EXTENDED REPORT

The incidence and prevalence of systemic lupus erythematosus in the UK, 1999–2012

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ABSTRACT

Objectives To estimate the incidence and prevalence of systemic lupus erythematosus (SLE) in the UK over the period 1999–2012.

Methods A retrospective cohort study using the Clinical Practice Research Datalink (CPRD). The incidence was calculated per 100 000 person-years and the prevalence was calculated per 100 000 people for the period 1999–2012 and stratified by year, age group, gender, region and ethnicity. Three definitions of SLE were explored: (1) systemic lupus, (2) a fully comprehensive definition of lupus including cutaneous only lupus and (3) requiring supporting evidence of SLE in the medical record.

Results Using our primary definition of SLE, the incidence during the study period was 4.91/100 000 person-years (95% CI 4.73 to 5.09), with an annual 1.8% decline (p<0.001). In contrast, the prevalence increased from 64.99/100 000 in 1999 (95% CI 62.04 to 67.93) (0.065%) to 97.04/100 000 in 2012 (95% CI 94.18 to 99.90) (0.097%). SLE was six times more common in women. The peak age of incidence was 50–59 years. There was regional variation in both incidence and prevalence. People of Black Caribbean ethnicity had increased temporal trend.11–13

The aim of this study was to examine the incidence and prevalence for SLE in the UK for the most recent decade, including temporal trends, variation with age, gender, region and ethnicity. To better understand the nature of SLE, three different definitions were examined: (1) systemic lupus, (2) a fully comprehensive definition of lupus including cutaneous only lupus and (3) any lupus case with supporting evidence of SLE in the medical record.

METHOD

Study design and study population

A retrospective cohort study was conducted using the CPRD, a longitudinal database of UK general practice records incepted in 1987 and deemed to be representative of the UK population.14 As of January 2013, there were 660 practices contributing anonymised records for approximately 12 million people in all four countries of the UK. Practices record patient demographics, consultations, hospitalisations, specialist referrals, prescriptions, test results, immunisations and diagnoses using Vision Practice Management Software. Clinical information is entered using Read codes, a standard clinical terminology system used in general practice in the UK. Prescriptions are recorded using Multilex (or British National Formulary (BNF)) codes. Researchers access data through the CPRD website or annual flat files. The quality of the data is regulated by the Medicines and Healthcare Products Regulatory Authority (MHRA).

Participants were males and females contributing data during the period 1 January 1999 to 31 December 2012. Participants were eligible from the date their practice was deemed to be contributing
Incidence and prevalence were presented by age, gender and region and expressed per 100 000 person-years or people with 95% CIs. Ten year age-bands were selected for age stratification. Direct standardisation by age and gender used the 2012 population as a reference. The trend with time and region was analysed using Poisson regression adjusting for age and gender with an interaction term. A subsequent analysis of prevalence over time also adjusted for length of data contribution.

Using only individuals from HES-linked practices, the SLE incidence and prevalence in different ethnic groups were estimated.

Data management and analysis were performed using StataMP4 software V.13 (Statacorp, Texas, USA). The significance level was 0.05.

**RESULTS**

**Incidence**

A total of 8 681 127 people contributed 55 839 080 person-years of follow-up, of whom 2740 had a new diagnosis of SLE (incidence 4.91 per 100 000 person-years; 95% CI 4.73 to 5.09).

**Temporal trend**

Table 1 shows the trend in incidence by year. There was an annual 1.8% decline (95% CI 0.8 to 2.7, p for trend <0.001) with a peak incidence in the year 2000 (5.93/100 000 person-years (95% CI 5.14 to 6.83)) and a minimum in the year 2010 (4.00/100 000 person-years (95% CI 3.45 to 4.64)). For comparison, when using a minimum of 3 years of registration, the annual incidence declined by 1.9% (95% CI 0.9 to 2.9, p for trend<0.001) with a peak of 5.34/100 000 person-years (95% CI 4.60 to 6.20) in 2000 and a minimum of 3.51/100 000 person-years (95% CI 3.00 to 4.12) in 2010.

**Age and gender**

The incidence in females was 5.8 times that of males (8.34/100 000 person-years (95% CI 8.01 to 8.69) vs 1.44/100 000 person-years (95% CI 1.30 to 1.58), p<0.001). The mean age at diagnosis was 48.9 years (SD 16.9). Women were significantly younger than men at diagnosis (48.3 years (SD 16.8) vs 52.1 years (SD 17.2), p<0.001). The peak age of incidence for women (40–49 years) was earlier than that for men (60–69 years) (figure 1A). The incidence was greater in females compared with males for all ages.

