

disproportionate increases in the excision of benign skin lesions.²⁷ Some of these lesions will appear and be treated as invasive melanoma, with all that it signifies to the patient, even though they have no capacity to metastasise.¹⁷ Thus an adequate evaluation of the cost effectiveness of different approaches to promoting the early diagnosis of melanoma should be a high priority for research.

Funding: None.

Conflict of interest: None.

- 1 Armstrong BK, Kricger A. Cutaneous melanoma. *Cancer Surv* 1994;19/20:219-40.
- 2 Giles GG, Dwyer T, Coates M, Bonnett A, Ring I, Hatton WM, et al. Trends in skin cancer in Australia: an overview of the available data. *Menzies Foundation* 1989;15:143-7.
- 3 Beardmore GL. The epidemiology of malignant melanoma in Australia. In: WH McCarthy, ed. *Melanoma and skin cancer*. Sydney: Government Printer, 1972: 39-64.
- 4 McCarthy WH, Martyn AL, Roberts G, Dobson A. Melanoma in New South Wales 1970-1976: confirmation of an increased incidence. *Med J Aust* 1980;2:137-40.
- 5 Lancaster HO. Some geographical aspects of the mortality from melanoma in Europeans. *Med J Aust* 1956;1:1082-7.
- 6 Holman CDJ, James IR, Gattley PH, Armstrong BK. An analysis of trends in mortality from malignant melanoma in Australia. *Int J Cancer* 1980;26:703-9.
- 7 Giles GG, Thursfield VJ, Staples MP. The bottom line: cancer mortality trends in Australia 1950-1991. *Cancer Forum* 1994;18:12-23.
- 8 Venzon DJ, Moolgavkar SH. Cohort analysis of malignant melanoma in five countries. *Am J Epidemiol* 1984;119:62-70.
- 9 Elwood JM, Swerdlow AJ, Cox B. Trends in incidence and mortality from cutaneous melanoma in England and Wales. *Transactions of the Menzies Foundation* 1989;15:131-5.
- 10 Swerdlow AJ. International trends in cutaneous melanoma. *Ann NY Acad Sci* 1990;609:235-51.
- 11 Thorn M, Bergstrom R, Adami H-O, Ringborg U. Trends in mortality rates from malignant melanoma in Sweden 1953-1987 and forecasts to 2007. *Br J Cancer* 1992;66:563-7.
- 12 Smith T. The Queensland melanoma project—an exercise in health education. *BMJ* 1979;ii:253-4.
- 13 Armstrong BK, Howell CM. Trends in mortality from melanoma in Australia. *Med J Aust* 1987;147:150.
- 14 Jelfs P, Giles GG, Shugg D, Coates M, Durling G, Fitzgerald P, et al. Cutaneous malignant melanoma in Australia 1989. *Med J Aust* 1994;161:182-7.

