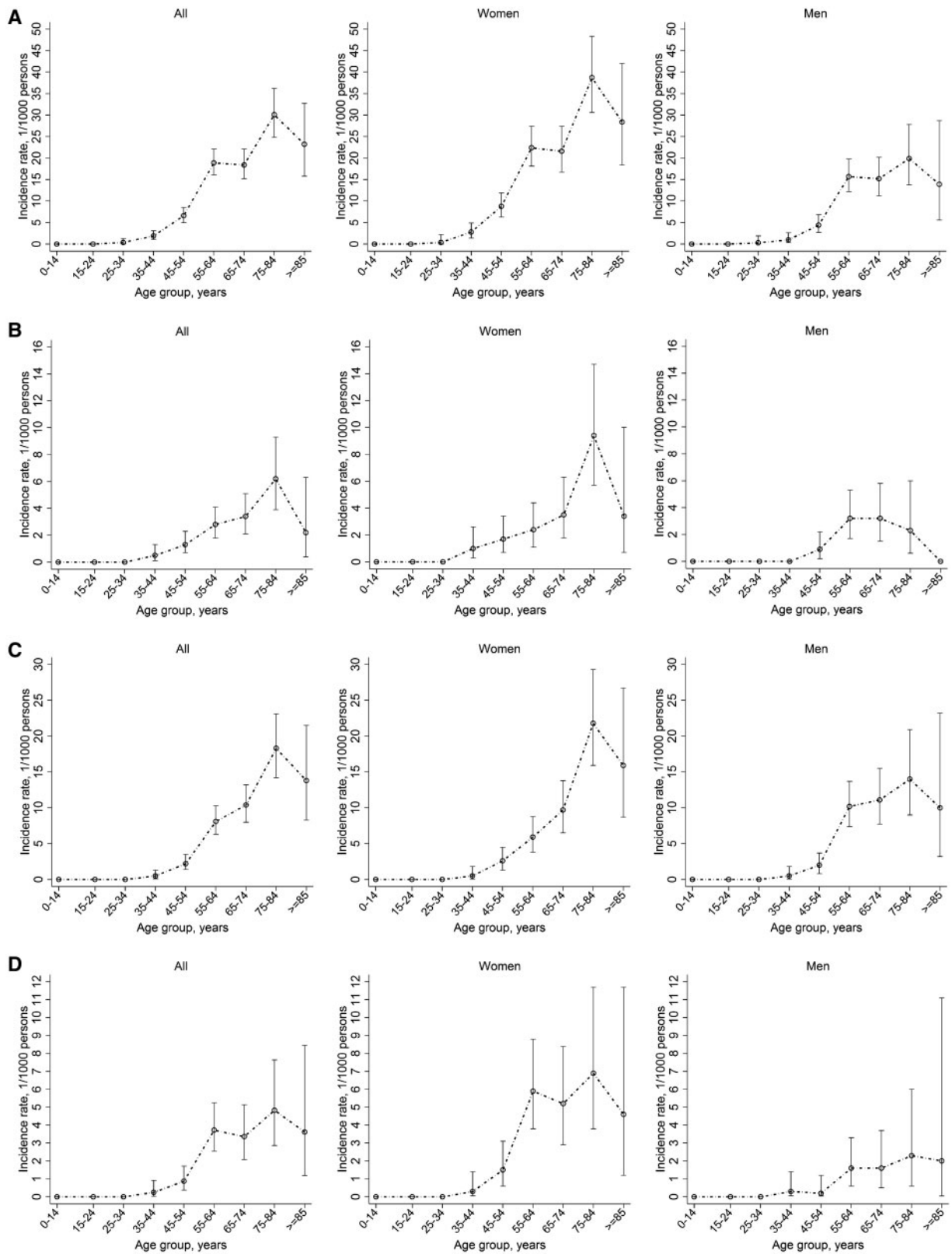
Trend in consultation incidence of OA in younger adults

The annual consultation incidence proportions of OA in those 35–44 years of age from 2003 to 2010 are presented graphically in Fig. 2, along with the 3 year moving average. The trend analysis revealed a linear increasing trend ($P=0.0173$) from an estimated annual consultation incidence of 0.3/1000 (95% CI 0.1, 0.8) in 2003 to 2.0/1000 (95% CI 1.3, 2.9) in 2010.