**Region**

There was geographic variation in incidence, with the lowest in the West Midlands (3.87/100 000 person-years (95% CI 3.31 to 4.43)) and the highest in the East of England (6.03/100 000 person-years (95% CI 5.36 to 6.69)) (figure 2A). There was no obvious trend with latitude, but the overall effect of region was statistically significant (p<0.001).

**Ethnicity**

Of the 8 681 127 participants, 5 408 624 were attending HES-linked practices, including 1643 incident cases (60%). Table 2 illustrates the variation in SLE incidence by ethnicity. People of Black Caribbean ethnicity had the highest incidence (31.46/100 000 person-years (95% CI 22.48 to 44.03)). In comparison, the incidence in White people was 6.73/100 000 person-years (95% CI 6.35 to 7.14). The effect of ethnic group on incidence was statistically significant (p<0.001).

**Alternative definitions**

Using the fully comprehensive definition, the incidence was 6.23/100 000 person-years (95% CI 6.03 to 6.44). In cases with

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**Case definition**

Read codes for SLE were selected from the CPRD medical dictionary browser using the search terms ‘lupus’ and ‘SLE’ (see online supplementary table S1). Three definitions were used. The first definition ‘systemic lupus’ included all codes that represented SLE or a subtype of SLE excluding cutaneous only lupus. Codes relating to a diagnostic test, a scoring system, tuberculosis, lupus pernio, drug-induced lupus or neonatal lupus were excluded. Selected codes were verified by 6 co-investigators (3 rheumatologists (FR, MD, PL), 1 epidemiologist (WZ), 1 statistician and CPRD expert (MG) and 1 general practitioner (GP) (GD)). The second definition was fully comprehensive including cases with cutaneous only subtypes. The third definition required a record of either a positive immunological blood test result (ANA, anti-dsDNA, ENA or anti-Ro), a prescription for a drug used in SLE (hydroxychloroquine, azathioprine, methotrexate, mycophenolate mofetil, ciclosporin, cyclophosphamide, rituximab, prednisolone) or a referral to a rheumatologist from cases identified in definition 2.

The date of first code was taken as the date of diagnosis. Individuals with a code for SLE but without a diagnosis date were excluded (n=49).

External validation of SLE diagnosis was not conducted because: (1) there was no additional funding or time available for this; (2) GPs would be unlikely to give a patient a Read code for SLE unless it had been confirmed by a hospital specialist (personal communication, GD); and (3) validation of other autoimmune diseases on the CPRD has shown positive predictive values of >90%.16 17

**Outcome measures and analysis**

Incidence was calculated by dividing the number of new cases of SLE by the number of person-years from eligible participants during the study period. For each year of follow-up, the entry date was the latest of 1 January, UTS date or the date of registration plus 365 days. The exit date was the earliest date of incident SLE diagnosis, death, transfer-out of a participating CPRD practice, date of last data collection from a practice or 31 December of the specified year.

Prevalent cases were all live cases of SLE on the database meeting the eligibility criteria on the 1 July of each year. Point prevalence was calculated by dividing the number of prevalent cases by the number of people on the CPRD who had UTS and ‘acceptable’ data on the 1 July of each year.

‘up-to-standard’ (UTS) data and their individual record was of ‘acceptable’ research quality as verified by the CPRD. When calculating incidence at least 12 months registration prior to study entry was required. A minimum period of 1 year following registration has been suggested to minimise the risk of prevalent cases being coded as incident cases for chronic diseases15 and was used in a previous SLE study.16 A minimum period of 3 years postregistration was analysed for comparison.

For consenting English practices, ethnic group data were available through linkage to Hospital Episode Statistics (HES). HES records patient demographics, primary diagnosis, procedures, intensive care stays and maternity data for hospital admissions since 1997 using International Classification of Diseases (ICD)-10 codes. Linked data were only used to obtain ethnic group as this was not available on the January 2013 release of the CPRD dataset.

Independent Scientific Advisory Committee for MHRA Database Research approval was obtained for this study on 4 June 2013 (Protocol 13_092).
supporting evidence, the incidence was 4.58/100,000 person-years (95% CI 4.40 to 4.76). The trends in age, gender, region, year and ethnicity were similar between definitions (see online supplementary table S2).

Prevalence
During the study period, there were 7732 eligible prevalent cases with SLE of which 6634 were female (86%).

