Epidemiology

Survival following a diagnosis of heart failure in primary care

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Abstract

Background. Heart failure is a common long term condition affecting around 900,000 people in the UK and patients commonly present to primary care. The prognosis of patients with a code of heart failure in their primary care record is unknown.

Objective. The study sought to determine the overall survival rates for patients with heart failure in a primary care population from the time of diagnosis.

Methods. Survival analysis was carried out using UK primary care records from The Health Improvement Network (THIN) between 1 January 1998 and 31 December 2012. Patients age 45 or over with a first diagnostic label of heart failure were matched by age, sex and practice to people without heart failure. Outcome was death in the heart failure and no heart failure cohorts. Kaplan-Meier curves were used to compare survival. Age-specific survival rates at 1, 5 and 10 years were determined for men and women with heart failure. Survival rates by year of diagnosis and case definition were also calculated.

Results. During the study period, 54,313 patients had a first diagnostic code of heart failure. Overall survival rates for the heart failure group were 81.3% (95% CI 80.9–81.6), 51.5% (95% CI 51.0–52.0) and 29.5% (95% CI 28.9–30.2) at 1, 5 and 10 years respectively and did not change over time.

Conclusions. In a primary care population, the survival of patients diagnosed with heart failure did not improve over time. Further research is needed to explain these trends and to find strategies to improve outlook.

Key words: Community, heart failure, prognosis, primary care, survival.

Introduction

Heart failure is an important public health problem associated with significant morbidity and mortality for patients and high costs for healthcare systems (1). An estimated one to two in every 100 adults in Western populations are living with heart failure (2). In the UK, the costs of heart failure to the National Health Service (NHS) are second highest for any disease after stroke (3). Accurate estimates of heart failure prognosis are vital to healthcare commissioners to allow appropriate allocation of resources, to physicians in making management decisions and, perhaps most importantly, to patients to allow informed choices about treatments and end of life care (4).

Heart failure prognosis has previously been reported as worse than many cancers although the outcome of patients with heart failure is not always reliably recorded (5). Cancer registries help to identify patterns in both incidence and survival of common cancers...
and have reported significant improvements in prognosis over time (6). Global heart failure registries are lacking so outcomes have been established from hospital data and screening studies.

Analysis of hospital records from a cohort of 5.1 million patients in Scotland reported median survival rates of 1.33 years (95% CI 1.17–1.50) in men following a first hospitalisation for heart failure in 1986. With an increase in prescribing of evidence-based treatments (angiotensin converting enzyme inhibitors (ACE-I), B blockers and mineralocorticoid receptor antagonists) survival at 1 year increased to 2.34 years (95% CI 2.15–2.56) by 2003 (7).

Table 1. Characteristics of participants with heart failure and matched participants without heart failure from The Health Improvement Network (THIN) database between 1 January 1998 and 31 December 2012

| Characteristic | Heart failure | No heart failure |
|----------------|---------------|-----------------|
| Age (years)    | n = 54313     | n = 254809      |
| 45–54          | 1738 (3.2%)   | 8685 (3.4%)     |
| 55–64          | 5268 (9.7%)   | 26323 (10.3%)   |
| 65–74          | 12127 (22.3%) | 60308 (23.7%)   |
| 75–84          | 20942 (38.6%) | 100184 (39.3%)  |
| 85–94          | 13021 (24.0%) | 55311 (21.7%)   |
| ≥95            | 1217 (2.2%)   | 3998 (1.6%)     |
| Missing        | 27983 (51.5%) | 130413 (51.2%) |
| Townsend score |               |                 |
| 1              | 10935 (20.1%) | 59001 (23.2%)   |
| 2              | 11156 (20.5%) | 55974 (22.0%)   |
| 3              | 11089 (20.4%) | 50893 (20.0%)   |
| 4              | 10864 (20.0%) | 45959 (18.0%)   |
| 5              | 7940 (14.6%)  | 31956 (12.5%)   |
| Not known      | 2329 (4.3%)   | 11026 (4.3%)    |
| BMI            |               |                 |
| Mean (SD)      | 27.9 (5.9)    | 26.4 (4.7)      |
|Missing         | 8664 (16.0%)  | 51692 (20.3%)   |
| Smoking status |               |                 |
| Smoker         | 7656 (14.1%)  | 32803 (12.9%)   |
| Not current smoke | 43198 (79.5%) | 196407 (77.1%)  |
| Missing        | 3459 (6.4%)   | 25599 (10.1%)   |
| Diabetes       | 10729 (19.8%) | 27483 (10.8%)   |
| Hypertension   | 28820 (53.1%) | 106661 (41.9%)  |
| Angina         | 12522 (23.1%) | 28656 (11.3%)   |
| Myocardial infarction | 12282 (22.6%) | 16339 (6.4%)   |
| Ischaemic heart disease | 23308 (42.9%) | 43803 (17.2%) |
| Stroke         | 4969 (9.2%)   | 14983 (5.9%)    |
| Atrial fibrillation | 15473 (28.5%) | 17969 (7.1%)   |
| Valve disease  | 5322 (10.2%)  | 6281 (2.5%)     |

