Colorectal cancer in the north and south of Ireland 1950–1984

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Abstract

Study objective—Northern Ireland has the highest standardised mortality ratios for colon cancer in the United Kingdom and the Republic of Ireland has some of the highest mortality rates for cancer in the world. The aim of the study therefore was to investigate trends in colorectal cancer in the north and south of Ireland over the period 1950 to 1984.

Design—The study was a cohort analysis of deaths from colorectal cancer for ages 35–74 years by five year age groups, divided by sex.

Setting—This was a population study involving all cases reported to the Registrar General of Northern Ireland and the Eire Vital Statistics and Central Statistical Office during the study period.

Measurements and main results—As in mainland Britain, rectal cancer mortality declined in the north and the south during the study period, but the fall began sooner for males than females. Colon cancer mortality fell in the late 1950s but subsequently rose to its previous high levels.

Conclusions—The observation that there were declines in mortality in the north and south of Ireland in the late 1950s does not support the hypothesis that altered diet due to war rationing in Great Britain and Northern Ireland underlay the fall in British colon cancer mortality after the war. The very high standardised mortality ratios for colon cancer in Northern Ireland highlight a continuing major public health problem in the region.

Mortality from colorectal cancer in the British Isles has changed considerably during this century. There have been important declines in age standardised death rates in England, Wales and Scotland.1–3 Though broadly parallel trends have emerged, the mortality rate in Scotland has remained higher than in England and Wales for 70 years.4 Northern Ireland has the highest standardised mortality ratios for colon cancer within the United Kingdom5 (fig 1), and the Republic of Ireland shares with New Zealand some of the highest national mortality rates in the world.6 7 Variations in mortality over the last 30 years in Northern Ireland have not previously been reported, but trends have been highlighted in the Republic of Ireland.7 8 Notably, colon cancer mortality in both sexes rose significantly between 1950 and 1982,9 with an apparent plateau and slight decline after the mid 1970s.7 However, although allowance may be made for a changing age structure in the population by age standardisation, the interpretation of trends may be complicated by the presence of cohort effects. That such effects may be important in colorectal cancer has already been shown in England and Wales.3

The present paper describes the time trends in colorectal cancer mortality for Ireland as a whole between 1950 and 1984 and findings are contrasted with the epidemiological data from mainland Britain.

Methods

For each year in the 1950–84 period, for ages between 35–74 years, numbers of deaths from cancer of the colon (ICD 153) and rectum (ICD 154) and age and sex specific mid-year population estimates (in five year age groups) were abstracted from the Annual Reports of the Registrar General of Northern Ireland, from the Eire Annual Reports on Vital Statistics and Central Statistical Office, Dublin, and from unpublished data on certain revisions from both of these sources. The total population of Northern Ireland over this time was approximately 1.5 million and of the south, approximately 3 million.

The changes in death rates over the period are displayed graphically, using directly age standardised mortality rates for the seven consecutive quinquennia between 1950–84, standardising to the world population structure.9 Standardised rates for England and Wales are shown for comparison.

The effects of age, period at death, and birth cohort on mortality rates have been investigated for each disease and sex, by using the age-period-cohort modelling approach10 for all deaths between 35 and 74 years. Briefly, the significance of age, period, and cohort effects may be assessed by fitting a series of nested models and using the changes in goodness of fit to obtain likelihood ratio $\chi^2$ statistics. Older age groups were not included because of the doubt about the coded cause of death in the very old.

Results

Figures 2 to 5 show the time trends in colorectal cancer in Ireland and England and Wales during the 1950–84 period. Important differences emerge between diseases, between the sexes, and to some extent between countries. Figures 2 and 3 show that, although death rates from colon cancer in 1950 differed little in the three regions, a marked differential in mortality between Ireland...
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and England and Wales had emerged by 1980–84. Ireland as a whole, in keeping with England and Wales, showed some decline in mortality during the 1950s, perhaps more marked among males, but by the early 1960s mortality rose in an almost parallel fashion in the north and south of Ireland. In females, mortality appears to plateau and then decline slightly in the early 1980s.

For rectal cancer figs 4 and 5 indicate that the mortality in Ireland and England and Wales has shown broadly comparable declines over the study period. In Northern Ireland, declines in rates among females began somewhat later than in the other countries.

Likelihood ratio χ² tests obtained from fitting various models to the Irish data are shown in the table.

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The first line of the table indicates that, for males, period effects were significant when added to the model containing terms for age. Only for males in the Republic were cohort effects significant (line 2). In Northern Ireland males, line 3 shows that period effects persisted after adjustment for cohort. In the Republic there was significant lack of fit in the full age-period-cohort model, and the period and cohort effects in lines 3 and 4 were no longer significant when adjusted for extra Poisson variation. For Republic of Ireland males, secular trends could therefore be explained equally well by either period or cohort effects, but for Northern Ireland males period effects were clearly preferable.