Temporal trend
There was a linear 3.1% increasing annual trend in prevalence from 64.63/100,000 in 1999 (95% CI 61.74 to 67.62) to 97.04/100,000 in 2012 (95% CI 94.19 to 99.94) (p for trend<0.001) (table 1). Adjusting for length of follow-up, in addition to age and gender there remained an increasing linear trend of 2.5% (95% CI 2.3 to 2.8, p<0.001).

Age and gender
In every year, females had a significantly higher prevalence of SLE than males. There was an increasing trend in both genders, which was more marked in females such that the ratio between genders increased from 5.07 (95% CI 4.48 to 5.73) in 1999 to 6.75 (95% CI 6.18 to 7.38) in 2012 (figure 3).

The overall peak age of prevalence was 60–69 years. The peak age in prevalence for men (80–89 years) was later than that of women (60–69 years) (figure 1B).

Region
In 1999, the North East had the lowest prevalence (45.88/100,000 (95% CI (30.21 to 61.56)) and Northern Ireland the highest (94.42/100,000 (95% CI 73.39 to 115.44)), whereas in 2012 the East Midlands had the lowest prevalence (65.16/100,000 (95% CI 48.52 to 85.68)) and Northern Ireland remained the highest (120.12/100,000 (95% CI 103.39 to 138.78)) (figure 2B).

Ethnicity
In all, 2,812,963 of the 4,547,778 denominator population in 2012 attended HES-linked practices (61.9%) and were eligible for this analysis. Table 2 shows the variation in prevalence with ethnicity on 1 July 2012. Similar to incidence, people of Black

Table 1  SLE incidence rate (IR) and prevalence by year, 1999–2012

<table>
<thead>
<tr>
<th>Year</th>
<th>Number of incident cases</th>
<th>Person-years</th>
<th>Crude IR, per 100 000 person-years (95% CI)</th>
<th>Standardised IR*, per 100 000 person-years (95% CI)</th>
<th>Number of prevalent cases</th>
<th>Crude point prevalence, per 100 000 (95% CI)</th>
<th>Standardised point prevalence*, per 100 000 person-years (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1999</td>
<td>138</td>
<td>2,708,850</td>
<td>5.09 (4.31 to 6.02)</td>
<td>5.10 (4.25 to 5.96)</td>
<td>1,875</td>
<td>64.63 (61.74 to 67.62)</td>
<td>64.99 (62.04 to 67.93)</td>
</tr>
<tr>
<td>2000</td>
<td>191</td>
<td>3,221,486</td>
<td>5.93 (5.15 to 6.83)</td>
<td>5.87 (5.03 to 6.71)</td>
<td>2,312</td>
<td>66.67 (63.98 to 69.44)</td>
<td>66.60 (63.87 to 69.33)</td>
</tr>
<tr>
<td>2001</td>
<td>189</td>
<td>3,558,271</td>
<td>5.31 (4.61 to 6.13)</td>
<td>5.30 (4.54 to 6.06)</td>
<td>2,674</td>
<td>69.78 (67.16 to 72.47)</td>
<td>69.97 (67.30 to 72.63)</td>
</tr>
<tr>
<td>2002</td>
<td>226</td>
<td>3,850,272</td>
<td>5.87 (5.15 to 6.69)</td>
<td>5.82 (5.06 to 6.59)</td>
<td>3,062</td>
<td>73.21 (70.64 to 75.85)</td>
<td>73.77 (71.15 to 76.38)</td>
</tr>
<tr>
<td>2003</td>
<td>209</td>
<td>4,010,316</td>
<td>5.21 (4.55 to 5.97)</td>
<td>5.19 (4.49 to 5.90)</td>
<td>3,319</td>
<td>76.24 (73.67 to 78.88)</td>
<td>76.99 (74.36 to 79.61)</td>
</tr>
<tr>
<td>2004</td>
<td>182</td>
<td>4,166,948</td>
<td>4.34 (3.75 to 5.02)</td>
<td>4.34 (3.71 to 4.98)</td>
<td>3,528</td>
<td>78.00 (75.45 to 80.62)</td>
<td>78.04 (75.46 to 80.62)</td>
</tr>
<tr>
<td>2005</td>
<td>236</td>
<td>4,285,893</td>
<td>5.53 (4.87 to 6.28)</td>
<td>5.54 (4.84 to 6.25)</td>
<td>3,775</td>
<td>81.23 (78.66 to 83.86)</td>
<td>81.26 (78.66 to 83.86)</td>
</tr>
<tr>
<td>2006</td>
<td>179</td>
<td>4,326,228</td>
<td>4.14 (3.57 to 4.79)</td>
<td>4.13 (3.53 to 4.74)</td>
<td>3,902</td>
<td>83.65 (81.05 to 86.32)</td>
<td>84.39 (81.74 to 87.05)</td>
</tr>
<tr>
<td>2007</td>
<td>213</td>
<td>4,365,260</td>
<td>4.88 (4.27 to 5.58)</td>
<td>4.85 (4.20 to 5.51)</td>
<td>4,076</td>
<td>86.31 (83.68 to 89.00)</td>
<td>87.18 (84.49 to 89.86)</td>
</tr>
<tr>
<td>2008</td>
<td>213</td>
<td>4,369,525</td>
<td>4.87 (4.26 to 5.58)</td>
<td>4.87 (4.22 to 5.53)</td>
<td>4,190</td>
<td>88.62 (85.96 to 91.35)</td>
<td>88.61 (85.93 to 91.29)</td>
</tr>
<tr>
<td>2009</td>
<td>195</td>
<td>4,385,184</td>
<td>4.42 (3.84 to 5.09)</td>
<td>4.42 (3.80 to 5.05)</td>
<td>4,316</td>
<td>91.16 (88.46 to 93.92)</td>
<td>91.17 (88.46 to 93.89)</td>
</tr>
<tr>
<td>2010</td>
<td>173</td>
<td>4,324,739</td>
<td>4.02 (3.47 to 4.67)</td>
<td>4.03 (3.43 to 4.63)</td>
<td>4,382</td>
<td>93.45 (90.70 to 96.26)</td>
<td>94.61 (91.80 to 97.43)</td>
</tr>
<tr>
<td>2011</td>
<td>209</td>
<td>4,239,545</td>
<td>4.93 (4.30 to 5.65)</td>
<td>4.92 (4.26 to 5.59)</td>
<td>4,429</td>
<td>95.81 (93.01 to 98.67)</td>
<td>96.67 (93.81 to 99.53)</td>
</tr>
<tr>
<td>2012</td>
<td>187</td>
<td>4,026,564</td>
<td>4.64 (4.02 to 5.36)</td>
<td>4.64 (3.98 to 5.31)</td>
<td>4,413</td>
<td>97.04 (94.19 to 99.94)</td>
<td>97.04 (94.18 to 99.90)</td>
</tr>
</tbody>
</table>

