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The role of private care in the interval between diagnosis and treatment of breast cancer in Northern Ireland: an analysis of Registry data

Patricia Carney, Anna Gavin, Ciaran O'Neill

ABSTRACT

Objective: To examine the differences in the interval between diagnosis and initiation of treatment among women with breast cancer in Northern Ireland. Design: A cross-sectional observational study. Setting: All breast cancer care patients in the Northern Ireland Cancer Registry in 2006. Participants: All women diagnosed and treated for breast cancer in Northern Ireland in 2006. Main outcome measure: The number of days between diagnosis and initiation of treatment for breast cancer.

Results: The mean (median) interval between diagnosis and initiation of treatment among public patients was 19 (15) compared with 14 (12) among those whose care involved private providers. The differences between individual public providers were as marked as those between the public and private sector—the mean (median) ranging between 14 (12) and 25 (22) days. Multivariate models revealed that the differences were evident when a range of patient characteristics were controlled for including cancer stage.

Conclusions: A relatively small number of women received care privately in Northern Ireland but experienced shorter intervals between diagnosis and initiation of treatment than those who received care wholly in the public system. The variation among public providers was as great as that between the public and private providers. The impact of such differences on survival and in light of waiting time targets introduced in Northern Ireland warrants investigation.

INTRODUCTION

Breast cancer incidence rates have risen in Britain over the past 40 years from 73.8/100 000 women in 1975 to 126.2/100 000 in 2010. As the number of women with the disease has increased, so too has pressure on the healthcare system to develop effective strategies in respect of its management. While debate continues as to the role that should be accorded screening as part of the overall management strategy, an early detection and treatment have been shown in a number of studies to be significant determinants of the cost and outcome. Previous work in the Republic of Ireland with respect to screening and in the USA with respect to treatment have demonstrated that the possession of private medical insurance contributes to variations in uptake and services provided, respectively. Similarly, a work in the USA with respect to outcomes has shown these to vary between the public and private care facilities. Evidence from the USA also suggests that women in possession of private health insurance have a more favourable stage of disease at breast cancer diagnosis than non-insured women. While this may be expected in healthcare systems where there exists a significant role for private finance or provision (in the USA and Ireland approximately 50% of the adult population hold private medical insurance), in systems where private finance and/or delivery of core services are the exception rather than the norm, one might not expect policymakers to be as sanguine as to the existence of such differences or for such difference to be evident as a result. As healthcare budgets come under increasing pressure, so evidence of an apparent superiority of private medicine in contexts such as the UK may receive keen attention among policymakers.
struggling to provide quality services in an efficient manner. While evidence that delays in initiation of treatment are material in a clinical sense are mixed\textsuperscript{15,16} that delays may well impact on anxiety levels among patients is evident.\textsuperscript{17} In this paper, we examine the differences in the interval between diagnosis and initiation of care in patients with breast cancer among the private and public providers in Northern Ireland. We examine the intervals between diagnosis and treatment among and between the public and private providers and discuss the implications this may have for the commissioning of services.

**METHODS**

Data for all women diagnosed with breast cancer in Northern Ireland in 2006 were obtained from the Northern Ireland Cancer Registry (NICR). The Registry is a state-funded body that registers all cancer diagnoses in Northern Ireland and contains a range of data on the patient, their diagnosis and treatment. A total of 951 women were recorded by the registry as being diagnosed with breast cancer in 2006. Data on 848 women in the NICR were extracted for analysis. Data included the details on the care provider (healthcare and social care Trust) in which the woman was treated and whether the woman had a record of being diagnosed or treated at any stage of the care pathway privately. In addition, data on the cancer stage (I–IV), the woman’s age at diagnosis, the socioeconomic deprivation score for the area in which she resided, whether she resided in a rural or urban area, whether or not she experienced comorbid conditions and the number of symptoms experienced. Data on the source of the referral and the Trust where the woman presented were also extracted. The complete details of all variables are presented in online supplementary appendix 1. The interval used here is the duration time between diagnosis of breast cancer and the initiation of any treatment, that is, pre-treatment in terms of the Aarhus Statement.\textsuperscript{18}

A Cox’s proportional hazards regression model\textsuperscript{19} was used to examine whether women who were seen at any stage of their cancer journey privately had a different interval between diagnosis and initiation of treatment when other variables were controlled, than women who were dealt with entirely within the publicly funded system. Duration was measured in days. Women whose treatment was initiated on the same day as diagnosis were treated as having an interval between treatment and diagnosis of 1.

A secondary analyses examined the variation among the public providers using a Cox regression model.\textsuperscript{19} The proportional hazards assumption for the Cox regression model is tested using Schoenfeld residuals.\textsuperscript{20}

**RESULTS**

In table 1, descriptive statistics on a range of variables for women included in the Cox proportional hazards regression model are presented.

