Our study has several limitations. Answering questions about a sensitive topic on the telephone can be difficult, a questionnaire with a fixed choice of answers prevented doctors from qualifying or justifying their responses, and we lacked detailed information about doctors who did not respond.

Previous studies found similar patterns, but the French counterparts to Italian general practitioners and US oncologists were more in favour of legalising euthanasia.^{3 4} Our findings contradict the argument that opinions on euthanasia are related to cultural differences in English speaking countries; comparative studies are needed.² In France, the support shown for euthanasia may be due to a lack of professional knowledge on palliative care.⁵ Improving such knowledge would improve end of life care and may also clarify the debate over euthanasia.

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Case fatality rates for meningococcal disease in an English population, 1963-98: database study

Michael J Goldacre, Stephen E Roberts, David Yeates

Unit of Health-Care Epidemiology, Department of Public Health, University of Oxford, Oxford OX3 7LF Michael J Goldacre professor of public health Stephen E Roberts statistician David Yeates computer scientist

Correspondence to: M Goldacre michael.goldacre@ dphpc.ox.ac.uk

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Meningococcal septicaemia and meningitis are common causes of death in children and young adults. In fatal cases, the time from onset to death is often short. We analysed case fatality rates for meningococcal disease between 1963 and 1998 to determine whether they have decreased.

Methods and results

We used data on meningococcal disease from the Oxford record linkage study database, which includes anonymised statistical abstracts of records of admission to hospital and death certificates in a defined population of 0.35 million people from 1963, 0.9 million from 1966, 1.9 million from 1974, and 2.5 million from 1987 to 1998. We calculated incidence of menin-

gococcal disease and case fatality rates and assessed the significance of trends over time with logistic regression.

From 1963 to 1998, 1223 people had a record of admission to hospital for meningococcal meningitis or meningococcal septicaemia and 25 for other diagnoses—mainly meningitis or septicaemia without specification of an organism and a death certificate that specified meningococcal disease. The median age of these 1248 was 6 years; 255 (20%) were < 1 year old, 422 (34%) were 1-9 years old, 279 (22%) were 10-19 years old, 290 (23%) were ≥ 20 years old, and the ages of two were unknown; 116 died within 30 days and five more within 365 days of admission.

Analysis of the database showed that a further 25 people had died from meningococcal disease. Fifteen

Cases of meningococcal disease* from the Oxford record linkage study database, number of deaths within 30 days and case fatality rate per 100 000, 1963-98

	Cases admitted to hospital			All cases			
Time period	No of cases	No of deaths	Case fatality rate (95% CI)	No of cases	Incidence per 100 000 population (95% CI)	No of deaths†	Case fatality rate (95% CI)
1963-8	41	5	12.2 (2.2 to 22.2)	42	1.23 (0.86 to 1.60)	6	14.3 (3.7 to 24.9)
1969-73	72	6	8.3 (1.9 to 14.7)	74	1.72 (1.33 to 2.11)	8	10.8 (3.7 to 17.9)
1974-8	108	12	11.1 (5.2 to 17.0)	111	1.24 (1.01 to 1.47)	15	13.5 (7.2 to 19.9)
1979-83	113	11	9.7 (4.3 to 15.2)	113	1.09 (0.89 to 1.29)	11	9.7 (4.3 to 15.2)
1984-8	147	16	10.9 (5.8 to 15.9)	152	1.33 (1.12 to 1.54)	21	13.8 (8.3 to 19.3)
1989-93	303	25	8.3 (5.2 to 11.3)	308	2.40 (2.13 to 2.67)	30	9.7 (6.4 to 13.1)
1994-8	464	41	8.8 (6.3 to 11.4)	473	3.55 (3.23 to 3.87)	50	10.6 (7.8 to 13.3)
1963-98	1248	116	9.3 (7.7 to 10.9)	1273	1.97 (1.86 to 2.07)	141	11.1 (9.4 to 12.8)
P value for trend‡			0.31		<0.001		0.28



†Includes 25 people who died outside hospital or who had no record of hospital admission.