Discussion

To our knowledge, this is the first study to estimate the consultation incidence for OA using UK population-based health care data. We found that 1 in 100 adults is newly diagnosed with OA during the course of a year, rising to 3% of adults aged 75–84 years. Our novel age-stratified method for determining incidence revealed a trend of increasing incidence of recorded cases of OA among adults aged 35–44 years.

Fig. 1 Annual age-specific consultation incidence (per 1000 persons) of OA (any joint and by selected body region), overall and by gender: CiPCA 2010 [calculated using the maximum available (10 year) run-in period method]



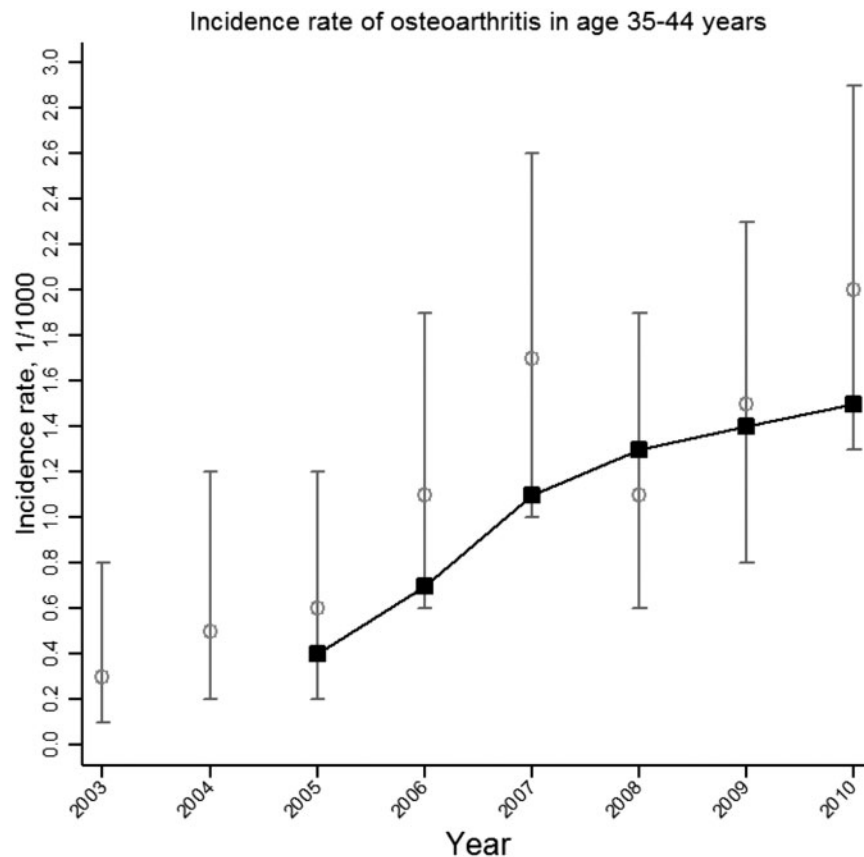
(A) OA (any joint); (B) hip OA; (C) knee OA; (D) hand OA. CiPCA: Consultations in Primary Care Archive.

TABLE 2 Comparison of crude consultation incidence estimates (per 1000 persons) with previously published international estimates

