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How is place of death from cancer changing and what affects it? Analysis of cancer registration and service data

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We aimed to compare trends in place of cancer death with the growth of palliative care and nursing home services, and investigate demographic, disease-related and area influences on individual place of death, using registration data for 216404 patients with breast, lung, colorectal and prostate cancer and aggregate data on services in South East England. Between 1985 and 1994 there was a trend away from hospital death (67-44%), to home (17-30%) and hospice death (8-20%). After 1995, this partly reversed. By 2002, hospital death rose to 47%, home death dropped to 23%, hospice death remained stable and nursing home death rose from 3 to 8%. Numbers of palliative care services increased, but trends for hospice and nursing home deaths most clearly followed the beds available. Cancer diagnosis and treatment influenced individual place of death, but between 1998 and 2002, age and area of residence were associated with most variation. Older patients and those living in more deprived areas died more often in hospitals and less often at home. Despite more palliative care services the proportion of people dying at home has not increased. Variation by age, deprivation and area of residence is unlikely to reflect patient preference. More active surveillance and planning must support policies for choice in end of life care.

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Most people say they would prefer to die at home, but in reality most patients in the UK spend their final days in hospital (WHO, 2004a). Over the last 30 years, hospices, palliative care teams and units have developed with the aim of improving care towards the end of life, and allowing people to die where they wish, if this is possible. Services have initially focused on patients with cancer, primarily because of the relative ease of predicting the course of this disease, and a national policy for supportive and palliative cancer care is now in place (House of Commons Health Committee, 2004; NICE, 2004a, b). There is, therefore, increasing interest in Europe about whether data on place of death can be used as an interim measure of the success of services provided (WHO, 2004a). Figures for England and Wales revealed a trend away from death in hospital or nursing homes to hospices between 1985 and 1994, but very little change in home deaths, which remained around 26%. However, this proportion varied between regions and was lowest in South East England across all age and cancer types (Higginson et al, 1998). For common cancers, individual, disease-related and area of residence factors were consistently associated with, but not strongly predictive of place of death. Men, patients aged under 74 years, those with lung or colorectal cancer or living in more affluent areas were more likely to die at home than women, patients aged over 75 years, those with breast cancer or those living in less affluent areas (Higginson et al, 1998, 1999).

The Thames Cancer Registry covers a population of 14 million people in South East England, an area with one of the highest concentrations of hospice and palliative care services in the UK (Hospice Information, 2006). We used Registry data to describe trends in place of death for common cancers and compared these to the growth of palliative care services and nursing homes between 1985 and 2002. We then investigated the relationship between demographic, disease-related factors and individual place of death throughout the period, and the additional influence of area of residence between 1998 and 2002.

METHODS

In the UK cancer registries record the occurrence of cancer in their residential populations as well as treatments given in the first 6 months after diagnosis. Information about death is provided by the National Health Service Central Register through the Office for National Statistics. Death certificates routinely record place of death and assign cancer as a main or contributing cause of death in part I of the certificate.

We extracted data on 216 404 residents in South East England who had been diagnosed with breast, lung, colorectal and prostate cancer between 1985 and 2002, and who died from their disease between 1985 and 2002. Cases where the only registration information was from the death certificate were not included. From death certificates, we classified death as occurring in NHS acute hospitals, hospices, long stay hospitals or nursing homes, private hospitals, at home or as unknown. We could identify nursing homes by their address, but death certificates do not

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distinguish deaths in hospital palliative care units from those in other wards.

We extracted data on hospice and palliative care services from Hospice Information directories for 1985-2002 (Hospice Information, 2006) and calculated the number of hospice beds, home care teams, day care services, hospital palliative care or support teams and hospital support nurses in our area. We summed home care services regardless of their funding (independent, NHS and Macmillan Cancer Relief) or base (hospices, NHS hospital or community), although there was insufficient detail about Marie Curie home services to include these. We could not deduce team size or caseload. We also obtained aggregate data on numbers of beds in registered nursing homes from the Department of Health where this was available for 1991-2001. We first plotted the proportion of deaths occurring in each of home, hospital, hospice and nursing home against the growth of different services over time. Data on acute and general hospital beds in our area were available only between 1996 and 2002 and were not plotted.