‡P values refer to the change in annual rates 1963-98 from logistic regression modelling. Visual inspection of the annual incidence and further modelling showed little change in incidence during 1963-88 (P=0.10) and a significant increase during 1989-98 (P<0.001).</p>

P+

Tables for specific age groups are on bmj.com died in a hospital in the region but had no record of admission; they had probably been brought in dead or died soon after arrival at hospital without a formal admission. Ten had died at residential addresses. The median age of the 25 was 16 years, and 11 (44%) were aged at least 20 years. Of all 146 people who died, 25 (17%) died without admission to hospital, 41 (28%) died on the day of admission, and 40 (27%) died the day after admission.

Incidence has risen substantially in recent years (table). Case fatality rates have not significantly declined over time (P=0.31 for cases admitted to hospital, and P=0.28 for all cases). We found no significant trends in case fatality rates within individual age groups (see tables on bmj.com). For those admitted to hospital, case fatality rates at 30 days were similar in the teaching hospital (8.9%; 95% confidence interval 5.6% to 12.2%) and non-teaching hospitals (9.4%; 7.6% to 11.3%), and in males and females.

Comment

Case fatality rates for meningococcal disease in the Oxford region have not fallen since the late 1960s. Others have also reported no recent reduction in case fatality rates in population based studies,¹⁻³ although recent declines in a specialist paediatric unit have been described.⁴ In our study, a relatively high percentage of people who died outside hospital, or on arrival, were adults and perhaps had been less closely observed than children in their illness outside hospital. Our case fatality rate of 11% is the same as that found in a study based on multiple source case ascertainment and case

note review in an adjacent health region in 1969-74.⁴ In that study, 22% of all deaths, compared to 17% in this study, occurred without admission to hospital or in people who were dead on arrival.⁵

The time from onset to death is usually rapid. This, and the fact that death rates have not declined, emphasises the need for vigilance in making the diagnosis and the importance of prevention through immunisation and, when appropriate, chemoprophylaxis for contacts.

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Prophylaxis for venous thromboembolism during treatment for cancer: questionnaire survey

C C Kirwan, E Nath, G J Byrne, C N McCollum

Venous thromboembolism is common in patients with cancer and is often the cause of death.¹ Patients receiving treatment for cancer are at even greater risk of thromboembolism. Thromboembolism occurs in 5% of patients receiving chemotherapy for early breast carcinoma,² and up to 17.6% of patients receiving chemotherapy for metastatic breast disease are affected.³ Patients with node-negative breast cancer taking tamoxifen were six times more likely to develop venous thromboembolism.⁴

Adjuvant use of tamoxifen carries a relative risk of 1.22 compared with no treatment. Combining methods of treatment further increases the risk of thromboembolism. Chemotherapy with tamoxifen increases risk by 3.5 times compared with chemotherapy alone,⁵ and preoperative radiotherapy for rectal carcinoma doubles the postoperative risk of venous thrombosis.⁵ Low doses (1 mg) of warfarin throughout chemotherapy for metastatic breast cancer are associated with a relative risk reduction of 85% with no increase in serious bleeding complications.¹

Participants, methods, and results

We sent a postal questionnaire to all oncologists in northern England, identified by internet search and in the *Medical Directory 2002*. We used a scoring system to establish specialty, main type of cancer treated, main method of treatment (chemotherapy, hormone therapy, or radiotherapy), and current prophylaxis practice and estimate of risk of venous thrombembolism.

Of the 123 responses to the 166 questionnaires we sent, 106 (64%) were acceptably completed. Half the oncologists (56) specialised in clinical oncology, 31 in medical oncology, seven in surgery, five in gynaecological oncology, five in paediatrics, one in urology, and one in radiology. We have no information about the specialties of oncologists who did not respond. The most common treatment was chemotherapy, used by 41 (39%) oncologists; 10 (9%) used hormone therapy and 44 (42%) used radiotherapy. The oncologists treated many types of tumour. Education and Research Building, Wythenshawe Hospital, Manchester M23 9LT C C Kirwan surgical research fellow E Nath research assistant G J Byrne senior lecturer in surgical oncology C N McCollum professor of surgery Correspondence to: G I Byrne gedbyrne@

compuserve.com

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