Database Location	Current study ^a	Prieto-Alhambra <i>et al.</i> [8]	van den Dungen <i>et al.</i> [23]	Kopec <i>et al.</i> [7]	Sun <i>et al.</i> [9]	Van der Waal <i>et al.</i> [6]	Kopec <i>et al.</i> [10]
Year	CiPCA North Staffordshire 2010	SIDIAP Catalonia 2006–10	Various GPRN The Netherlands 2007	MSP British Columbia 2003–04	AHCIP Alberta 2002	DNSGP-2 The Netherlands 2001	MSP British Columbia 2000–01
Consultation incidence estimate							
OA, total population	7.2		5.4–9.7	—	—		11.7
OA, total population, men	5.2			12.2	9.3 (11.14)		10.0 (13.93)
OA, total population, women	9.1			17.4	11.0 (12.70)		13.4 (19.31)
OA, population aged ≥ 15 years	8.6						14.7
OA, population aged ≥ 45 years	16.1						29.7
Hip OA, total population	1.2						
Hip OA, total population, men	0.9					0.9	
Hip OA, total population, women	1.5					0.6	
Hip OA, population aged ≥ 45 years	2.4	2.1 ^b (2.22)				1.2	
Hip OA, population aged ≥ 45 years, men	2.0	1.7 ^b (1.84)					
Hip OA, population aged ≥ 45 years, women	2.8	2.4 ^b (2.56)					
Knee OA, total population	2.9						
Knee OA, total population, men	3.3					1.5	
Knee OA, total population, women	3.4					0.9	
Knee OA, population aged ≥ 45 years	6.5	6.5 ^b (6.92)				2.0	
Knee OA, population aged ≥ 45 years, men	7.4	4.6 ^b (4.83)					
Knee OA, population aged ≥ 45 years, women	7.4	8.4 ^b (8.89)					
Hand OA, population aged ≥ 45 years	1.1	2.4 ^b (2.62)					
Hand OA, population aged ≥ 45 years, men	0.5	1.3 ^b (1.36)					
Hand OA, population aged ≥ 45 years, women	1.7	3.5 ^b (3.77)					

^aCiPCA estimates are those using the maximal available run-in period method. ^bEstimates for population aged ≥ 40 years. In SIDIAP [8], AHCIP [9] and MSP [10], age-stratified incidence was available, allowing indirect standardization for comparison with CiPCA by using the age-stratified population size in CiPCA. The standardized overall estimates are presented in the parentheses. AHCIP: Alberta Health Care Insurance Plan; CiPCA: Consultations in Primary Care Archive; DNSGP-2: Second Dutch National Survey of General Practice; GPRN: General Practice Research Networks; MSP: Medical Service Plan; SIDIAP: Sistema d'informació per al Desenvolupament de l'Investigació en Atenció Primària;

Fig. 2 Annual consultation incidence (per 1000 persons) of OA (any joint) among people aged 35–44 years between 2003 and 2010: CiPCA 2010 (calculated using the age-stratified run-in period method)



CiPCA: Consultations in Primary Care Archive. Grey bars represent annual consultation incidence; black line represents 3-year moving average.

Our estimates were generally within the range reported by previous international studies, although an area of disagreement concerns the relationship between OA consultation incidence and age. Previous studies have reported conflicting findings, showing either a steep, continuous increase in incidence with age [10] or a marked decline from the age of 60–79 years [9]. Our study suggested a plateau in incidence rates for men at age 65–74 years and an increase in women up to age 75–84 years. The exact reason for the differences in findings between studies is unclear and further studies in other databases using comparable methods are required. Differences in coding behaviour between UK primary care and Canadian administrative data, differences in the size of the denominator population for the very old age groups, the inclusion of comprehensive linked hospital data [10] and the use of a power function to model very long run-in periods [9] in the Canadian datasets may each contribute. In the UK, health conditions that are incentivized within the General Medical Services contract are likely to be more frequently diagnosed and recorded when present than non-incentivized conditions [27]. A consequence of this could be under-recording of OA and an apparent reduction in the consultation incidence proportion of OA, particularly in

older adults with multimorbidity [28]. However, a levelling off or decline in incidence rates for hand, knee and hip OA after the age of 80 years has been reported in US data using case definitions based on symptoms and radiographs [29]. Nevertheless, as has been argued before [15, 30], this pattern of declining incidence in later life must be cautiously interpreted since it may reflect biases due to competing risks, that is, individuals who would have been at high risk of incident OA in later life [30] may also have a higher mortality rate and thus may be lost to follow-up.

