Seasonal variation in diagnosis of invasive cutaneous melanoma in Eastern England and Scotland

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Background: Worldwide, the incidence of cutaneous melanoma has been reported to be highest in the summer and lowest in the winter. Northern Irish data suggested seasonal variation for women only, especially those with thinner melanomas, sited on limbs. We interrogated two larger UK cancer registries for temporal differences in melanoma diagnosis and associated patient characteristics.

Methods: Melanomas diagnosed from 2006 to 2010 in the Eastern England and Scottish cancer registries (n = 11,611) were analysed by month of diagnosis, patient demographics and melanoma characteristics, using descriptive and multivariate modelling methods.

Results: More patients with melanoma were diagnosed in the summer months (June 9.9%, July 9.7%, August 9.8%) than the winter months (December 7.2%, January 7.2%, February 7.1%) and this pattern was consistent in both regions. There was evidence that the seasonal patterns varied by sex (p = 0.015), melanoma thickness (p = 0.002), body site (p = 0.006), and type (superficial spreading melanomas p = 0.005). The seasonal variation was greatest for diagnosis of melanomas occurring on the limbs.

Conclusion: This study has confirmed seasonal variation in melanoma diagnosis in Eastern England and Scotland across almost all population demographics and melanoma characteristics studied, with higher numbers diagnosed in the summer months, particularly on the limbs. Seasonal patterns in skin awareness and related help-seeking are likely to be implicated. Targeted patient interventions to increase sun awareness and encourage year-long skin inspection are warranted.

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1. Introduction

Cutaneous melanoma is now the fifth commonest cancer diagnosed in the United Kingdom, following a rapid rise in incidence over the last few decades. While 11 cases per 100,000 population were diagnosed in 1999–2001, the age-standardised rate had increased by 55% to around 17 cases per 100,000 population in 2008–2010 [1]. The increase has been seen across sex and age groups, but most significantly among older men [2]. Importantly, more than a quarter of new melanoma cases were diagnosed in people aged less than 50 years in 2010, in contrast to only 11% among all cancers combined [1].

It is possible that some of the increase in incidence may be due to improved surveillance and earlier detection, as well as changes in diagnostic criteria [3]. However, most of the increasing incidence trends are considered to be due to exposure to ultra-violet radiation through increased frequency of sunbed use, intermittent unaccustomed exposure especially in childhood, and leisure-time exposure including holidays abroad and outdoor sport [4–6]. Approximately 86% of melanomas diagnosed in the UK in 2010 were estimated to be linked to exposure to ultra-violet radiation from the sun and sunbeds [7].

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There is well documented seasonal variation in rates of melanoma diagnosis worldwide. Among light-skinned populations the incidence of melanoma is highest in summer and lowest in winter, irrespective of latitude [8,9]. However, detailed analysis of seasonal variation within the UK is limited to small datasets, and only data collected up until 2006: in the Oxford Region using data routinely collected between 1952 and 1975 [10], and in Northern Ireland where an analysis of data collected between 1984 and 2006 confirmed seasonal variation for women only, especially those with thinner melanomas and tumours diagnosed on the limbs [11]. We conducted an analysis of routinely collected clinical data from the Eastern England and Scottish cancer registries between 2006 and 2010, and compared diagnosis by month of the year, taking into account key factors including patient demographics as well as melanoma characteristics relevant to predicting disease outcome.

2. Materials and methods

2.1. Patient cohorts

We analysed routinely collected clinical data regarding invasive cutaneous melanoma registered by the National Cancer Registration Service- Eastern Office and the Scottish Cancer Registry between 1st January 2006 and 31st December 2010. While the regions have a comparable population size at approximately 5.7 and 5.2 million respectively, they are geographically distinct and have diverse socio-demographic characteristics. Primary sources of information included electronic and paper-based reports and clinical notes from hospitals and pathology laboratories. Recent reports have highlighted the completeness of recorded Breslow thickness for all melanomas, at more than 85% across both registries since 2006 (C15 Vol X).

We abstracted data items related to patient, disease and temporal factors. Demographic variables included sex, age group at diagnosis (two groupings were used: <30 years, 30–49, 50–64 year and ≥65 years for comparison with crude incidence rates, and <50 years, 50–64 years, 65–74 years, ≥75 years for other analyses), and national quintile of small area measures of deprivation (using income domain of either the English or Scottish Index of Multiple Deprivation as applicable).