- 15 MacLennan R, Green AC, McLeod GRC, Martin NG. Increasing incidence of cutaneous melanoma in Queensland, Australia. *J Nail Cancer Inst* 1992;84:1427-32.
- 16 Jones ME, Shugg D, Dwyer T, Young B, Bonnett A. Interstate differences in incidence and mortality from melanoma: a re-examination of the latitudinal gradient. *Med J Aust* 1992;157:373-8.
- 17 Burton R, Armstrong B. Recent incidence trends imply a non-metastasising form of invasive melanoma. *Melanoma Res* 1994;4:107-13.
- 18 Kricger A, Armstrong BK, Jones ME, Burton RC. *Health, solar UV radiation and environmental change*. Lyons: International Agency for Research on Cancer, 1993. (IARC Technical Report 13.)
- 19 Scotto J, Pitcher H, Lee JAH. Indications of future decreasing trends in skin melanoma mortality among whites in the United States. *Int J Cancer* 1991;49:490-7.
- 20 Roush GC, McKay L, Holford TR. A reversal in the long-term increase in deaths attributable to malignant melanoma. *Cancer* 1992;69:1714-20.
- 21 Bonnett A, Roder D, Esterman A. Epidemiological features of melanoma in South Australia: implications for cancer control. *Med J Aust* 1989;151:502-9.
- 22 Coates M, McCredie M, Taylor R. *Cancer in New South Wales; incidence and mortality 1990*. Sydney: Cancer Council, 1994.
- 23 Giles GG, Farrugia H, Thursfield V, Staples M. *Canstar: skin cancer*. Melbourne: Anti-Cancer Council, 1994.
- 24 Streetly A, Markowe H. Changing trends in the epidemiology of malignant melanoma: gender differences and their implications for public health. *Int J Epidemiol* 1995;24:897-907.
- 25 Cooke K. Primary malignant melanoma in four regions of New Zealand. *N Z Med J* 1992;105:303-6.
- 26 Mackie RM, Hole D. Audit of public education campaign to encourage earlier detection of malignant melanoma. *BMJ* 1992;304:1012-5.
- 27 Burton RC, Coates MS, Hersey P, Roberts G, Chetry MP, Chen S, et al. An analysis of a melanoma epidemic. *Int J Cancer* 1993;55:765-70.
- 28 Holman CDJ, Armstrong BK. Cutaneous malignant melanoma and indicators of total accumulated exposure to the sun: analysis by histogenetic types. *J Nail Cancer Inst* 1984;72:257-66.
- 29 Khlal M, Vail A, Parkin M, Green A. Mortality from melanoma in migrants to Australia: variation by age at arrival and duration of stay. *Am J Epidemiol* 1992;135:1103-13.
- 30 Davis NC, Herron JJ. Queensland melanoma project: organisation and a plea for comparable surveys. *Med J Aust* 1966;1:643-4.
- 31 Green A. Incidence and reporting of cutaneous melanoma in Queensland. *Australian Journal of Dermatology* 1982;23:105-9.
- 32 Cristofolini M, Bianchi R, Boi S, Decarli A, Micciolo R, Cristofolini P, et al. Effectiveness of the health campaign for the early detection of cutaneous melanoma in Trentino, Italy. *J Dermatol Surg Oncol* 1993;19:117-20.
- 33 Hill D, White V, Marks R, Borland R. Changes in sun-related attitudes and behaviours, and reduced sunburn prevalence in a population at high risk of melanoma. *Eur J Cancer Prev* 1993;2:447-56.

(Accepted 24 January 1996)

Incidence and thickness of primary tumours and survival of patients with cutaneous malignant melanoma in relation to socioeconomic status

Rona M MacKie, David J Hole

Abstract

Objective—To study incidence of and survival from cutaneous malignant melanoma in relation to socioeconomic status.

Design—Application of Carstairs deprivation score to all malignant melanoma patients diagnosed in a geographically defined area over a 15 year period.

Setting—West of Scotland (area population 2 716 900).

Subjects—3142 patients first diagnosed with malignant melanoma in the period 1979-93.

Interventions—Surgical excision of primary malignant melanoma with additional treatment as appropriate and follow up until December 1994.

Main outcome measures—Malignant melanoma incidence, primary tumour thickness and five year survival by socioeconomic status

Results—From 1979 to 1993, the age standardised incidence rate for cutaneous malignant melanoma was 9.1/100 000 for the most affluent men and 2.4/100 000 for the least affluent men and 16.1/100 000 and 5.0/100 000 respectively for most and least affluent women ($P < 0.001$ for trend in both). The incidence increased steadily over time in both sexes in all socioeconomic groups. Good prognosis tumours (<1.5 mm thick) were most common in the most affluent men and women, and over the

study period the proportion of such tumours increased most in the intermediate affluence group (both sexes) and in the least affluent group (both sexes) and in the least affluent women. Five year disease free survival from melanoma for the sexes combined was 81% for most affluent, 77% for intermediate, and 73% for least affluent groups. Even after adjustment for known prognostic factors of tumour thickness, ulceration, age, and body site of primary melanoma, the more affluent the group, the better the survival.

