Deprivation, distance and death in lung cancer

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There are wide differences in survival rates for lung cancer both within the UK and when comparing the UK with many countries in the western world. Socio-economic inequalities have been shown to have a significant impact on survival for the large majority of cancers in adults in the UK, and many studies have reported lower lung cancer survival rates in patients of lower socio-economic status. There is also wide variation in treatment rates of patients with lung cancer by geographical region, and the paper by Forrest et al in this issue adds to the broadly consistent finding of lower treatment rates in patients from lower socio-economic backgrounds. In addition, many patients with lung cancer die very soon after diagnosis, a significant proportion with a particularly poor prognosis first presenting to secondary care as an emergency admission, a route to diagnosis that is more common in patients of lower socio-economic status. This would imply opportunities for earlier recognition of patients at high risk of lung cancer in primary care. Four important studies appear in this issue of Thorax, three from the UK and one from Australia, that address the issues of earlier diagnosis, early death, undertreatment and geographical distance from specialist centres, all of which have significant implications for optimising service provision for patients with lung cancer.

Despite major improvements to cancer care in England over the last 15 years, the ‘deprivation gap’, in other words, the survival difference between individuals from the least-deprived compared with the most-deprived groups, has not improved significantly in recent years, a finding confirmed and startlingly quantified in a recent report from Public Health England’s National Cancer Intelligence Network. This analysis estimated that both incidence and mortality for all cancers combined (excluding non-melanoma skin cancer) were higher in the more-deprived quintiles than the least-deprived, around 15 300 fewer cancers would have been diagnosed per year in the most recent period examined (2006–2010). Similarly, there was a yearly excess of around 19 200 deaths from cancer in the period 2007–2011. Lung cancer had, by far, the largest number of excess cases (11 700 persons per year) and deaths (9900 persons per year) in the most recent periods.

The study from Forrest et al identifies that the dominant factor in the poorer outcomes for more-deprived patients with lung cancer in the Northern and Yorkshire region of England is that they receive lower rates of treatment than their more affluent counterparts. They had postulated that timeliness of treatment from the point of referral from primary care might be a factor, linked to socio-economic status, but were not able to show that. Khakwani et al, also in this issue, have demonstrated that lung cancer centres in England with specialist thoracic surgical services have higher resection rates for patients referred directly to them than for those referred from the wider and much larger catchment areas that they serve. Reporting in this issue too, Tracey et al, from New South Wales in Australia, found that the further patients live from a specialist thoracic surgical centre, the less likely they are to receive surgical treatment for their lung cancer. In both these studies and others from the UK, the outcomes from larger specialist thoracic surgical centres with greater number of specialist thoracic surgeons were superior to smaller units. Important also is that a clear link between surgical and overall treatment rates and survival at population level has been demonstrated.

This is all very pertinent to the debate about the centralisation of cancer services. There is a clear logic in bringing the best available expertise into a smaller number of centres, with the hope and expectation that the care and survival outcomes of patients treated there would improve, but with this centralisation comes increasing geographical distances that are likely to further widen this deprivation gap. Studies in several cancers have shown that geographical distance from specialist centre and the related transport issues are associated with lower treatment rates and poorer outcomes.

Such geographical and socio-economic factors could well be part of the explanation for the lower survival rates demonstrated by Khakwani et al for the very much more widely spread populations served by larger thoracic surgical centres, although the authors suggest that the explanation might lie in referral practices and service configuration. In the UK, most patients with lung cancer are managed by multidisciplinary teams (MDTs) in District General Hospitals and referred on to larger centres for radical treatment, particularly surgery. While thoracic surgeons do attend these local MDTs, evidence suggests that there are still some patients who might have been considered suitable for radical therapy when discussed at a larger, more specialised MDT, who are being denied these treatments.

Looking at the characteristics of patients seen in primary care that are associated with early death from lung cancer, O’Dowd et al identified low socio-economic status and ‘rurality’ with a risk of early death (multivariate ORs 1.16 and 1.22, respectively), which would be entirely consistent with the importance of deprivation and distance from major (urban) specialist centres. They also showed that, paradoxically, patients who died early after diagnosis had attended their general practitioners (GPs) more often, suggesting that there were missed opportunities for earlier referral.

This plethora of recent inter-related research needs to be heeded by those responsible for the commissioning and delivery of lung cancer services and by those concerned with the public health aspects of healthcare delivery. O’Dowd’s paper points to the likelihood of missed
opportunities for earlier diagnosis in primary care and suggests that better clinical decision aids would be worth investigating. The aim of the government’s ‘Be Clear on Cancer’ campaigns raising public awareness of persistent cough as an early symptom of lung cancer\(^\text{24}\) is to promote earlier presentation to GPs and has tried to target social grades C2, D and E. However, efforts to improve the care for the more deprived are not without their unintended consequences. As long ago as 1971 Tudor Hart described what he called the Inverse Care Law—that the availability of good quality health care is inversely related to need in the population served.\(^\text{25}\) In Dr Forrest’s paper,\(^\text{10}\) she refers to work by White et al\(^\text{10}\) who have described the phenomenon of intervention-generated inequalities, where such things as measures to increase public awareness, promote early presentation and partake of screening programmes are likely to be more actively taken up by people in higher socio-economic status. So while such interventions may improve the health outcomes of the population as a whole, the ‘deprivation gap’ only widens.