We then took death in hospice, nursing home, NHS acute hospital and at home as our four dependent variables and fitted logistic regression models to identify individual demographic, disease-related and area of residence factors predicting in turn each of these outcomes versus the others. Our first analysis for the entire period 1985-2002 included sex, age at diagnosis, whether the diagnosis was based on clinical or microscopic evidence, primary site of cancer and treatment with surgery, radiotherapy, chemotherapy or hormone therapy. We also adjusted for year of death and years since diagnosis to examine trends over time. We grouped age into four bands: <65 years, 65-74 years, 75-84 years and 85 years plus. Our second analysis explored the additional influence of area of residence for the years 1998-2002. For this, we assigned each individual to an electoral ward and a cancer network using their postcode of residence. We calculated the deprivation score for each ward using the income domain of the Indices of Multiple Deprivation (IMD) 2000 for England (Department of Environment, 2000) and assigned individuals to a quintile of deprivation ranging from most (1) to least affluent (5) wards.

We present the results of logistic regression analyses as proportions of deaths occurring in each place for each factor. Proportions are easier to interpret than odds ratios, and were derived from a back calculation from the odds ratios obtained from the logistic regression analyses. We present unadjusted and adjusted proportions to show the effect of controlling for all other factors. Our large sample size means that many small differences reach statistical significance. We draw attention only to those factors producing at least five percentage points difference - a difference which we believe a clinical service might be interested to explore further.

RESULTS

The average age of death for patients in this cohort increased from 71.3 years in 1985 to 72.7 in 2002. The proportion dying at age of 85 years and over increased from 8 to 12% while the proportion dying between the age of 65 and 74 years dropped from 34 to 28%.

How has place of death changed?

Figure 1 suggests that the period 1985-2002 is best considered in two phases - before and after 1994. In the first phase, hospital deaths declined from 67 to 44% - a trend that appeared to be mirrored by a combined increase in home death from 17 to 30% and in hospice death from 8 to 20%. In the second phase, however, the movement away from hospital death appeared to partly reverse. Between 1995 and 2002 hospital death rose to 47%, nursing home death to 8%, hospice death remained stable and home deaths dropped to 23%. In 2002 - the last year of the study -



Figure I Trends for place of death for patients with breast, colorectal, lung and prostate cancer in South East England 1985-2002. Note: Figure excludes the proportion dying in private hospitals and patients where place of death was not known.

home death and nursing home death home appear to have increased slightly and hospital death to have decreased. During 1992 and 1995 there were changes in processing and receipt of our registry data which may be responsible for the 'mirroring' of trends in hospital and home deaths during this period. This artefact overlies but does not explain the reversal of overall trends which is also seen in national data for this period.

Figure 2 shows that during the first phase, while home and hospice death increased, the provision of home care services and hospice beds also increased. From 1995 onwards while nursing home death and hospital death increased, nursing home beds and, to a lesser degree, the sum of hospital palliative care services (teams and nurses) also increased. For nursing home deaths, unlike hospice deaths, there is a lag of several years between the rise of available beds in these services and deaths within them. The decline in home death occurring after 1995 did not appear to follow a substantial drop in the provision of palliative home care or day care services, which both remained stable, although during this period the availability of nursing home beds was increasing.

Which individual and disease-related factors affect place of death?

Table 1 shows unadjusted and adjusted proportions of deaths in each place for individual demographic and disease-related factor over the entire study period. Hospital death was more likely for patients aged over 75, those with lung or breast cancer, a clinical rather than microscopic diagnosis, and those not receiving radiotherapy. Home death was more likely for those with colorectal cancer and those aged less than 75 years. Hospice death was also more likely for colorectal cancer and for those aged less than 75 years. Nursing home death increased with older age (4% for those aged 65-74 years and 12% of those aged over 85 years).

Did place of residence affect place of death between 1998 and 2002?