Conclusion—Although the incidence of cutaneous malignant melanoma is higher among more affluent people, the prognosis is better in this group than for less affluent individuals. Early diagnosis campaigns should be targeted particularly to less affluent men and primary prevention campaigns should emphasise the greater risk in more affluent women.

Introduction

Cutaneous malignant melanoma has been reported to be a disease of those in higher paid occupations.^{1 2} This could relate to differences in exposure to the sun and to affluence: short intense episodes of burning sun exposure, as occur on foreign holidays, are currently believed to be important in the development of malignant melanoma.³

Department of Dermatology, University of Glasgow, Glasgow G12 8QQ
Rona M MacKie, professor of dermatology

West of Scotland Cancer Surveillance Unit, Ruchill Hospital, Glasgow
David J Hole, principal epidemiologist, Greater Glasgow Health Board

Correspondence to: Professor MacKie.

BMJ 1996;312:1125-8

Table 1—Age standardised incidence rates of melanoma by socioeconomic category for men and women; based on cases diagnosed 1979-93

Socioeconomic group:	Men		Women	
	No of cases (n=1134)	Rate (SE)	No of cases (n=2008)	Rate (SE)
1 (affluent)	115	9.1 (0.9)	210	16.1 (1.1)
2	141	7.2 (0.6)	266	12.6 (0.8)
3	256	6.5 (0.4)	418	9.8 (0.5)
4	238	5.5 (0.4)	441	9.5 (0.5)
5	209	5.3 (0.4)	332	7.9 (0.4)
6	134	4.2 (0.4)	244	7.0 (0.4)
7 (deprived)	40	2.4 (0.4)	90	5.0 (0.5)
Not known*	1		7	
Test for trend	P < 0.001		P < 0.001	

Population denominators for each postcode sector are available from the 1981 and 1991 national censuses. A 1986 population was estimated by averaging appropriate age groups from the 1981 and 1991 censuses, allowing for the 10 year aging between censuses. Incidence rates for 1979-83 used the 1981 population, for 1984-8 the 1986 population, and for 1989-93 the 1991 population. All incidence rates have been age standardised to the Scottish 1981 population.

*Lived in very small postcode areas or had imprecisely quoted addresses.

No analysis has been yet been made of melanoma survival by socioeconomic status with control for important predictors of prognosis such as age, sex, ulceration, and tumour thickness of the primary melanoma at the time of initial surgery.⁴ We carried out such an analysis, looking at all patients presenting with malignant melanoma over a 15 year period in the west of Scotland where there is available, as a result of the activities of the Scottish Melanoma Group, detailed information on clinical and pathological details of all patients presenting with cutaneous melanoma in this area (population 2 716 900). In addition, each patient's postcode enables an area based classification of their socioeconomic status to be made. Throughout the west of Scotland area, groups of very different socioeconomic status experience the same daily climatic conditions, living within 2-3 km of each other.

Methods

Details of melanoma patients were drawn from the records of the west of Scotland section of the Scottish Melanoma Group, which records details of all patients presenting with primary cutaneous malignant melanoma in Scotland.⁵ A total of 3142 patients (2008 women and 1134 men) diagnosed as having invasive primary cutaneous malignant melanoma (level 2 or deeper) between 1979 and 1993 were identified and were allocated an economic grouping by using the

method of Carstairs and Morris.⁶ This classifies areas according to four measures derived from census data: unemployment, car ownership, overcrowding, and percentages of households where the occupation of its head is classified as social class IV and V. Thus, given a postcode sector of residence, it is possible to allocate a person to one of seven categories from 1, the most affluent, to 7, the most deprived. All deaths up to December 1994 were recorded.

STATISTICAL METHODS

We examined trends in incidence across deprivation categories, comparing observed and expected numbers of cases and applying the χ^2 test for trend. Expected numbers of cases for each affluence category were calculated by applying the overall incidence rates for each age group, sex, and year of diagnosis to the appropriate age-sex distribution for each deprivation category for each year of the study; data were then aggregated. Trends in incidence over time were assessed by fitting linear regressions of age standardised rates on year of diagnosis for the affluence-sex subgroups. The gradients derived from the regression models represent average annual increases in rates and were tested statistically, assuming a *t* distribution.