A total of 795 women were seen solely in public hospitals and 89 at some stage in private hospitals. The mean duration between diagnosis and initiation of treatment had a record of being diagnosed or treated at any stage of the care pathway privately. In addition, data on the cancer stage (I–IV), the woman’s age at diagnosis, the socioeconomic deprivation score for the area in which she resided, whether she resided in a rural or urban area, whether or not she experienced comorbid conditions and the number of symptoms experienced. Data on the source of the referral and the Trust where the woman presented were also extracted. The complete details of all variables are presented in online supplementary appendix 1. The interval used here is the duration time between diagnosis of breast cancer and the initiation of any treatment, that is, pre-treatment in terms of the Aarhus Statement.\textsuperscript{18}

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**RESULTS**

In table 1, descriptive statistics on a range of variables for women included in the Cox proportional hazards regression model are presented.
Table 2 Duration between diagnosis of breast cancer and beginning of treatment

| Number of days between diagnosis and treatment of breast cancer | Public patients (N=759) | Private patients (N=89) |
|---------------------------------------------------------------|-------------------------|-------------------------|
| Treatment initiated on the same day                          | 4%                      | 5%                      |
| Treatment initiated between 1 and 10 days after diagnosis      | 27%                     | 41%                     |
| Treatment initiated between 11 and 15 days after diagnosis     | 23%                     | 20.5%                   |
| Treatment initiated between 16 and 25 days after diagnosis     | 23.4%                   | 20.5%                   |
| Treatment initiated between 26 and 75 days after diagnosis     | 22%                     | 13%                     |
| Treatment initiated between 76 and 121 days after diagnosis    | 0.6%                    | 0%                      |
| Average time between diagnosis and initiation of treatment    | 19 days                 | 14 days                 |
| Median time between diagnosis and initiation of treatment     | 15 days                 | 12 days                 |
| 10th centile                                                 | 5 days                  | 3 days                  |
| 25th centile                                                 | 9 days                  | 6.5 days                |
| 50th centile                                                 | 15 days                 | 12 days                 |
| 75th centile                                                 | 24 days                 | 18 days                 |
| 90th centile                                                 | 36 days                 | 28 days                 |
| 95th centile                                                 | 48 days                 | 33 days                 |

Table 3 shows each of the number of women in the dataset by trust in which they lived and the trust in which they presented, for example, 179 women diagnosed with breast cancer in 2006 resided in the Belfast Trust area but 237 women presented through the Belfast Trust. As illustrated the Trust with the highest number opting for private healthcare is the South Eastern Health and Social Care Trust (HSCT) with the lowest number coming from the Western Trust.

Figure 1 illustrates the variation in interval for both groups of patients. The y-axis represents the number of days between diagnosis and treatment; the whiskers plot the CI.

The results from Cox’s proportional hazards regression model in which we focus on the distinction between public and private controlling for a range of variables—including age, stage, source of referral, comorbid status and deprivation score—are presented in table 4. Coefficients in the table may be interpreted as follows: if the coefficient is greater than 1, treatment is likely to be initiated sooner after diagnosis; if the coefficient is less than 1, the interval between diagnosis and initiation of treatment is likely to be longer. As can be seen, controlling for confounding variables, women diagnosed or treated at any stage privately are likely to have a shorter interval between diagnosis and initiation of treatment than those whose diagnosis and treatment are wholly within the public system. The p value for the Schoenfeld residual test is presented at the end of table 3. Obtaining a p value greater than 0.05 indicates that the proportional hazards assumption for the Cox regression model is not violated.

Examining the variations in the interval between diagnosis and initiation of treatment among providers reveals that, on an average, the interval was 15 days for Belfast HSCT (n=207), 18 days for the Northern HSCT (n=109), 14 days for private hospitals (n=89) or the South-Eastern HSCT n=(100), 25 days for the Southern HSCT (n=102), 19 days for the Western HSCT (n=98) and 24 days for the breast screening programme (n=143). This indicates wide variations among publicly funded care providers. In table 5, results of a Cox regression that controls for individual trusts are presented, as well as confounding variables are presented. As can be seen while the coefficient on private hospitals remains for women seen at some stage in private hospitals was 14 days and that for those seen solely in publicly funded hospitals is 19 days. The difference in the interval between diagnosis and initiation of treatment was statistically significant before other variables were controlled for, p<0.01. The differences in the percentages of private versus public patients in terms of the intervals between diagnosis and initiation of treatment are reported in table 2 at various durations.
significant, so too is that on several of the publicly funded care providers.