Our analysis for the most recent years included area of residence as assessed by cancer network of residence and deprivation of ward of residence (Table 2). The results for demographic and diseaserelated factors were broadly similar to those in Table 1, although nursing home deaths become more likely for those with breast and prostate cancer. However, much more striking was the variation by



Figure 2 Trends in place of death for patients with breast, lung, colorectal and prostate cancer and the growth of services for care towards the end of life in South East England 1985–2002. Note: Department of Health Data is only available for 1991–2001 and Hospice Information Directories are not available for 1986 and 1989.

area of residence. Concentrating on the nine of 13 cancer networks that we completely cover, the adjusted proportion of patients dying in hospital ranged from 39% in Sussex to 60% in West London. Home deaths ranged from 16% in Surrey, West Sussex and Hampshire to 27% in South Essex. Hospice death ranged from 10% in West London to 31% in Surrey, West Sussex and Hampshire. Nursing home deaths ranged from 4% in North London to 13% in Sussex. Of London networks, South East London had the lowest rate of hospital death (49%) and the highest rate of home death (23%). Patients from more deprived areas died more often in hospital and less often at home. There was no important deprivation gradient for nursing home or hospice death.

DISCUSSION

Summary of main findings

This study of 216404 patients diagnosed and dying from four common cancers in South East England found an initial trend

away from hospital death (67-44%) to home (17-30%) and hospice death (8-20%) between 1985 and 1994. After 1995 this trend partly reversed. By 2002, the proportion of hospital deaths rose to 47%, hospice deaths remained stable, home deaths dropped to 23% and nursing home deaths rose from 3 to 8%. The number and range of palliative care services increased but trends for hospice and nursing home death most clearly followed the numbers of beds available. Analysis of individual data showed that throughout the period disease-related factors had a modest influence on place of death. Patients with colorectal cancer were more likely to die at home and in hospices while patients with lung or breast cancer, no microscopic diagnosis and no radiotherapy were more likely to die in hospitals. However, between 1998 and 2002, age and place of residence were associated with most variation. Older patients were more likely to die in hospitals and nursing homes and less likely to die at home or in hospices. Patients from deprived areas were more likely to die in hospitals and less likely to die at home. There was significant variation in each place of death by cancer network of residence.



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Table I	Associations of individual demographic and disease-related factors with place of death for cancer patients who died 1985–2002 in South East
England	