There was no evidence of significant period or cohort effects for females either in Northern Ireland or in the Republic.
Table 4: Likelihood ratio $\chi^2$ tests for period and cohort effects

<table>
<thead>
<tr>
<th>Effect</th>
<th>Degrees of freedom</th>
<th>Northern Ireland</th>
<th>Republic of Ireland</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Males</td>
<td>Females</td>
</tr>
<tr>
<td>Colon</td>
<td>Period/Age</td>
<td>6</td>
<td>14.6*</td>
</tr>
<tr>
<td></td>
<td>Cohort/Age</td>
<td>12</td>
<td>10.9</td>
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<tr>
<td></td>
<td>Period/Age, Cohort</td>
<td>5</td>
<td>11.2*</td>
</tr>
<tr>
<td></td>
<td>Cohort/Age, Period</td>
<td>12</td>
<td>7.5</td>
</tr>
<tr>
<td>Rectum</td>
<td>Period/Age</td>
<td>6</td>
<td>34.5*</td>
</tr>
<tr>
<td></td>
<td>Cohort/Age</td>
<td>13</td>
<td>46.7*</td>
</tr>
<tr>
<td></td>
<td>Period/Age, Cohort</td>
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</tr>
<tr>
<td></td>
<td>Cohort/Age, Period</td>
<td>12</td>
<td>15.3</td>
</tr>
</tbody>
</table>

* Effect: Period/Age—The effect of fitting period after fitting age
  Cohort/Age—The effect of fitting age after fitting period
  Period/Age, Cohort—The effect of fitting period and cohort

* p < 0.05; t p < 0.01; # non-significant after allowing for extra Poisson variation.

The limited evidence available would therefore appear to support period effects in preference to cohort effects for the explanation of Irish mortality trends.

Discussion

The purpose of this paper is to describe the trends in colorectal cancer in Ireland, north and south, and to put these trends in the context of the mainland Britain trends over the same period. It is not the intention to enter into the continuing debate on the usefulness of particular methods of separating the effects of age, periods or cohort from a vital rate matrix.12

As has been indicated elsewhere1 2 the trends are not thought to be attributable to changes in coding and certification practices, especially when the “very old” deaths have been excluded. Any such changes would be expected to have the same effect on males and females.3 An alternative may be that changing diagnostic practices could account for the observed trends. This point warrants careful consideration. The most important diagnostic innovation during the period (which may have had greater effects on more proximal colonic tumours) was the introduction of double contrast barium enema in the late 1960s. It is hard to concede that this was the explanation for the rise in colon cancer mortality in Ireland in the late 1960s, as colonic and rectal disease mortality followed downward trends on mainland Britain where diagnostic access presumably increased during a roughly equivalent period. Indeed the near parallelism of the rise in colon cancer mortality north and south in the late 1960s would be difficult to explain on the basis of diagnostic access. It is unlikely that such access diffused at the same rate in the south as in the north of Ireland since, for much of this period, there was no National Health Service in the Republic of Ireland.13

These trends would be greatly clarified if accurate local incidence data were available. Some of the decrease in mortality that occurred in the UK mainland after 1950 has been attributed to improved survival.2 Differences in survivorship between small areas have been found elsewhere to underlie geographical mortality differentials,14-16 but without high quality incidence data broken down by subsite,17 such a premise would be hard to evaluate further in Ireland.

More interesting perhaps, in view of a renewed emphasis on the primary prevention of cancer, are the putative effects of changing exposures to dietary risk factors. It has been contended that the declines in colorectal cancer mortality observed on mainland UK in the 1950s were attributable to a large degree to the healthier diet adopted by most of the population during war rationing 10 years earlier.1 2 18 During this period, consumption of meat and fat products decreased and crude fibre intake increased.2 In general, it is accepted that dietary factors affect the risks of colorectal disease more than rectal disease,19 but although the decline in rectal cancer mortality in Ireland seems more consistently to reflect similar mainland declines, the contrast with colon cancer may be significant. The similar falls in the North Island diet reflected those of the mainland during the war, no such rationing was introduced in the South. Indeed a recent analysis of the 1948 National Food Survey in the Republic of Ireland clearly indicates that intakes of fat and meat products were well above those of UK citizens at that time (M Crawford, personal communication).

However the weaknesses of such ecological inference in epidemiology are well known.20 21 Though a similar anomaly was described in New Zealand,6 where excess dietary risks were similar for both Maoris and non-Maoris during a period when colorectal cancer was falling among Maoris and rising among Europeans, it is possible that genetic variation in New Zealand could have accounted for some of the inter population differences in responses to either “risk” or “protective” factors.22

The fact that the age-period model was the most statistically parsimonious lends support to the use of a simple graphical plot of age standardised rates to display the data. It is unwise, however, on the basis of such statistical models, to make too far reaching assumptions about the origin of regional or gender variations in period or
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cohort trends. Osmond et al have inferred that the origin lies in dietary practices. Other factors such as a changing parity distribution may be important but have been given scant attention. In view of the growing evidence for cancer as a multistep process at the cellular level the identification of period effects, traditionally relying on the detection of roughly equivalent changes in log transformed incidence rates across age groups, may be conceptually inadequate. Age may well be an effect modifier. Changes in diet, whether adverse or beneficial, may have different effects on different age groups or different subsites within the bowel and may well interact with genetic factors in undiscernible ways. Excesses such as these can at best be descriptive. The use of statistical analyses of epidemiological data to infer biological processes can be misleading. Even when acting independently, the actions of some carcinogenic factors may fit an additive statistical model, a multiplicative model, or neither.

Trends in colorectal cancer mortality both in Northern Ireland and the Republic of Ireland do not mirror the changes in mainland Britain. In particular, the Irish data are characterised by continuing high male colon cancer mortality rates and only a very recent decline in female colorectal cancer mortality rates. The explanation for these differences must await a more adequate paradigm than ecological inference can provide.

15 Piantadosi S, Byar DP, Gail M. Screening geographic areas for unusual survival experience or stage at diagnosis with application to breast and colon cancer. J Natl Cancer Inst 1985; 75: 269–75.