Disease data included melanoma Breslow thickness, histological type and body site of occurrence. Breslow thickness data were split into two categories: <2 mm and ≥2 mm. Breslow thickness was reported in millimetres for 94% of the Scottish cases, and 89% of the Eastern England cases (17 Eastern England cases had Breslow thickness reported as group only). Histological types of melanomas were categorised using the ICDO(3) morphology codes: superficial spreading melanoma (SSM), nodular melanoma (NM), lentigo maligna melanoma (LMM), unclassified/unknown melanoma (UM) (codes 8720 NOS, 8723 NOS regressing), and others (OM) (codes 8722 Balloon cell, 8730 Amelanotic, 8740 Malignant melanoma in junctional naevus, 8741 Malignant melanoma in precancerous melanosis, 8744 Acral lentiginous, 8745 Desmoplastic, 8746 Mucosal lentiginous melanoma, 8761 Melanoma arising in congenital melanocytic naevus, 8770 Mixed epithelial and spindle cell, 8771 Epithelioid cell melanoma, 8772 Spindle cell NOS, 8780 Blue naevus malignant). Anatomical site was coded using the International Classification of Diseases (ICD) 10 (four digit) as follows: C430 (lip), C431 (eyelid), C432 (ear), C433 (other and unspecified parts of face), and C434 (scalp and neck) were located on ‘Head and Neck’, C435 corresponded to ‘Trunk’, C436 to ‘Upper Limb’, C437 to ‘Lower Limb’, C438 (other specified sites of skin) and C439 (site unspecified) to ‘Other’ sites. Melanomas occurring in the eye or genital organs, and in situ melanomas were excluded.

Temporal differences were characterised by the month and year of melanoma diagnosis.

2.2. Statistical analysis

Initial descriptive analyses were undertaken to compare cases per year between the two regions, and to compare stratified Breslow thickness by patient demographics, tumour characteristics, and month of diagnosis. Multivariate analyses were then undertaken using a negative binomial regression to model the number of cases diagnosed in each month (outcome). The main exposure of interest was seasonal variation which was modelled using sine and cosine components with a period of one year. The model further included fixed effect exposures for sex, age, deprivation, region, Breslow thickness (as a binary variable <2 mm or ≥2 mm), site and histological type. Finally a longer term trend in incidence was accounted for with a cubic spline with 3 knots (a pragmatic choice providing more flexibility than a linear trend whilst remaining reasonably parsimonious). A negative binomial model was used as initial investigations suggested that more variation existed than suggested by the Poisson distribution (i.e. there was over-dispersion). As with Poisson regression, the negative binomial framework models counts rather than rates and so an offset equal to the log of the person time at risk needs to be included such that the outputs from the model may be interpreted as rate ratios. To do so we use the population at risk in each age by sex by deprivation group. This was calculated using 2008 national statistics aggregated up from the lower super output area level (a lower super output area is a geographic region defined for reporting of UK census data, each containing a population of around 1500 people). When initially specifying the model more complex parametrisations of seasonal variation were considered, however, early investigations suggested that higher order Fourier components did not significantly improve the fit of the model.

In addition to considering the overall seasonal trend we were also interested in whether the seasonal variation in incidence was dependant on other factors. In order to investigate this we considered interactions between the sine and cosine components and other variables retaining only those found to be statistically significant. Where more than one variable was found to have a significant interaction, interactions between those variables were also considered. As a supplementary analysis we repeated the final regression model treating Breslow thickness as a 4 category model (≤1 mm, 1–1.99 mm, 2–3.99 mm and ≥4 mm). Data analysis was undertaken using Stata 13 (Stata Corporation, College Station, TX).

3. Results

3.1. Sample description and incidence

A total of 11,611 invasive cutaneous melanoma cases were registered in both regions from 2006 to 2010: 5998 in Eastern England and 5613 in Scotland. The number of cases and crude incidence is shown in Table 1 by year, patient demographics and tumour characteristics. Analysis of overall melanoma incidence rates demonstrated a steady rise in numbers detected over the five years by 21% from 92 to 112 per 100,000 people per year. The incidence of melanomas was slightly higher in women than in men (109 versus 102 per 100 000), with higher incidence of diagnosed melanomas in the older age groups (<30 years 13 per 100,000 compared to 253 per 100,000 for ≥65 years). The incidence of melanoma decreased with increasing levels of deprivation (‘least deprived’ group 132 per 100,000 versus 71 per 100,000 in the ‘most deprived’ group). The majority (67%) of melanomas were
relatively thin, and the commonest histological type was superficial spreading melanoma (54%). Slightly more melanomas were located on the trunk (29%) and lower limbs (27%) than on the upper limbs (21%) and head/neck areas (20%).