	A	cute hospi	tal		Home Hospice Nursing hor				ne				
	Number	%	% Adj	Number	%	% Adj	Number	%	% Adj	Number	%	% Adj	Total
Age group <65 65-74 75-84 85+	28 818 34 777 33 811 10 512	47 49 52 55	47 47 48 50	17332 18104 13960 3717	28 26 22 19	28 26 23 22	2 032 3 045 0 566 2046	20 18 16 11	20 20 18 12	956 2467 4721 2375	2 3 7 12	2 4 8 12	61 670 70 665 64 921 19 148
Test for Trend	χ^2 (I df)	574.1 <0.001	51.1 <0.001		1036. 6 <0.001	410.7 <0.001		753.7 <0.001	271.2 <0.001		4322.2 <0.001	3206.3 <0.001	
Sex Male Female	60 626 47 292	51 48	51 49	29 552 23 56 I	25 24	25 23	19 793 17 896	7 8	17 19	5005 5514	4 6	4 6	8 630 97 774
Test for Heterogeneity	χ^2 (I df) P	60.5 < 0.00	46.7 <0.001		9.2 <0.00	42.8 <0.001		97.6 <0.001	92.0 <0.001		232.3 <0.001	39.3 < 0.00	
Basis of diagnosis Clinical Microscopic	29 382 78 536	57 48	57 52	10763 42350	21 26	21 23	6697 30 992	3 9	3 6	3283 7236	6 4	6 5	51936 164468
Test for Heterogeneity	χ^2 (I df) P	222.9 <0.00	219.3 <0.001		536.2 <0.001	28.9 <0.00		959.3 <0.001	212.1 <0.001		311.8 <0.001	55.1 < 0.001	
Site Colorectal Lung Breast Prostate	23 203 54 794 17 375 12 546	46 54 47 48	46 54 56 52	3 38 24 738 9 40 5854	26 24 24 22	26 23 21 21	9873 16166 6904 4746	19 16 18 18	19 16 14 18	2705 3575 2304 1935	5 4 6 7	5 4 5 6	50 937 102 071 37 340 26 056
Test for Heterogeneity	χ ² (3 df) P	68. <0.00	837.7 <0.001		47.8 <0.00	320.6 <0.001		354.4 <0.001	294.2 <0.001		910.8 <0.001	5.9 <0.00	
Had noninvestigat No Yes	ive surgery 78 702 29 216	51 46	51 54	36 944 16 169	24 26	24 22	25 189 12 500	16 20	6 7	7527 2992	5 5	5 5	53 570 62 834
Test for Heterogeneity	χ^2 (I df) P	402.2 <0.001	09.0 <0.00		67.6 <0.001	61.6 <0.001		376.8 <0.001	0.3 0.618		1.9 0.171	1.1 0.298	
Had radiotherapy No Yes	79 090 28 828	52 44	52 43	34962 18151	23 28	23 27	24 472 13 217	16 20	16 20	8182 2337	5 4	5 5	5 223 65 8
Test for Heterogeneity	χ^2 (I df) P	83.5 <0.00	240.9 <0.00		548.2 <0.001	329.8 <0.001		528.9 <0.001	425.9 <0.001		323.6 <0.001	2.5 0.117	
Had chemotherap No Yes	94814 13104	50 46	50 49	45 22 I 7892	24 28	24 25	31 764 5 925	17 21	7 8	9976 543	5 2	5 3	187856 28548
Test for Heterogeneity	χ^2 (1 df) P	206.7 <0.001	2.7 <0.00		70.4 <0.00	20.9 <0.001		253.8 <0.001	0.6 0.00		568.8 <0.001	25.6 <0.00	
Had hormone the No Yes	rapy 85818 22100	50 48	50 49	41 640 11 473	24 25	24 25	29 243 8446	17 18	17 18	7631 2888	4 6	4 5	70 033 46 37
Test for Heterogeneity	χ^2 (I df) P	5.2 <0.00	23.8 <0.001		1.3 0.263	8.8 0.003		26.1 <0.001	9.1 0.003		236.5 <0.001	46.6 < 0.001	
Year of death 1985 1986 1987 1988 1989 1990 1991	3278 5473 5487 5976 6485 6558 6392	67 62 57 54 53 52 50	67 64 59 58 57 56	847 1792 2294 2660 3212 3415 3506	17 20 24 24 26 27 28	17 19 22 22 24 25 26	388 996 1183 1592 1718 1926 2074	8 11 12 14 14 15 16	8 10 11 12 12 13 14	191 328 330 396 425 420 453	4 4 3 4 3 3 4	4 3 3 3 3 3	4866 8833 9644 10989 12215 12659 12738

Table I (Continued)

	Acute hospital				Home			Hospice		N			
	Number	%	% Adj	Number	%	% Adj	Number	%	% Adj	Number	%	% Adj	Total
1992	6274	50	55	3318	26	24	2296	18	16	440	3	3	12660
1993	6536	45	50	4345	30	28	2834	19	17	507	3	3	14681
1994	5342	44	50	3730	30	28	2505	20	17	339	3	2	12279
1995	6055	47	54	3170	25	22	2242	17	15	398	3	2	12827
1996	4917	46	53	2625	25	22	2135	20	17	583	5	5	10704
1997	5606	47	54	2765	23	21	2188	18	15	631	5	4	11930
1998	6463	48	55	3062	23	20	2709	20	17	927	7	6	13501
1999	6653	48	55	3247	24	21	2675	19	17	920	7	5	13798
2000	6849	49	56	3008	22	20	2685	19	17	1019	7	6	13893
2001	6812	49	55	2892	21	19	2737	20	17	1019	7	6	13919
2002	6762	47	54	3225	23	21	2806	20	17	1193	8	7	14268
Test for Trend	χ^2 (I df) P	906.0 <0.001	353.0 <0.001		68.3 < 0.001	9.2 <0.00		949.9 <0.001	672.1 <0.001		1087.3 <0.001	648.8 <0.001	

Adjusted model includes: age, sex, basis of diagnosis, site, treatment (surgery, radiotherapy, chemotherapy or hormone therapy), year of death and years since diagnosis.