Dates of death were obtained through contact with the patient's general practitioner and with the registrar general for Scotland through linkage with the west of Scotland cancer registry. Deaths due to melanoma were identified.

The trend in disease free survival across deprivation categories for the different age and sex groups was tested using the log rank statistic.⁷ Calculation of hazard ratios, to compare disease free survival between affluence categories while allowing for the major recognised prognostic factors (tumour thickness, ulceration, etc), was carried out using Cox's proportional hazards model.⁸ All variables included in the model were entered as categorical variables with stage at presentation with melanoma in six categories: primary melanoma <1.5 mm, 1.5-2.49 mm, 2.5-3.49 mm, 3.5-4.99 mm, >5 mm thick, and melanoma spread beyond the primary site.

Results

Table 1 presents the average annual age standardised incidence by deprivation category over the 15 year period 1979-93. For both sexes there was a strong ($P < 0.001$) trend in incidence ranging from 2.4/100 000 for least affluent men to 9.1/100 000 for most affluent men and from 5.0/100 000 for least affluent women to 16.1/100 000 for most affluent women. Figure 1 shows that for all six subsets—men and women in most affluent, intermediate, and least affluent circumstances—there was a steady increase in incidence over the 15 years studied. The slopes of the six curves are similar, implying that the factors responsible for the rising incidence are acting in all six groups.

Figure 1 and table 1 also show the clear difference in incidence of malignant melanoma between the sexes, with a female:male ratio of 3:2. This relative over-representation of women is maintained across the socioeconomic spectrum.

Table 2 lists the percentage of patients presenting with tumours less than 1.5 mm thick divided by socioeconomic status and over time. For men in the affluent and intermediate categories, there has been a significant increase in the proportion presenting with thin melanomas between 1979-84 and 1991-3. The highest proportion of thin melanomas in men was seen in the affluent group (61%) in the period 1985-90. For women, the overall trends are similar, and there have been significant increases over time in the proportion of thin tumours in the intermediate and least affluent

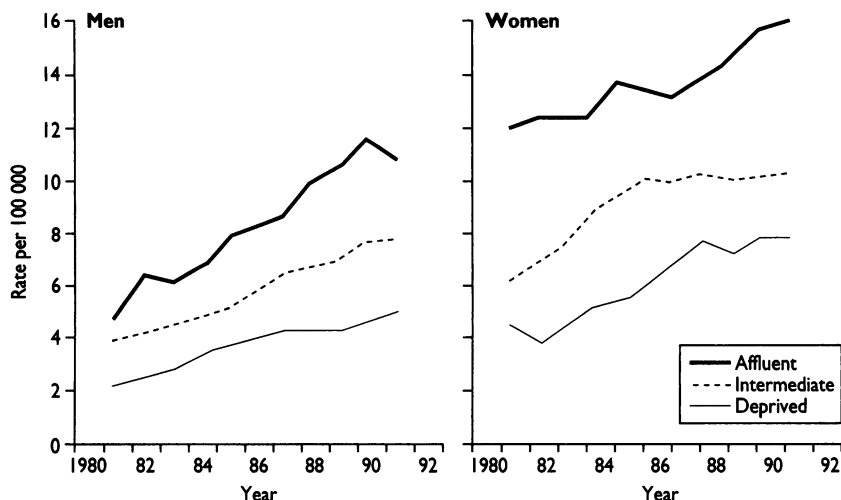


Fig 1—Incidence of melanoma (five year moving averages) in men and women in affluent, intermediate, and deprived categories, 1979-93

Table 2—Percentage (number) of patients presenting with primary melanoma <1.5 mm thick by socioeconomic category over time*