### 3.2. Seasonal variation in incidence

For the whole patient cohort, more melanomas were diagnosed in the summer months (June 9.9%, July 9.7%, August 9.8%) than the winter months (December 7.2%, January 7.2%, February 7.1%) and this pattern was consistent for both regions studied. Seasonal variation was evident when analysed by patient sex (particularly among women) and age (most marked in those aged less than 50 years) (Table 2). Seasonal variation was also evident when analysed by Breslow thickness, histological type and body site of occurrence (Table 3).

Multivariate analysis showed that after adjusting for patient and melanoma characteristics, there was very strong evidence ($p < 0.001$) of seasonal variability in melanoma diagnosis. The average seasonal variability predicted from a model without interaction terms is shown in Fig. 1 as rate ratios compared to 1st January each year. There was substantial variation in incidence over a year with rates over 30% higher in the summer months than the winter months and a peak incidence occurring in July (peak to peak rate ratio = 1.35 95% confidence interval [CI] 1.27, 1.43). This analysis excluded those patients with missing Breslow thickness, but similar patterns were seen in a model excluding tumour factors on the full data set, indicating no obvious bias by using a complete case analysis.

In multivariate analysis using interaction terms we found no evidence that the seasonal variability was different for patients of different ages ($p = 0.51$), deprivation ($p = 0.56$) or region ($p = 0.56$). The seasonal variation by sex, Breslow thickness, body site and histological type is shown in Fig. 2; there is evidence of each being a moderating factor ($p = 0.015$ for sex, $p = 0.002$ for Breslow thickness, $p = 0.006$ for body site, and $p = 0.005$ for histological
Table 2
Distribution of melanoma incidence (%) by month of diagnosis and patient demographics (n=11,611).

| Month of diagnosis | Jan | Feb | Mar | Apr | May | Jun | Jul | Aug | Sep | Oct | Nov | Dec |
|--------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| Frequency (%)      | 834 (7.18) | 829 (7.14) | 918 (7.91) | 849 (7.31) | 979 (8.43) | 1151 (9.91) | 1129 (9.72) | 1138 (9.80) | 971 (8.36) | 1036 (8.92) | 940 (8.10) | 837 (7.21) |
| Sex                |     |     |     |     |     |     |     |     |     |     |     |     |
| Men, %             | 390 (7.12) | 396 (7.22) | 436 (7.95) | 382 (6.97) | 452 (8.25) | 529 (9.65) | 502 (9.16) | 555 (10.13) | 441 (8.05) | 489 (8.92) | 465 (8.48) | 444 (8.10) |
| Women, %           | 444 (7.24) | 433 (7.06) | 482 (7.86) | 467 (7.62) | 527 (8.60) | 622 (10.15) | 627 (10.23) | 583 (9.51) | 530 (8.65) | 547 (8.92) | 475 (7.75) | 393 (6.41) |
| Age group          |     |     |     |     |     |     |     |     |     |     |     |     |
| <50 yrs            | 228 (7.26) | 213 (6.79) | 272 (8.67) | 248 (7.90) | 280 (8.92) | 287 (9.14) | 333 (10.61) | 308 (9.81) | 267 (8.51) | 262 (8.35) | 232 (7.39) | 209 (6.66) |
| 50–64 yrs          | 227 (6.83) | 254 (7.64) | 265 (7.97) | 241 (7.25) | 259 (7.79) | 339 (10.20) | 314 (9.44) | 342 (10.29) | 289 (8.69) | 295 (8.87) | 263 (7.91) | 237 (7.13) |
| 65–74 yrs          | 185 (7.82) | 169 (7.14) | 173 (7.31) | 155 (6.35) | 195 (8.24) | 236 (9.97) | 216 (9.13) | 242 (10.22) | 177 (7.48) | 229 (9.67) | 216 (9.13) | 174 (7.35) |
| 75+ yrs            | 194 (6.98) | 193 (6.94) | 208 (7.48) | 205 (7.37) | 245 (8.81) | 289 (10.40) | 266 (9.57) | 246 (8.85) | 238 (8.56) | 250 (8.99) | 229 (8.24) | 217 (7.81) |
| Region             |     |     |     |     |     |     |     |     |     |     |     |     |
| Scotland           | 402 (7.16) | 400 (7.13) | 459 (8.18) | 404 (7.20) | 487 (8.68) | 559 (9.96) | 505 (9.00) | 550 (9.80) | 472 (8.41) | 494 (8.80) | 476 (8.48) | 405 (7.22) |
| Eastern England    | 432 (7.20) | 429 (7.15) | 459 (7.65) | 445 (7.42) | 492 (8.20) | 592 (9.87) | 624 (10.40) | 588 (8.90) | 499 (8.32) | 542 (9.04) | 464 (7.74) | 432 (7.20) |
| Deprivation quintiles |     |     |     |     |     |     |     |     |     |     |     |     |
| 1st most affluent  | 227 (7.32) | 209 (6.94) | 243 (7.84) | 229 (7.39) | 270 (8.71) | 300 (9.68) | 309 (9.97) | 315 (10.16) | 245 (7.90) | 293 (9.45) | 244 (7.89) | 216 (6.97) |
| 2%                 | 206 (7.19) | 213 (7.43) | 228 (7.96) | 225 (7.85) | 226 (7.89) | 300 (10.47) | 272 (9.49) | 269 (9.39) | 231 (8.06) | 239 (8.34) | 254 (8.86) | 203 (7.08) |
| 3%                 | 188 (6.98) | 196 (7.28) | 203 (7.54) | 188 (6.98) | 244 (9.06) | 273 (10.14) | 282 (10.47) | 257 (9.54) | 247 (9.17) | 241 (8.95) | 187 (6.94) | 187 (6.94) |
| 4%                 | 140 (7.43) | 135 (7.16) | 145 (7.69) | 144 (7.64) | 162 (8.59) | 180 (9.55) | 180 (9.55) | 165 (8.85) | 187 (9.92) | 156 (8.82) | 168 (8.91) | 159 (8.44) |
| 5th most deprived  | 73 (6.84)  | 76 (7.12)  | 99 (9.28)  | 63 (5.90)  | 77 (7.22)  | 98 (9.18)  | 101 (9.47) | 110 (10.31) | 92 (8.62)  | 95 (8.90)  | 96 (9.00)  | 87 (8.15)  |