In summary, there is overwhelming evidence that patients with lung cancer of lower socio-economic status receive less treatment and have poorer short-term and long-term survival outcomes compared with their more affluent counterparts. The number of excess deaths linked to deprivation is large and the gap between the least and most deprived has not lessened over time. Improved targeting of public awareness campaigns to specific social groups is important so as not to widen this gap even further. Better ways of supporting clinicians in primary care in their difficult task of identifying very high-risk patients to refer for specialist assessment need to be developed and should probably include more accurate clinical decision support tools. Travel times and the availability of public transport need to be taken into account when decisions are being made about service configuration. Every patient should have access to the most expert of multidisciplinary opinion, but clearly the number of highly specialised centres has to be limited, so much more effective hub-and-spoke relationships need to be developed to make decision making as consistent and effective as possible. It has to be our aim to find ways of ensuring equitable access to the highest quality of care for all patients with lung cancer wherever they live and whatever their social background.

Competing interests None.

Provenance and peer review Not commissioned; internally peer reviewed.


Published Online First 13 October 2014

http://dx.doi.org/10.1136/thoraxjnl-2014-205517

http://dx.doi.org/10.1136/thoraxjnl-2014-205554

http://dx.doi.org/10.1136/thoraxjnl-2014-205841

http://dx.doi.org/10.1136/thoraxjnl-2014-205692

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Nearly one in three UK lung cancer patients dies within 3 months of diagnosis, despite serial visits to their doctor

More help needed for family doctors to spot signs and request appropriate and timely tests, say researchers

[What characteristics of primary care and patients are associated with early death in patients with lung cancer in the UK? Online First doi 10.1136/thoraxjnl-2014-205692]
[Deprivation, distance and death in lung cancer Online First doi 10.1136/thoraxjnl-2014-206153]

Nearly one in three lung cancer patients in the UK dies within three months of diagnosis, despite having visited their family doctor several times beforehand, reveals an analysis of primary care data, published online in the journal Thorax.

The findings suggest that family doctors may not be picking up the signs of lung cancer and investigating them as appropriately as they might, or promptly enough, say the researchers.

The UK lags behind other countries in survival rates for lung cancer, a gap that has largely been attributed to late diagnosis of the disease.

In a bid to find out more about the factors in primary care associated with these figures, the researchers analysed family doctors’ (GPs’) investigation of lung cancer between 2000 and 2013.

They used data from The Health Improvement Network (THIN), which contains the anonymised health records of millions of primary care patients across the UK.

They analysed 20,142 cases of lung cancer recorded by 444 general practices during the study period. Of these, one in 20 (5%; 1071) was recorded only on the death certificate.

Of the remainder, one in 10 patients (2036) died within a month of diagnosis; and around one in seven (15%) died within 3 months. These were classified as ‘early’ deaths.

Over half (57%) of all lung cancer deaths were in men: they were 17% more likely to die early than women. The average age at diagnosis was 72. And those aged 80 and above were 80% more likely to die early than those who were diagnosed at a younger age.

Current smokers were also 43% more likely to die early than those who had never smoked, but former smokers were less likely to do so than those who had never smoked.

And an early death was 16% more likely in areas of high deprivation than in more affluent areas. Living in a rural area also increased the odds of an early death from lung cancer.

Further analysis indicated differences in consultation and investigation patterns for patients who died early compared with those who survived for more than three months.

Patients who died early went to see their GP more frequently in the few months before their diagnosis, visiting their general practice an average of five times.

The odds of an early death were lower among those who had had a chest x ray carried out in primary care, irrespective of how many times they had seen their GP before diagnosis. But the odds of an early death rose in tandem with the number of GP consultations they had had.

And patients registered at a practice with high referral rates for chest x-rays were 41% more likely to die early than those registered at practices with low referral rates, even after taking account of other influential factors.

This does not mean that chest x-rays are of questionable value, the researchers emphasise. “Like any investigation, however, it needs to be used at an appropriate time, in a selected patient group,” they write.

And somewhat unexpectedly, a patient living in an area where lung cancer rates were high was no more likely to survive for at least three months than one living in an area where rates were low, the analysis showed.

The researchers point out that the average family doctor is likely to see only one new case of lung cancer a year.

“For this reason we need to promote better use of risk assessment tools, and use software prompts to help GPs to identify and investigate in a timely manner those at risk,” they write. “This has potential to increase the proportion of patients who are diagnosed at an early stage and are, therefore, suitable for treatment with curative intent.”

In a linked editorial, Dr Michael Peake, honorary consultant and senior lecturer at the University of Leicester’s Glenfield Hospital, agrees that GPs need better tools to support their clinical decision making, so that they can pick up patients at risk earlier on.

And he insists: “The number of excess deaths linked to deprivation is large and the gap between the least and most deprived has not lessened over time. Improved targeting of public awareness campaigns to specific social groups is important so as not to widen this gap further.”