Time period	Socioeconomic category		
	Affluent	Intermediate	Deprived
Men:			
1979-84	34 (23/68)	29 (54/188)	26 (13/50)
1985-90	60 (73/121)	45 (138/310)	40 (32/81)
1991-93	54 (36/67)	54 (111/205)	40 (17/43)
P value†	<0.05	<0.001	>0.05
Women:			
1979-84	52 (80/154)	38 (136/358)	29 (28/96)
1985-90	61 (121/199)	57 (310/544)	51 (77/152)
1991-93	59 (72/123)	58 (167/289)	57 (49/86)
P value†	>0.05	<0.001	<0.01

*156 patients included above presented with stage 2 disease (involvement of regional lymph nodes with no detectable primary tumour).
†Change in proportion of small tumours over time for each sex/deprivation category.

groups of women. Affluent women had a significantly higher proportion of thin tumours than affluent men (52% v 34%) during the 1979-84 period. As with men, the highest proportion of thin tumours in women (61%) was seen in the most affluent group in the 1985-90 period. In all groups except the least affluent men, the proportion of primary melanomas <1.5 mm at excision is now over 54%.

Table 3 and figure 2 show the relation between five year survival and socioeconomic status. For the most affluent men, five year survival is 74% and for the most affluent women it is 84%, compared with 68% and 82% for the intermediate men and women and 61% and 79% for least affluent men and women. The difference in survival between the most affluent group and the other groups was significant (P for trend <0.001), as was the difference in survival between the most affluent men and women (P<0.001).

Table 3—Five year survival by socioeconomic category, sex, and age group

	Socioeconomic category			P value (test for trend)*
	Affluent	Intermediate	Deprived	
Sex				
Men	64 (256)	58 (703)	56 (174)	<0.001
Women	79 (476)	72 (1191)	70 (334)	<0.001
Age (years)				
<45	90 (222)	84 (520)	79 (128)	0.02
45-54	77 (150)	78 (320)	73 (73)	<0.01
55-64	83 (132)	70 (356)	67 (107)	<0.001
65-74	65 (111)	55 (373)	62 (102)	0.51
≥75	36 (117)	38 (325)	40 (98)	0.81

*Trend in survival rates across seven deprivation categories; age and sex adjusted where appropriate.

Table 4—Relative hazard ratios derived from Cox's proportional hazards model for survival differences between socioeconomic categories for melanoma patients under 65 years of age. Based on 450 deaths among 2008 patients

	Socioeconomic category			P value (test for trend)
	Affluent	Intermediate	Deprived	
Unadjusted	0.60 (0.46 to 0.77)***	1.00	1.34 (1.05 to 1.70)*	<0.001
Adjusted for age and sex	0.61 (0.47 to 0.78)***	1.00	1.38 (1.09 to 1.75)*	<0.001
Adjusted for age, sex and tumour thickness	0.75 (0.57 to 0.98)*	1.00	1.32 (1.04 to 1.68)*	<0.001
Adjusted for age, sex, tumour thickness and ulceration	0.76 (0.58 to 0.99)*	1.00	1.30 (1.02 to 1.64)*	<0.001

*P<0.05, ***P<0.001 for comparison with intermediate group.