*S by age and gender to the population of 2012.
SLE, systemic lupus erythematosus.

Figure 1  Line chart of age and gender-specific (A) incidence rate 1999–2012 and (B) point prevalence in 2012 for systemic lupus erythematosus.

Figure 3  Figure 3
Caribbean ethnicity had the highest prevalence (517.51 per 100,000 people, 95% CI 398.54 to 660.84).

Alternative definitions
The prevalence of SLE in 2012 with the fully comprehensive definition was 125.51/100,000 (95% CI 122.26 to 128.76) and in cases with supporting evidence was 88.09/100,000 (95% CI 85.38 to 90.86). The trends for year, age, gender, region and ethnicity were similar to that of the first definition, with variation in the regional differences (see online supplementary table S2).

DISCUSSION
For people with our primary definition of SLE, the incidence for 1999–2012 was 4.91 per 100,000 person-years (95% CI 4.73 to 5.09) and the prevalence in 2012 was 97.04 per 100,000 people (95% CI 94.19 to 99.94). Our findings suggest that although the prevalence is increasing, the incidence is decreasing. We confirm the female predominance, which is approximately six times that of males. The peak age of incidence is 50–59 years. There is regional variation in incidence and prevalence of SLE with the highest incidence in the East of England and the highest prevalence in Northern Ireland. Black Caribbean is the ethnic group with the highest incidence and prevalence. Similar trends were found with all 3 definitions of SLE.