Table 2. One-, five- and ten-year survival rates in participants with heart failure and matched participants without heart failure from The Health Improvement Network (THIN) database between 1 January 1998 and 31 December 2012

| Age (years) | Survival (%) |
|-------------|--------------|
|             | 1 year (95% CI) | 5 years (95% CI) | 10 years (95% CI) |
| Heart failure | No heart failure | Heart failure | No heart failure | Heart failure | No heart failure |
| 45–54        | 94.0 (92.7–95.0) | 100.00 (99.5–99.8) | 80.9 (78.6–83.0) | 97.8 (97.4–98.2) | 68.2 (64.4–71.6) | 95.4 (94.6–96.1) |
| 55–64        | 91.5 (90.7–92.3) | 99.1 (99.0–99.3) | 75.0 (73.6–76.4) | 94.7 (94.4–95.0) | 58.2 (56.0–60.3) | 87.8 (87.1–88.5) |
| 65–74        | 87.6 (87.0–88.2) | 97.5 (97.4–97.7) | 65.4 (63.4–65.4) | 86.6 (86.3–86.9) | 40.4 (39.0–41.8) | 71.4 (70.8–71.9) |
| 75–84        | 81.9 (81.3–82.4) | 94.2 (94.0–94.3) | 49.5 (48.7–50.3) | 71.9 (71.6–72.2) | 23.8 (22.8–24.9) | 46.0 (45.5–46.6) |
| 85–94        | 70.7 (69.8–71.5) | 87.5 (87.2–87.8) | 28.5 (27.5–29.6) | 49.6 (49.0–50.1) | 7.9 (6.8–9.0) | 19.2 (18.4–19.9) |
| ≥95          | 53.8 (50.7–56.7) | 77.5 (76.0–78.8) | 9.0 (6.5–12.0) | 24.3 (22.3–26.3) | 0 — | 0 — |
| Overall      | 81.3 (80.9–81.6) | 94.0 (93.9–94.1) | 51.5 (51.0–52.0) | 74.0 (73.8–74.3) | 29.5 (28.9–30.2) | 54.1 (53.8–54.4) |
Practices that contributed at least 1 year of clinical data were included in the study. In addition, because there may be under recording of mortality in earlier years, data were only included after each practice’s acceptable mortality reporting date (13). This is the date after which recorded mortality in the practice is consistent with predicted mortality. This was to ensure that the main outcome of interest was being accurately recorded by the participating practices.

Population
The dataset extracted from THIN database included all persons aged 45 years and over, registered at a practice for at least 12 months during the study period. This age cut-off was chosen because the types of heart failure affecting children and younger people are pathologically distinct from those found in older adults. Eligible cases had a clinical code of heart failure. The index date was the first date of a recorded heart failure code. Patients with a previous diagnosis of heart failure, either prior to age 45 or the study start date, were excluded. Cases were matched with up to five comparator patients who were registered in the same practice on the index date and did not have a diagnosis of heart failure on that date (but could become a case later). Comparators were also matched on sex and age ±5 years.

Clinical codes
Patients with a diagnosis of heart failure were identified using clinical codes input by GPs to record new diagnoses in the medical record. The NHS Clinical Terminology Browser and Quality and Outcomes Framework (QOF) guidelines were used to generate a comprehensive list of clinical codes used to record a diagnosis (Supplementary Table S1). Heart failure is a clinical syndrome and the diagnosis requires the presence of symptoms and objective evidence of a structural or functional abnormality of the heart. Patients with a clinical code of heart failure and either an echocardiograph report or a hospital letter were classified as being a confirmed case of heart failure and those with just a clinical code alone as an unconfirmed case.