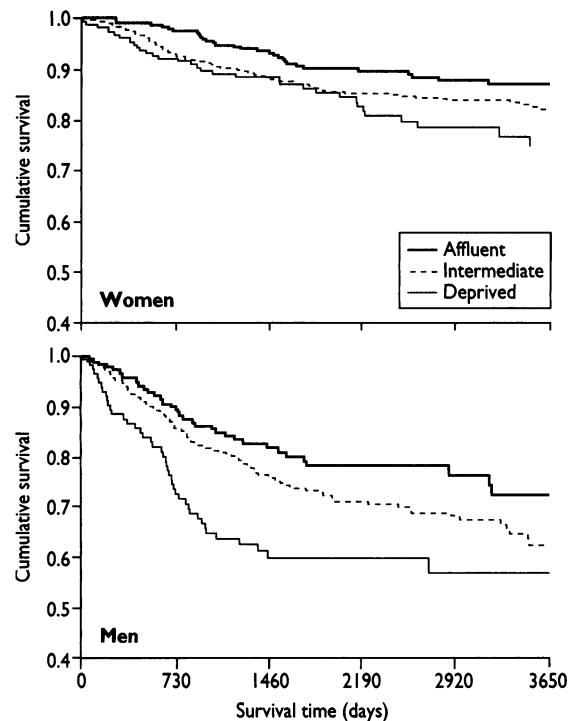


Fig 2—Kaplan-Meier survival curves for men and women in affluent, intermediate, and least affluent groups. Only deaths related to melanoma were included for analysis

Table 3 further divides five year survival by affluence and by age group. In the most affluent groups the five year survival for younger people was 91% while in the least affluent group it was 80% (P=0.04). In people aged 55-64, survival was better in the more affluent groups (P<0.001).

The relative hazard ratios for patients under 65 years presenting with melanoma are shown in table 4. After adjustment for all known prognostic factors such as age, sex, site, ulceration, and primary tumour thickness, the most affluent patients continued to have a survival advantage over the intermediate group, which in return had a better survival than the least affluent group (P for trend <0.001).

Discussion

This study is the first to show a relation between thickness at presentation and socioeconomic status, and also the first to relate socioeconomic status to survival. It shows that affluent individuals have more favourable five year survival prospects. Even after the major prognostic factors (tumour thickness, ulceration, age at diagnosis, and site of primary tumour) were controlled for, this difference remained significant. Tumour thickness is commonly regarded as a measure of early diagnosis; when patients were matched for tumour thickness, socioeconomic status continued to be a factor influencing survival. This suggests that a greater knowledge of the features of early melanoma

Key messages

- Affluent people have a significantly higher risk of developing malignant melanoma than deprived individuals
- Paradoxically, five year disease free survival prospects are better for affluent people
- Variations in incidence could be explained by differences in sun exposure related to socio-economic status
- Variations in nutrition and possibly immune function by socioeconomic status could explain survival differences
- In public education campaigns, deprived men should be targeted to encourage earlier diagnosis
- Primary prevention campaigns should be targeted to more affluent socioeconomic groups

in the affluent group, leading to earlier self referral and thus earlier diagnosis and treatment, does not wholly explain our results.

Recent results from the same geographic area also showed poorer survival for the less affluent patients with breast cancer.⁹ We speculate that a common factor, such as poor nutrition leading to low levels of antioxidants, or immunological defects, could be responsible in both tumour types.

The pattern of higher incidence but better survival for the most affluent compared with the least affluent of both sexes is mirrored in the greater incidence of melanoma but better survival in women. The greatest risk of developing melanoma is seen in affluent women, and the poorest five year disease free survival prospects in

deprived men. It is possible that behavioural differences with regard to sun exposure between the sexes and between affluent and deprived individuals may explain some of the differences in incidence, but this would not explain the differences in survival.

Table 2 shows clearly that over the three time periods studied the proportion of thin tumours has increased in all groups, and that this increase is significant in intermediate and deprived women and affluent and intermediate men. Over 50% of all tumours are now <1.5 mm thick, but among deprived men the proportion is only 39%, indicating that early detection activities need to be targeted particularly at less affluent men.

Funding: The Scottish Melanoma Group has been funded by the Cancer Research Campaign and is currently funded by the Scottish Home and Health Department.

Conflict of interest: None.