4.2. Comparison with existing literature

Our results contrast notably with those of Chaillol and colleagues [11], based on Northern Ireland Cancer Registry data, which only found seasonal variation in women with thin melanomas arising on the limbs. They did not observe seasonal variation in older men, thinner melanomas arising on the trunk, head or neck, or any thicker melanomas irrespective of age, sex and body site. The reasons for this contrast with our results are unclear, although the sample size of the Northern Irish study was five-fold smaller, therefore there may have not been enough power to detect such associations. The issue of power was further compounded by the stratified approach they took rather than the approach of using interactions that we have employed. Furthermore, our data were collected between 2006 and 2010 (compared with Chaillol et al’s analyses using data from cases diagnosed between 1984 and 2006), so that important differences in more recent patterns of UV exposure, social conventions regarding exposure of the skin, and help-seeking behaviour for skin changes in earlier decades, may also play a role.

Chaillol et al. also undertook a systematic literature review of studies examining whether seasonal variations were associated with sex, age, body site, histological type or Breslow thickness. They identified 15 studies (Europe (Norway, Germany, Greece, Italy, UK) 7, USA 5, Australia 2, Brazil 1) reporting data from 281 to 32,868 melanoma cases diagnosed between 1960 and 2006 [8–10, 12–23]. Overall, they found that seasonal variations were more marked in women than men, in younger ages, and on the limbs more than the trunk or head and neck; our findings are strikingly similar. Seven studies reported greatest seasonal variation in superficial spreading melanomas, with three studies also reporting seasonal variation with lentigo maligna melanomas. Five studies found seasonal variation in thin lesions with none in thick lesions, while one Greek study found no seasonal variation with Breslow thickness. A more recent study by Keller and colleagues [24], set in Bavaria in Germany and analysing 11,501 malignant melanomas registered from 2003 to 2008 again shows contrasting results: in this population seasonal variation was most evident in thicker (>2 mm) melanomas on the limbs. The only other previous study set in the UK reported the month of first attendance (rather than...
Table 3
Distribution of melanoma incidence (%) by month of diagnosis and melanoma characteristics.