Table 2	Associations of individual	demographic,	disease-related	and area	of residence	with place of	of death	for patients	who d	ied from	Breast,	lung,
colorectal	or prostate cancer betwee	n 1998 and 20	02 in South Eas	st England								

	Αсι		Home			Hospice		Νι					
	Number	%	% Adj	Number	%	% Adj	Number	%	% Adj	Number	· %	% Adj	Total
Age group <65 65-74 75-84 85+	8756 10018 10790 3975	45 48 50 54	45 46 48 50	5085 4963 4180 1206	26 24 20 16	26 24 21 18	4578 4454 3727 853	23 21 17 11	23 22 19 13	411 1146 2263 1258	2 5 11 17	2 5 10 14	19521 21058 21377 7423
Test for Trend	χ^2 (I df) P	219.0 <0.001	55.6 <0.001		421.17 <0.001	2 3. <0.00		553.8 <0.001	268.8 < 0.001		2016.8 <0.001	293. <0.00	
Sex Male Female	18074 15465	49 47	49 47	8369 7065	23 22	23 22	7068 6544	19 20	19 20	2305 2773	6 8	6 10	36 629 32 750
Test for Heterogeneity	χ^2 (I df) P	31.2 <0.001	24.0 <0.001		6.3 <0.00	10.0 0.002		5.2 0.023	5.7 0.017	7	9.7 <0.00	22.8 < 0.00	
Basis of diagnosis Clinical Microscopic	8462 25 077	56 46	56 49	2755 12679	18 23	18 21	1977 11635	13 21	3 8	1472 3606	10 7	10 9	15 002 54 377
Test for Heterogeneity	χ^2 (1 df) P	495.0 <0.001	92.4 <0.00		65.9 <0.00	56.3 < 0.001		493.1 <0.001	59.9 <0.00		73.2 <0.00	6.5 0.01	
Site Colorectal Lung Breast Prostate	7528 15992 5853 4166	45 53 45 45	45 54 51 44	3799 6833 2821 1981	23 23 22 21	23 22 19 20	3564 5429 2722 1897	21 18 21 20	21 18 18 22	370 460 184 064	8 5 9 11	8 5 10 12	16714 30306 13026 9333
Test for Heterogeneity	χ² (3 df) P	421.7 <0.001	326.2 <0.001		2. 0.007	50.3 < < 0.001		02.9 <0.00	06.7 <0.00		565.1 <0.001	315.6 <0.00	
Had noninvestigative surgery No Yes	24 850 8689	50 44	50 50	10836 4598	22 23	22 22	9110 4502	18 23	18 20	3646 1432	7 7	7 7	49 630 19 749
Test for Heterogeneity	χ^2 (I df) P	208.3 <0.001	1.1 0.300)	7. <0.00	0.1 0.792		76.0 <0.00	2. 0.00		0.2 0.663	0.9 0.34	
Had radiotherapy No Yes	25 03 I 8508	51 42	51 41	10122 5312	21 26	21 25	8981 4631	18 23	18 23	3937 1141	8 6	8 8	49 42 20 237

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Table 2 (Continued)