- 1 Lee JAH and Strickland D. Malignant melanoma, social status and outdoor work. *Br J Cancer* 1980;41:757-63.
- 2 Kirkpatrick CS, Lee JAH, White E. Melanoma risk by age and socioeconomic status. *Int J Cancer* 1990;46:1-4.
- 3 Nelemans PJ, Groenendal H, Kiemeneij L, Rampen F, Ruiter D, Verbeek A. Effect of intermittent exposure to sunlight on melanoma risk among indoor workers and sun sensitive individuals. *Environ Health Perspect* 1993;101:252-5.
- 4 Breslow A. Tumour thickness, level of invasion, and node dissection in stage I cutaneous melanoma. *Ann Surg* 1975;18: 572-5.
- 5 MacKie RM, Hunter JAA, Aitchison TC, Hole D, McLaren K, Rankin R, et al. Cutaneous malignant melanoma in Scotland 1979-1989. *Lancet* 1992;339:971-5.
- 6 Carstairs V, Morris R. *Deprivation and health in Scotland*. Aberdeen: Aberdeen University Press, 1991.
- 7 Peto R, Peto J. Asymptotically efficient rank invariant test procedures. *Journal of the Royal Statistical Society* 1972;135A:185-94.
- 8 Cox DR. Regression models and life tables. *Journal of the Royal Statistical Society* 1972;34:187-220.
- 9 Carnon AG, Ssemwogerere A, Lamont DW, Hole DJ, Mallon EA, George WD, et al. Relation between socioeconomic deprivation and pathological prognostic factors in women with breast cancer. *BMJ* 1994;309:1054-7.

(Accepted 24 January 1996)

Association between incidence of non-Hodgkin's lymphoma and solar ultraviolet radiation in England and Wales

Graham Bentham

Abstract

Objectives—To examine whether the incidence of non-Hodgkin's lymphoma in different areas of England and Wales is associated with levels of solar ultraviolet radiation.

Design—Geographically based study examining the association between incidence of non-Hodgkin's lymphoma and estimated levels of solar ultraviolet radiation, controlling for social class and employment in agriculture.

Setting—59 counties in England and Wales.

Subjects—All registered cases of non-Hodgkin's lymphoma during the period 1968-85.

Main outcome measure—Age and sex adjusted odds ratio for non-Hodgkin's lymphoma in each county.

Results—Incidence of non-Hodgkin's lymphoma was significantly associated with solar ultraviolet radiation levels ($P < 0.001$), even after social class and employment in agriculture were controlled for ($P = 0.004$). In a comparison of counties in the highest and lowest quarters of solar ultraviolet radiation, the relative risk of non-Hodgkin's lymphoma was 1.27 (95% confidence interval 1.24 to 1.29), rising to 1.34 (1.32 to 1.37) after adjustment for social class and employment in agriculture.

Conclusions—The incidence of non-Hodgkin's lymphoma in different areas of England and Wales is positively associated with levels of solar

ultraviolet radiation. These results are consistent with the hypothesis that exposure to solar ultraviolet radiation increases the risk of non-Hodgkin's lymphoma.

Introduction

The incidence of non-Hodgkin's lymphoma has increased greatly in many countries,¹ including England and Wales.² An established risk factor is chronic immunosuppression³⁻⁴—as in individuals with congenital immunodeficiency syndromes,⁵ patients treated with immunosuppressive drugs,⁶ and those infected with HIV.⁷ Particular types of non-Hodgkin's lymphoma are also associated with infection with the Epstein-Barr virus and with HTLV-I.⁸ Risk is also raised in a number of occupations,^{9,10} notably in agriculture, where exposure to pesticides may be a factor.¹¹ However, trends in known risk factors and in diagnostic practices can account for only a small part of the observed increase in non-Hodgkin's lymphoma.^{12,13} Part of the rise may have been the result of increased population exposure to some other widespread risk factor.

It has been hypothesised that exposure to solar ultraviolet radiation may be a cause of non-Hodgkin's lymphoma.¹⁴ This is an important cause of skin cancers,¹⁵ which have been increasing even more rapidly than non-Hodgkin's lymphoma,¹⁶ probably as a result of the adoption of lifestyles involving increased exposure to the sun.¹⁵ Increased exposure to sunlight might also

Centre for Social and Economic Research on the Global Environment, School of Environmental Sciences, University of East Anglia, Norwich NR4 7TJ
Graham Bentham, senior lecturer

Correspondence to: c.bentham@uea.ac.uk

BMJ 1996;312:1128-31