Baseline variables
Demographic variables including age, sex, ethnicity, area deprivation quintile, cardiovascular risk factors and co-morbidities were extracted. The latest deprivation quintile prior to the index date was used or, if unavailable, the most recently recorded after the index date. Cardiovascular risk factors (smoking, blood pressure, cholesterol, body mass index (BMI)) were the most recently recorded prior to the index date. Cardiovascular co-morbidities (angina, myocardial infarction (MI), ischaemic heart disease, diabetes, hypertension, stroke, atrial fibrillation (AF), valve disease), were defined by the presence of a clinical code at any time prior to the index date.

Outcome measure
The outcome measure was overall survival rate for the cohort which was determined using the date of death for patients who had died from any cause (all-cause mortality).

Statistical analyses
Data were extracted directly from THIN database using clinical codes. Analysis was carried out using Stata versions 10 and 11. The absolute number of heart failure cases, and overall incidence rate by age and sex, was calculated.

Heart failure patients and matched patients without heart failure, included in the survival analyses were followed until the earliest of the following dates: patient died, patient left (de-registered from) their practice, the practice ceased contributing data to THIN, or the study ended. Kaplan-Meier curves were used to compare survival in the heart failure and no heart failure cohorts. Log rank tests were used to compare survival between groups.

One-, five- and ten-year survival rates for the heart failure cohort were calculated by 10-year age band from the age of 45. For comparison, survival rates of the no heart failure cohort were also determined. To examine trends over time, age-sex specific and overall survival rates were calculated by year of diagnosis. Case definition was also explored by calculating overall survival rates by year of diagnosis for the confirmed and unconfirmed heart failure groups separately.

Results
Five hundred and sixty four practices contributed at least 1 year of data between 1 January 1998 and 31 December 2012. A total
of 2,728,841 patient records were included in the dataset; 54,313 patients had a new clinical code of heart failure during the study period. Overall heart failure incidence was 3.00 (95% CI 2.98–3.03) per 1000 person-years, increasing with age and higher in men than women in every age band. Incidence of heart failure in age group 65 and older was 6.78 (95% CI 6.72–6.84) per 1000 person-years and in the age group 75 years and older was 10.86 (95% CI 10.74–10.97) per 1000 person-years.

Survival in heart failure and no heart failure cohorts

The characteristics of patients with and without heart failure are shown in Table 1. All five Townsend deprivation quintiles were similarly represented except for the most deprived group which had around one-third less cases than the other four groups. BMI was similar but there were 1.2% more smokers in the heart failure group. Cardiovascular co-morbidities, particularly ischaemic heart disease, AF and valvular disease were more common in patients with heart failure.

Overall, heart failure cases had a significantly worse prognosis than comparators (log rank test, chi-square (1) 12,732.59, 1, P < 0.0001). The survival rates for the heart failure cohort, and age, sex and practice matched cohort without heart failure, are shown in Table 2. Overall survival rates in the heart failure group were 81.3% (95% CI 80.9–81.6) at 1 year from date of diagnosis, 51.5% (95% CI 51.0–52.0) at 5 years and 29.5% (95% CI 28.9–30.2) at 10 years.

Survival over time

The survival rates of all heart failure cases by year of diagnosis are shown in Figure 1. One- and five-year survival rates remained stable during the study period at 80% and 50%, respectively. Ten-year survival rates for those diagnosed between 1998 and 2002 were 28%.
One-, five- and ten-year survival rates for men and women by age group are shown in Figure 2a–c, respectively. Survival rates for all groups did not change significantly over time.

Survival by case definition

The number of confirmed cases (13,773) made up just over a quarter (26.5%) of the total number of heart failure cases (54,313) in the dataset overall and did not increase significantly over a 10-year period. Cases with confirmed heart failure were younger and 56% were male compared to 50% in the unconfirmed group. The proportion of patients in each Townsend quintile was similar for confirmed and unconfirmed cases. There were 2% more smokers in the confirmed compared to the unconfirmed heart failure group. Ischaemic heart disease, angina and MI were all more common in the confirmed group.