	Acute hospital			Home				Hospice		Nu	_		
	Number	%	% Adj	Number	%	% Adj	Number	%	% Adj	Number	%	% Adj	Total
Test for Heterogeneity	χ² (Ι df) Ρ	452.2 <0.001	525.8 <0.001		263.5 < 0.001	43. <0.00		92.3 <0.00	62.9 <0.00		7.8 <0.00	2.4 0.122	
Had chemotherapy													
No Yes	27 748 579 I	49 44	49 47	11977 3457	21 26	21 23	10 409 3203	19 24	19 20	4764 314	8 2	8 5	56216 13163
Test for Heterogeneity	χ^2 (I df) P	22.7 <0.00	25.4 <0.001		50.9 <0.00	25.9 <0.001		227.3 <0.001	24.1 <0.001		508.2 <0.001	97.0 <0.001	
Had hormone therapy	27720	50	50	12 220	22	22	10.000	10	10	2/55	-	7	
No Yes	27739 5800	50 43	50 46	12320 3114	22 23	22 25	10.839 2773	19 21	19 20	3655 1423	/	/ 7	55958 13421
Test for Heterogeneity	χ² (Ι df) Ρ	74.6 <0.00	28.8 <0.001	8.8 21.9 0.003 <0.001			.5 0.00	2.3 0.130	1	259.4 <0.001	8.9 0.003	1	
Network of residence													
North East London	3946	56	56	1353	19	19	1346	19	19	347	5	5	7080
North London	3314	54	55	1224	20	19	11/3	19	20	261	4	4	6128
South East London	2222	47 49	47 51	167/	19	17	1347	22	22	313	5	5	7204 5884
West London	3988	58	60	1391	20	19	693	10	10	417	6	6	6879
TCB part of Mount Vernon	2459	50	53	1415	20	26	488	10	10	458	9	9	4924
South Essex	2121	52	54	1165	29	20	415	10	ii ii	338	8	8	4067
Kent & Medway	3663	42	42	2139	24	23	2186	25	26	688	8	7	8793
Surrey, West Sussex and Hampshir	re 2688	43	48	1232	20	16	1771	28	31	349	6	5	6262
Sussex	2642	38	39	1515	22	20	1625	23	25	1046	15	13	7001
TCR part of Central South Coast	327	25	25	249	19	18	367	28	32	150	12	10	1301
TCR part of Mid Anglia	1734	52	53	791	24	22	473	14	15	328	10	9	3360
TCR part of West Anglia	201	41	45	140	28	24	48	10	11	48	10	9	492
Other/NK	24	46	55	12	23	21	6	12	6	7	13	12	52
Test for	χ² (12 df)	1361.2	1219.7		399.9	401.8		1613.5	1687.6		943.1	690.8	
Heterogeneity	Γ	< 0.001	< 0.001		< 0.001	< 0.001		< 0.001	< 0.001		< 0.001	< 0.001	
IMD	5000	10	10				0.450						
Most affluent I	5020	42	42	2926	25	25	2458	21	21	896	8	8	11850
2	5965	46	45	3106	24	23	2449	19	21	1125	9	8	130/1
3	6/54 7/02	48 51	47	31//	23	22	2620	19	22	1099	8	7	13736
T Least affluent 5	8105	53	49	20222	21 19	18	2071	17 21	24	834	5	7	15 330
NK	3	27	23	2700	18	17	4	36	72		9	5	15 550
Test for	χ^2 (I df)	362.7	92.2		132.4	152.0		0.8	58.9		67.5	4.5	
Trend	Р	< 0.00	< 0.00		< 0.00	< 0.00		0.387	< 0.001		< 0.00	0.035	

Adjusted model includes: age, sex, basis of diagnosis, site, treatment (surgery, radiotherapy, chemotherapy or hormone therapy), cancer network of residence and deprivation.

Limitations of this study

This population-based study used data collected from medical records and death certificates for routine cancer registration. Coding officers may have missed some deaths in new nursing homes and hospices when their addresses were unfamiliar in the early part of the study period. Lack of information on death certificate on deaths in hospital palliative care units and lack of data on hospital beds meant we could not explore these trends and it is possible that excluding patients for whom we had only death certificate data from the analysis introduced some bias. Important information on patient preference for place of death, functional status, presence of a carer at home, family support, and hospital and community services received in the weeks before death (Grundy *et al*, 2004; Gomes and Higginson, 2006) is not routinely collected and is therefore missing from the individual analyses.

Comparison to other findings

No other large UK studies have compared overall trends in place of death with the growth of services that might support patients to die in different places. However, one study of North West England between 1993 and 2000 found that proximity to a hospice or hospital increased the chances of dying there (Gatrell *et al*, 2003). Studies in the US have also found that the availability of beds and physicians affects death in hospital (Fisher *et al*, 2003a, b). National bed data available for 1987 to 1994 when hospital deaths decreased showed a decline of 19% in the numbers of acute and general hospital beds (Department of Health, 2006). It, therefore, seems likely that the initial trend for increasing home death was in part driven by the decreased availability of hospital beds and the growth of hospice and palliative home care services. However, it is more difficult to explain the reversal of the trend for home death

after 1995 using the routine service data that is available. The number of home care teams did not decline, hospital palliative care services were only just beginning to increase and hospital beds did not increase nationally until 2001. We can speculate that the decline in home death was due to other changes in care at home including the ability of families to provide care, the prior move of some older adults into nursing homes and the move to GP cooperatives for out of hours care. These factors could all have led to increased hospital admission and fewer home deaths.