| Month of diagnosis | Jan | Feb | Mar | Apr | May | Jun | Jul | Aug | Sept | Oct | Nov | Dec |
|--------------------|-----|-----|-----|-----|-----|-----|-----|-----|------|-----|-----|-----|
| Breslow thickness   |     |     |     |     |     |     |     |     |      |     |     |     |
| Total %            | 834 (7.18) | 829 (7.14) | 918 (7.91) | 849 (7.31) | 1151 (9.91) | 1129 (9.72) | 1138 (9.80) | 971 (8.36) | 1036 (8.92) | 940 (8.10) | 837 (7.21) |     |
| <2 mm %            | 547 (7.02) | 536 (6.88) | 590 (7.58) | 572 (7.35) | 644 (8.27) | 765 (9.82) | 746 (9.58) | 790 (10.15) | 690 (8.86) | 703 (9.03) | 642 (8.24) |     |
| ≥2 mm %            | 216 (7.53) | 217 (7.57) | 246 (8.58) | 207 (7.22) | 275 (9.59) | 290 (10.11) | 288 (10.04) | 258 (9.00) | 209 (7.29) | 242 (8.44) | 231 (8.05) |     |
| Unknown %          | 71 (7.43) | 76 (7.95) | 82 (8.38) | 72 (7.30) | 60 (9.44) | 95 (9.94) | 90 (9.41) | 72 (7.33) | 61 (9.52) | 67 (7.60) | 86 (8.90) |     |
| Distribution        |     |     |     |     |     |     |     |     |      |     |     |     |
| Superficial spreading %  | 419 (6.74) | 426 (6.85) | 462 (7.43) | 456 (7.33) | 501 (8.05) | 613 (9.85) | 641 (10.03) | 624 (10.03) | 532 (8.55) | 573 (9.21) | 512 (8.23) | 462 (7.43) |
| Nodular %          | 106 (7.26) | 90 (6.64) | 121 (8.31) | 108 (7.97) | 119 (8.78) | 136 (10.04) | 133 (9.82) | 128 (9.45) | 107 (8.73) | 113 (8.34) | 104 (7.68) | 89 (6.59) |
| Lentigo maligna %  | 83 (7.25) | 76 (6.64) | 94 (8.21) | 87 (7.60) | 111 (9.69) | 114 (9.69) | 98 (8.56) | 103 (9.0) | 101 (8.82) | 100 (8.73) | 98 (8.56) | 80 (6.99) |
| Unclassified/unknown % | 61 (7.61) | 171 (8.08) | 176 (8.32) | 156 (7.37) | 182 (8.60) | 210 (9.92) | 196 (9.26) | 208 (9.83) | 162 (7.66) | 195 (9.22) | 199 (7.51) | 140 (6.62) |
| Other %            | 65 (8.40) | 66 (8.53) | 65 (8.40) | 42 (5.43) | 66 (8.53) | 78 (10.08) | 72 (7.33) | 72 (7.33) | 72 (7.33) | 72 (7.33) | 67 (6.66) | 86 (8.53) |
| Histological type   |     |     |     |     |     |     |     |     |      |     |     |     |
| Nodular %          | 106 (7.26) | 90 (6.64) | 121 (8.31) | 108 (7.97) | 119 (8.78) | 136 (10.04) | 133 (9.82) | 128 (9.45) | 107 (8.73) | 113 (8.34) | 104 (7.68) | 89 (6.59) |
| Lentigo maligna %  | 83 (7.25) | 76 (6.64) | 94 (8.21) | 87 (7.60) | 111 (9.69) | 114 (9.69) | 98 (8.56) | 103 (9.0) | 101 (8.82) | 100 (8.73) | 98 (8.56) | 80 (6.99) |
| Other %            | 65 (8.40) | 66 (8.53) | 65 (8.40) | 42 (5.43) | 66 (8.53) | 78 (10.08) | 72 (7.33) | 72 (7.33) | 72 (7.33) | 72 (7.33) | 67 (6.66) | 86 (8.53) |
| Anatomical site     |     |     |     |     |     |     |     |     |      |     |     |     |
| Head/neck %        | 185 (7.81) | 186 (7.85) | 179 (7.56) | 188 (7.94) | 207 (8.74) | 216 (9.12) | 214 (9.04) | 222 (9.38) | 194 (8.19) | 194 (8.19) | 201 (8.49) | 182 (7.69) |
| Trunk %            | 257 (7.62) | 256 (7.59) | 270 (8.00) | 238 (7.66) | 273 (8.15) | 341 (10.11) | 314 (9.31) | 322 (9.55) | 270 (8.00) | 296 (8.78) | 270 (8.00) | 264 (7.83) |
| Upper limb %       | 165 (6.65) | 173 (6.08) | 183 (7.37) | 165 (6.65) | 210 (8.46) | 250 (10.07) | 255 (10.27) | 261 (10.52) | 222 (8.94) | 224 (9.02) | 196 (7.90) | 178 (7.17) |
| Lower limb %       | 205 (6.54) | 187 (5.96) | 260 (8.29) | 233 (7.43) | 271 (8.64) | 328 (10.46) | 322 (10.27) | 316 (10.08) | 269 (8.58) | 296 (9.44) | 259 (8.26) | 190 (6.06) |