Our estimates of incidence and prevalence of SLE are higher than those of previous UK studies. Although this could be due to increases with time, our incidence decreased with time. A more likely explanation is that our primary definition included subtypes of SLE such as lupus nephritis and cerebral lupus which may have been excluded from previous studies. For example, Hopkinson et al. invited cases for assessment, verified the diagnosis in person and required cases to meet the ACR criteria for SLE. This would have excluded minor cases, those with single organ disease and those unwilling to attend for assessment, which would have been included in our study. Our incidence estimate was similar to Somers et al. who used the CPRD during the period 1990–1999. However, they included cases with subacute cutaneous lupus. In contrast, Nightingale et al. had a significantly lower incidence of SLE despite including cases with discoid lupus as they required 3 years of registration before diagnosis plus supporting evidence within the medical record. Their estimate remained lower than our third definition which required supporting evidence (4.30/100,000 person-years (95% CI 4.13 to 4.48)). The incidence found with our fully comprehensive definition of lupus (6.23/100,000 person-years (95% CI 6.03 to 6.44)) is much higher than previous UK studies of SLE and is the first study in the UK to include all cutaneous and systemic subtypes enabling an estimate of the full burden of lupus in the community. Interestingly, this estimate is still lower than some international studies of SLE incidence, for example from Taiwan, Argentina and Brazil, demonstrating the variation between countries due to racial, genetic and environmental factors and also differences due to case definition and study methodology. With all 3 definitions, the estimated prevalence for SLE was above the European Union threshold for a ‘rare disease’. The observed 50% increase in prevalence with time may be true as it occurred with all 3 definitions, for example, due to improved survival in people with SLE or due to ageing of the entire UK population. However, the latter is less likely as our estimates were age standardised. It may be due to more accurate recording within electronic records as paper notes have been replaced. Previous database studies have observed that the prevalence of chronic diseases appears to increase with time due to length of data contribution, including one SLE study with a 61% increase in prevalence from 1992 to 1998. However, after adjusting for length of data contribution, our prevalence still increased with time. The decrease in incidence may be true.

or due to the possibility that prevalent cases were coded as incident cases in the early years of the study period. However, increasing the minimum registration period to 3 years there remained a small but significant negative trend with time, suggesting this was a true decrease.

When examining differences with age and gender, our study confirmed the known female predominance for people diagnosed with SLE, which was observed with every definition of SLE. The mean age of diagnosis of 49 years and peak age of incidence of 50–59 years is later than that which is conventionally taught, but is consistent with previous UK and some international studies. Males had a statistically significant later peak age of diagnosis compared with females in accord with findings from previous studies worldwide.

Although there was no clear trend with latitude, there were statistically significant regional differences in both SLE incidence and prevalence. Regional variation has been noted previously by Somers et al. who also found the West Midlands to have the lowest incidence and but in contrast to our study found Northern Ireland to have the highest incidence. Reasons for this regional variation are unclear but may be due to a combination of local environmental or genetic factors, variation in ethnic mix or socio-economic factors, differences in coding, consultation, diagnosis or referral rates or differences in health-seeking behaviour between the populations.

Ethnicity is recognised to be an important factor in the development of SLE. This is the first study to report ethnic variation in SLE incidence and prevalence using the CPRD. We found that people of Black Caribbean origin had the highest incidence and prevalence of SLE. Combining ethnic groups, people of Asian descent had the second highest incidence and prevalence after Black people, and White people had the lowest incidence and prevalence of SLE. This supports findings from previous UK studies.

The main limitations of our study are those inherent in the use of a large database such as the CPRD. The date of diagnosis and accuracy of diagnosis could not be verified which may have led to missing data and misclassification bias. Unfortunately, we were unable to externally validate our cases. However, GPs would be unlikely to Read code a diagnosis of SLE without confirmation from secondary care (personal communication, GD) and previous studies of other chronic autoimmune diseases have shown good accuracy in recording on the CPRD with positive predictive values >90%. Ideally, every case should be assessed in person, but this is impractical due to the confidentiality of the database. Another limitation was that ethnicity data were only available for individuals attending English HES-linked practices and within these practices 50% of individuals had missing ethnicity data. This may have caused an overestimation of the ethnic-specific incidence and prevalence as people with ethnicity recorded are those admitted to hospital and hence more likely to have a chronic condition such as SLE. However, this should not affect the rate ratios between ethnic groups and therefore the trends remain valid. In addition, to maintain the sample size for the analysis of region, age and gender, ethnicity could not be adjusted for. This would be an interesting future study.

In summary, the incidence of SLE in the UK is decreasing and the prevalence increasing. This increase in prevalence suggests that SLE is no longer a ‘rare’ disease, which has implications for healthcare planning. These data confirm the known predilection for SLE in women. The peak age of diagnosis is in middle age, contrary to the generally held belief that lupus mainly targets young people. There is UK-wide variation. Black Caribbean is the ethnic group with the highest incidence and prevalence. Future research on geographic variation and ethnicity is warranted.
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Contributors  All authors contributed to the conception, study design, interpretation of the data, and to the drafting and approval of the final manuscript. FR performed the analysis.

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Ethics approval  Independent Scientific Advisory Committee (ISAC) for MHRA Database Research (Protocol 13_092).

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REFERENCES


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