DISCUSSION

In the interests of brevity, we focus our discussion on the impact of public versus private care on the interval between diagnosis and initiation of treatment as well as among the public sector providers. The differences are evident between patients whose entire care was within the public sector and those who were seen at some point in the private sector. That private medicine (funding or provision) should impact on care in a mixed healthcare system is unsurprising. Self-interest would suggest the reason individuals hold private medical insurance is because of the superior service—including the speed with which individual’s treatment might be initiated— it affords and private providers operate because there exists an effective demand for the services they provide. As noted, the previous studies in the area of breast cancer have demonstrated the existence of difference whether in service uptake, provision or outcome in such systems. That differences in terms of the speed with which care is initiated appear to exist between the public and private sectors in a country whose publicly funded healthcare system extols equality of access based on need, and not the ability to pay is not perhaps what one might expect. With specific regard to the interval between diagnosis and initiation of treatment, general concerns have indeed been explicitly referenced by the Department of Health. Crucially though these have focused on variations per se and not on variations between public and private providers. This study has clearly demonstrated that such concerns (at least in 2006) were justified in the Northern Ireland.

Table 4 Cox’s proportional hazards regression model output

| Variable | HR (SE) |
|----------|---------|
| Urban/rural | 1.045 (0.075) |
| Age 41–50 | 1.079 (0.164) |
| Age 51–55 | 0.92 (0.149) |
| Age 56–60 | 1.21 (0.195) |
| Age 60–65 | 1.024 (0.172) |
| Age 66–70 | 1.167 (0.222) |
| Age 71 or greater | 1.128 (0.169) |
| Have a comorbidity | 0.995 (0.005) |
| Stage 2 | 1.017 (0.08) |
| Stage 3 | 0.972 (0.109) |
| Stage 4 | 0.909 (0.147) |
| Number of symptoms | 0.988 (0.019) |
| Quintile 2 | 0.885 (0.093) |
| Quintile 3 | 0.849 (0.095) |
| Quintile 4 | 0.8** (0.088) |
| Least deprived | 0.96 (0.108) |
| Belfast HSCT | 0.962 (0.15) |
| Northern HSCT | 0.732* (0.119) |
| Breast screening unit | 0.475*** (0.074) |
| South-Eastern HSCT | 0.957 (0.156) |
| Southern HSCT | 0.464*** (0.076) |
| Western HSCT | 0.68** (0.113) |

Schoenfeld residual test p value <0.05

Numbers in parentheses are SEs.

*, 90% significance level; **, 95% significance level; ***, 99% significance level.

HSCT, Health and Social Care Trust.

Table 5 Cox’s proportional hazards regression model output—trust of presentation

| Variable (N=848) | HR (SE) |
|------------------|---------|
| Urban/rural | 1.125 (0.082) |
| Age 41–50 | 1.079 (0.164) |
| Age 51–55 | 0.92 (0.149) |
| Age 56–60 | 1.21 (0.195) |
| Age 60–65 | 1.024 (0.172) |
| Age 66–70 | 1.167 (0.222) |
| Age 71 or greater | 1.128 (0.169) |
| Have a comorbidity | 0.995 (0.005) |
| Stage 2 | 1.017 (0.08) |
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Numbers in parentheses are SEs.

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variables are controlled for), the dichotomy between the public and private provision is false in the sense that best practice among public providers was on a par with that in the private sector. Poor performance among some public providers was not evident among all of them. It is unclear whether the difference between the best and worst performing providers in terms of the interval in the initiation of treatment was material with respect to the outcome, though others have found delays may be a source of anxiety to some patients and as noted with respect to survival. In their investigation into the effect of waiting times between localised breast cancer diagnosis and treatment, Redaniel et al found that the duration times have a very little impact on survival. However, as highlighted by Paul et al, longer interval times can have negative psychological impacts on patients with cancer. This also raises the question of the effect of longer duration times on non-localised breast cancer. Our results show that there is no stage differential in the interval between diagnosis and initiation of treatment.

The cost and outcomes do not form part of the analysis undertaken and no inference with respect to these is made. The difference in the speed with which the system responds may nevertheless be taken as a valid indicator of quality. What might constitute an appropriate policy response to such delays, this study suggests, will depend on the particular circumstances that explain poor performance in the case of individual providers. These may relate to the staffing issues unique to the period studied, other capacity issues or broader managerial deficiencies. That the issue is public sector provision per se is clearly not the case. As public expectations rise and healthcare budgets come under increasing pressure, debate as to how best to deliver care in breast cancer and other services will intensify. Privatisation will inevitably be mooted by some as a solution. These results, however, indicate that with respect to breast cancer services, there is no reason to believe it offers a superior service to that which can be delivered by the public sector.

Since 2006, waiting time targets for patients with cancer have been introduced in Northern Ireland. Whether this has impacted on the speed with which treatment is initiated (ie, delay in receiving treatment after a positive diagnosis) or differences among and between public and private providers in this regard require further examination. What, if any, impact such delays or failure to reach targets may have had on survival and cost of care are also areas that seem worthy of further examination.

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Competing interests None.

Ethics approval Study involves the secondary analysis of anonymised cancer registry data.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement The Cancer Registry maintains details of all incident cases of cancer in Northern Ireland. Details of the data and arrangements for access can be obtained from the Northern Ireland Cancer Registry.

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