The overall survival rates were lower in the unconfirmed heart failure group than in the confirmed heart failure group (log rank test, chi-square (1) 170.37, 1, P < 0.0001). Figure 3 shows 1-, 5- and 10-year survival rates in the confirmed and unconfirmed heart failure groups over time. The unconfirmed heart failure group had an overall 5% lower survival rate at 1, 5 and 10 years.

Conclusions

This study provides contemporary survival rate estimates for patients with a first diagnostic code of heart failure in their GP record. The outlook for patients with heart failure did not improve over time. Patients with a confirmed diagnosis had a slightly better outcome than those that were unconfirmed.

The cohort in this analysis included 54,313 patients with heart failure, much larger than most prospective studies, and the follow-up
period was over a 15-year period. The NHS provides healthcare to the entire population of the UK, free at the point of access and most patients are registered with a primary care provider. Unlike screening cohort studies, primary care databases do not rely on participants volunteering to take part in the study rather they represent a cross-section of the entire population. These results are therefore likely to be generalizable to the community population as a whole (14). The large number of patients in each age and sex category also improves the accuracy of the survival rate estimates.

The main limitation of the study is the reliability of GP coding. Research using general practice databases is reliant on the accuracy of clinical coding input by GPs during the consultation. Heart failure is a chronic condition which is often insidious in onset and can masquerade as other conditions making early and accurate diagnosis difficult. Over time the use and accuracy of clinical coding has improved, particularly since the introduction of the QOF in 2004 (15). The definition of heart failure has also changed over time leading to dysynchrony between epidemiological studies, making comparison difficult, and the meaning of a code for heart failure in an individual’s medical record may be based on different criteria depending on the accepted definition at the time of diagnosis. The benefit of THIN and similar primary care databases are that they provide an insight into real-life general practice and the survival rates of patients with a clinical code of heart failure in their record is likely to be an important statistic for practising GPs.

There is a paucity of contemporary epidemiological information on heart failure from a primary care setting. Annual QOF data shows a prevalence of 0.8–0.9% each year since the introduction of the heart failure indicator in 2006 but the number of new cases per year is not recorded (16). National heart statistics include heart failure incidence and mortality within an umbrella term of ‘cardiovascular disease’ so estimates for heart failure alone are lacking. Survival
rate estimates have therefore come from screened cohorts and hospital databases. The ECHOES study is one of the few studies to report 10-year, as well as 5-year, survival rates (10). All deaths in the cohort were collated from routinely collected mortality data. Overall the mortality rate for all-cause heart failure was 9% per year; survival was 53% at 5 years and 27% at 10 years. The overall 10-year survival rate of 29.5% for the heart failure group in the THIN study is therefore similar to the screened population in ECHOES.

Sweden has well-developed, high-quality health registries which allow more detailed examination of the epidemiology of diseases. A cross-sectional study of 2.1 million residents of the Stockholm region was carried out using an administrative health data register (17). Between 2006 and 2010, the incidence of heart failure was 3.8 per 1000 person-years. The overall 5-year survival rate in the Swedish study was 48%, similar to the THIN study results, however between 2006 and 2010 there was a 19% decrease in mortality in both men and women ($P < 0.001$). A study of the Framingham cohort in the US also found an overall improvement in survival of 12% per decade between 1950 and 2000 (18).

Cancer survival rates in the UK have doubled in the last 40 years (19), yet in this study the outlook for patients with a first diagnostic label of heart failure has not improved over time. More research is needed into the disease trajectory and management of heart failure in order to explain the lack of improvement in survival over time despite the availability of evidence-based therapies shown to improve outcomes in clinical trials. This study did not examine medication use following a heart failure diagnosis and this is an area of planned further work. The treatment used in heart failure with reduced ejection fraction have not shown the same benefit in patients with heart failure with preserved ejection fraction which may partly explain lack of survival improvement. It was not possible to differentiate the type of heart failure due to lack of specific coding based on ejection fraction. However, as heart failure with preserved ejection fraction is increasingly recognised, and therefore more likely to be coded, further analysis by subtype may become possible.

The results of this study are estimates at a population level but discussions with the individual patient require tailored statistics and information in a form that patients and their carers can understand. Prognostic estimates vary depending on age and sex but also other co-morbidities, treatments and variables such as blood pressure and ejection fraction (20). Prognostic modelling to include other variables from the THIN database to more accurately predict survival for individual patients is also an area where further research is planned.

Supplementary Material

Supplementary data are available at Family Practice online.

Declaration

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