The observed trend in incidence among those 35–44 years old is potentially significant but must be interpreted with caution. An OA diagnosis at this age is both uncommon and potentially significant, with the age threshold conventionally used for a working diagnosis of OA being 45 years [31], implying that this younger age group with early onset has the prospect of several decades of living with the condition. However, the consultation incidence of diagnosed OA plainly does not equate to the incidence of pathology or even of symptoms [32, 33]. Instead, consultation incidence represents an event in the course of a patient's experience of associated symptoms and is determined by their decision to consult and the GP's

decision to make and record the diagnosis of OA. Observed trends may be due to changes in the thresholds for these decisions and other aspects of data quality. Although there is evidence from other primary care databases in the UK that the number of recorded general practice consultations per person-year has been increasing [34, 35], this has been less marked in younger age groups and the magnitude of this appears unlikely to explain the trend we observed. More diligent recording of secondary diagnoses and a lowering of the threshold among practitioners for making the diagnosis of OA (as opposed to using non-specific codes such as knee pain) may contribute to this trend. However, we observed the same trend of increasing incidence in adults aged 35–44 years when restricting the analysis to diagnoses made in primary care (data not shown). Between 1996–97 and 2003–04, Kopec *et al.* [7] found no evidence of an increased consultation incidence of OA in persons <40 years of age in British Columbia. However, we cannot exclude the possibility that our findings reflect a true secular change between 2003 and 2010 in the incidence of OA in adults aged 35–44 years. These cohorts grew up during periods of increasing prevalence of obesity in England [36–38], which would be expected to increase the risk of OA [39]. The contribution of injury, another important determinant of OA in younger ages, is unclear due to a lack of available data on representative populations.

The higher incidence of knee OA in men than in women in the 55–64 year age band was unexpected. This may be a chance finding based on multiple comparisons and relatively small numbers of incident cases within each age–gender stratum, although it is consistent with previous consultation prevalence in this age group [40] and radiographical OA [41] in the population.

Administrative databases and electronic health care records data are increasingly recognized as key resources for chronic disease research and surveillance [42, 43]. OA is underrepresented in these fields given its importance to population health [1, 4]. One recurrent concern, shared with all secondary uses of data recorded principally for administrative or clinical purposes, is the validity of OA case definitions. A variety of approaches have been undertaken in our and other databases internationally, including manual review of anonymized free text in the health record, typically adjudicated by one or more clinicians [44–46], comparison of rates in different databases [17, 21], comparison with independent patient self-reports [24, 25, 47] and clinical or imaging assessment [48]. As with other data sources [49], incidence and prevalence estimates derived from administrative and health record data are sensitive to the particular case definition adopted [10]. While a single record of a consultation with an OA diagnostic code has been supported in some of these studies [24, 25, 47], and is the most commonly used case definition, it must nevertheless be acknowledged as naive in that it assumes no misclassification errors [50]. Uncertainty around our estimates is likely to be greater than implied by the CIs and, together with the relatively small size of the database, means that our

findings should be replicated in other similar databases. If confirmed, an increase in newly diagnosed cases of OA in younger adulthood may have important but quite different implications depending on whether it reflects a lowering of the diagnostic threshold or the earlier onset of OA. Distinguishing between these two interpretations warrants further investigation. Secondary care information (e.g. hospital letters) is recorded and coded in CiPCA, but the completeness of this information will vary by practice. While it is unlikely (due to the generally long run-in periods that we have used) that many patients with an existing diagnosis of OA during the run-in period would not be identified in CiPCA, the number of new cases of OA in 2010 may be conservative since there may be some patients diagnosed in secondary care who were not recorded with OA in CiPCA.

Our estimates of the rate and age–sex distribution of new OA diagnoses, showing broad consistency with incidence rates derived internationally, help to address an information gap on OA occurrence in the UK. A novel method, designed to more efficiently use data within routine electronic health care databases, has provided preliminary data on a trend likely to be of public health concern but which these data alone cannot answer conclusively.

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Supplementary data

Supplementary data are available at *Rheumatology* Online.

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