Turning to predictors of individual place of death, our finding that younger patients, patients with colorectal cancer and those living in more affluent areas died more often at home is consistent with (Higginson et al 1998, 1999) analyses of a partial national registration dataset up until 1994. However, we found that patients with breast and lung cancer were more likely to die in hospital and we were further able to show that hospital death was associated with lack of microscopic diagnosis, and no radiotherapy treatment. This suggests the late admission of patients with advanced stage of disease. Conversely our finding that a microscopically confirmed diagnosis and radiotherapy treatment were associated with home and hospice death suggests that some time within 'the system' may allow for referral to supportive services (Burge et al, 2003). A recent systematic review of factors predicting home death by Gomes and Higginson (2006) found that the six strongest predictors were patients' low functional status, their preferences, home care and its intensity, living with relatives and extended family support. Our new finding that cancer network is an important cause of variation in home death is consistent with this, and probably represents a combination of difference by area in access to home care services, and the nearness of relatives and extended family. It is very unlikely to represent underlying variation in patients' preference for place of death or functional status. Finally our finding that patients from more deprived areas were equally likely to die in hospices and nursing homes as those from affluent areas, contradicts the view that the latter may access these services more often. Inequalities in hospital and home death do, however, persist.

Implications for practice and policy

Our findings reveal that despite increased investment in and provision of palliative care services, cancer patients in South East England remain twice as likely to die in hospital (47%) than at home (23%). The proportion dying at home is now lower than a decade ago, lower than elsewhere in the UK, and far lower than most patients would prefer. Recent national policy has set out the evidence that coordinated palliative care services can allow more people to die at home if they wish (NICE, 2004a, b) and advocated equity of choice in final place of care. This study covers a period before most recent initiatives (Gold Standards Framework, 2006; Marie Curie, 2006) but the variation it finds underlines the need for much more active local surveillance to drive these policies. It also suggests that opportunities exist to learn from differing strategies, organisation and practice within cancer networks. For example, London networks might ask what it is about service **Clinical Studies**

provision in South East London that produces rates of home death similar to those outside London. Networks outside London might ask why hospice deaths are sometimes so high and whether nursing homes are preventing hospital admission and providing better symptom control. Our data also suggest that a good place for clinicians in primary care and acute trusts to start identifying patients in the palliative stage of disease and determining their preference for avoiding or planning admission would be the clinical diagnosis of lung or breast cancer in patients living in deprived areas for whom radiotherapy treatment is not planned. The effect of any change in practice across a network can be monitored easily by the routine work of cancer registries.

Further research

We do not yet fully understand why place of death varies across the UK, how the nexus of factors around the patient operate together to influence this (Gomes and Higginson, 2006) and why home deaths have declined and remain so low in South East England. The imaginative use of available routine data as part of the development of cancer intelligence could help us see more clearly what is happening. For example, trends within individual cancer networks could reveal the influence of different historical patterns of service provision. Ecological studies could show us what happens when a new service such as a hospice opens locally. Mapping rates geographically by primary care trust could show the influence of services (beds and teams) and workforce (district nurses (Shipman et al, 2005), Marie Curie nurses and out of hours care by general practitioners). Studies of how patients move between different services and the interdependence between services are also required. Linking hospital episode statistics data with cancer registration data will, for example, allow us to explore where patients with different cancers are admitted to hospital from, how long they stay and where they are discharged to in the last months of life. The influence that admission has on rate of death in different trusts or primary care trusts explored in a similar way to US studies have done (Wennberg et al, 2004). Qualitative case studies of selected areas could then focus on explaining how different patterns of care are perpetuated, or how change has occurred. Finally in the context of an ageing population, changes in migration and kinship patterns we need to determine older people's preference for death in institutions, ensure that the information we have on what currently occurs is available and public so that where possible people may make their own choices in planning care towards the end of life (WHO, 2004b).

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