*a Unspecified and other sites n=252 (Jan 22; Feb 27; Mar 26; Apr 25; May 16; Jun 16; Jul 24; Aug 17; Sep 16; Oct 26; Nov 14; Dec 23).
Fig. 2. (a) Seasonal variability in Superficial Spreading Melanoma incidence by melanoma thickness, body site and patient sex predicted from a model including interaction terms (rate ratios compared to 1st January). (b) Seasonal variability in melanoma incidence for types other than Superficial Spreading Melanoma by melanoma thickness, body site and patient sex predicted from a model including interaction terms (rate ratios compared to 1st January).
lesion on seasonal variations in melanoma diagnosis is complex, and other factors may also be important. Previous research from an Italian cohort has indicated an excess risk of melanoma and other skin cancer for people born in Spring or Autumn months [30], which nonetheless would not explain the patterns of excess risk in the summer months observed in our study.

4.3. Strengths and weaknesses of the study

The main strength of this study is that it used population-based data of high quality and completeness from two regions covering >10 million UK population. The most important prognostic factor, the Breslow thickness, was reported in millimetres for about 90% of the cohort. Furthermore, we were able to robustly account for potential confounding of seasonal variation by patient or tumour case-mix; in addition we examined the potential effect modification by the same factors.

This report has a number of limitations inherent in observational studies. Most importantly, the data are based on the accuracy of the information recorded at the National Cancer Registration Service and the Scottish Cancer Registry. We are not able to comment on time to presentation to the GP, or time from first seeing their GP to referral and histological diagnosis, as these data are not collected routinely by the registries – however it is highly unlikely that access to GPs and referral waiting times will exhibit such pronounced patterns of seasonal variation that would be necessary to explain the observed findings. We acknowledge that data from the two registries may not reflect national data, but suggest that the findings are unlikely to differ greatly in other UK regions.

4.4. Implications for practice and future research

Although there remains insufficient evidence for the introduction of national skin cancer screening programmes [31], a recent review suggested that melanoma early detection programmes aimed at high-risk individuals might be cost-effective [32]. Until then, raising awareness of skin cancer to promote timely diagnosis is vital [33]. Skin cancer awareness campaigns need not only to provide information on the early signs and symptoms that should prompt help-seeking (as well as encouragement to seek advice), but could capitalise on evidence of seasonal variation in diagnosis, which indicates that at least part of the population are more ‘skin-aware’ and show greater readiness for seeking medical help for suspected skin lesions during the summer months. This does not of course negate the need for ‘skin awareness’ throughout the year.

The beneficial role of friends and relatives in encouraging help-seeking behaviours in those who have a skin lesion has been documented [28,34]. There is evidence that such sanctioning of help seeking by others can be an important factor in reduced time to presentation [29,35], and again there is potential for skin cancer awareness campaigns to capitalise on this, particularly during the summer months. Similarly, there is scope for interventions among non-clinician professionals such as salon workers, chiropodists and others [36,37].

In conclusion, contemporary evidence from a large and geographically diverse population of UK patients with melanoma indicates notable seasonal variation in diagnosis, most likely demonstrating congruent seasonal variation in awareness of skin changes on self or others, and related help-seeking behaviour.

Author contributions

The study was initiated by FMW and CC in the context of a Cancer Research UK/National Awareness and Early Diagnosis Initiative funded project reference: C8640/A12226: ‘Why do some patients present with thicker melanomas? A qualitative exploration of patients’ symptom detection, help-seeking decisions, and experiences of the pathways to diagnosis’. Methods were further developed and modified in discussions with GL, GAA and all other authors. GAA performed most analyses, with JM and HB. DG and DB oversaw the provision of data from the National Cancer Registry Service- Eastern Region and the Scottish Cancer Registry respectively. All authors commented on the methods and findings, and saw and approved the final draft of the paper.

Ethics approval

Not required. An integral part of the Regional Cancer Registries was that anonymous data may be used for research purposes.

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Conflict of interests

The authors declare no conflicts of interest.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.canep.2015.06.006.

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