Sickness absence and disability pension among women with breast cancer: a population-based cohort study with a predictive model

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Abstract

Background: Women's return to work after diagnosis of breast cancer (BC) are becoming more prevalent. However, register-based national investigation on sickness absence (SA) and disability pension (DP) in BC women is lacking. Methods: The aim of the study was to explore SA and DP before and after a first BC diagnosis and the possibility to predict new cancer-related SA by using disease-related and sociodemographic factors. A longitudinal register study of the 3536 women in Sweden aged 19-64 with a first BC diagnosis in 2010 was conducted by linkage of multiple national registers. Particularly, information on SA and DP was obtained from the National Social Insurance Agency’s database. Descriptive statistics on SA and DP before and after the BC diagnosis were performed. The risk of being on sickness with a new SA spell due to BC or BC-related diagnoses was modeled using logistic regression. Results: The proportion of women with SA increased during the year following the BC diagnosis date and declined over the next two years to proportions before diagnosis. At the time of BC diagnosis, half of the women began a new SA spell >14 days with cancer, cancer-related, or mental diagnosis. Disease-related and sociodemographic factors including occupational sector, living area, age, cancer stage, educational level, and number of previous SA days showed statistical significance (p<0.05) in predicting a new SA around BC diagnosis. By using these factors, it was possible to correctly predict 67% of the new SA spell. Conclusion: SA among women with BC was elevated mainly in the first year after diagnosis. New SA following BC diagnosis can accurately be predicted.

Introduction

Breast cancer (BC) is a major health problem with 1.67 million new cases worldwide annually.[1] Due to early detection and better treatments, mortality has decreased, hence more knowledge is needed on potential adverse long-term social consequences of BC for the growing number of survivors.[2–4] About half of the women diagnosed with BC are of working age,[5, 6] thus, BC might imply sickness absence (SA) or even disability pension (DP) for many of them due to effects of BC and/or BC treatments. Studies indicate that many women with BC value paid work highly and want to continue working after diagnosis. [7–10] The detection of women with higher risk of SA is therefore important in order to facilitate part- or full-time (return to) work.

Studies of SA, DP, and return to work (RTW) among women with BC indicate that the majority RTW happens within two years.[5, 11, 12] Nevertheless, studies from Sweden and the Netherlands show that BC is associated with higher SA as long as five years after diagnosis[13–15] and that some are granted DP up to 10 years after diagnosis.[13, 15, 16] Advanced cancer stage,[11–13, 17] pre-diagnosis SA,[13–15, 18] comorbidity,[19] and several sociodemographic factors[12–15, 17–29] were negatively associated with RTW, alternatively positively with SA/DP, depending on the outcome used. However, studies of SA, DP, and RTW vary greatly in terms of study design, outcomes, selection of included women, national health insurance systems, and female employment frequency,[5, 11, 16, 19–21, 30–33] the latter implying different health-selection effects on outcomes. Sweden has a high female employment rate, also in
higher ages (>50 years old).[34] Thus, the healthy-selection effects on outcomes in Sweden are smaller which is an advantage when aiming at gaining knowledge on associations of BC with future SA/DP.

Most previous studies are hampered by short follow-ups, selected study populations, high drop-out, no information on DP or on pre-diagnosis SA/DP, or only have self-reported SA/DP.[19] Although elevated levels of post-diagnostic anxiety and depression are reported,[35, 36] detailed analyses of SA and DP due to mental diagnoses are seldom conducted. Thus, knowledge is limited on pre- and post-diagnosis diagnosis-specific SA and DP in women with BC, in nationwide population-based studies; knowledge that is needed to better understand the situation for women diagnosed with BC, as a basis to identify potential risk factors for SA/DP as a basis for preventive measures. Moreover, in healthcare, among employers, insurance organizations, and patients, there is a need to be able to predict future SA following a BC diagnosis in order to effectively direct prevention measures.[37] Using information on disease-related and sociodemographic factors is one way to achieve this goal.[38] To the best of our knowledge, this is the first study to test a model for prediction of SA following a first BC diagnosis.

The aims were: (a) to explore the annual prevalence of SA and DP due to cancer, other somatic, and/or mental diagnoses during the two years before and the three years following a BC diagnosis, and (b) to predict risk of a new SA spell following a BC diagnosis.

### Methods

A population-based longitudinal cohort study was performed.

We included all the 3536 women in Sweden, aged 19–64 who were diagnosed with a first malignant neoplasm of breast (International Classification of Diseases 10\textsuperscript{th} version (ICD–10) [39] code: C50) in 2010. Data were obtained from five nationwide registers as follows:

- The Board of Health and Welfare's Cancer Register (all BC cases 1958–2010, diagnosis date, type, T, N, and M classifications[40]), Patient Register (main diagnosis, dates of in- and specialized outpatient care 2008–2010), and Cause of Death Register (dates 2010–2013);

- Statistics Sweden's Longitudinal Integration Database for Health Insurance and Labor Market Studies (LISA) (age, educational level, marital status, family composition, birth country, occupational sector, geographical and type of living area in December 2009, emigration 2010–2012, not living in Sweden 2008 or 2009);

- National Social Insurance Agency’s Micro-data for Analyses of Social Insurance (MiDAS) (SA and DP benefits 2008–2013: dates, full- or part-time, main diagnosis).

Data were linked at individual level using the ten-digit personal identity numbers assigned to all residents in Sweden.
**SA and DP benefits in Sweden**

All people in Sweden ≥ 16 years, with an income from work or unemployment benefit, with reduced work capacity due to disease or injury can be granted SA benefit from the Social Insurance Agency. The employers usually provide reimbursement for the first 14 days of a SA spell, which is why we do not have information on most SA spells ≤ 14 days. From day 8, a medical certificate issued by the treating physician is required. All residents aged 19–64 can be granted with DP if having long-term or permanent work incapacity due to disease or injury. SA and DP can be granted for full-time (100%) or part-time (25, 50, or 75%) of ordinary working hours. SA benefits cover 80% and DP 64% of lost income, up to a certain level.

**Measures**

We investigated two types of outcomes; DP and SA (for spells >14 days). SA and DP days were transformed into net days; e.g., 2 days on half-time SA or DP was counted as one net day. SA and DP diagnoses were coded by the certifying physician who assessed the patient's condition and work capacity. Diagnoses were for some analyses classified into four categories: 1): BC (ICD10: C50), BC-related diagnoses (Z80, Z85, N61-N63), and other cancer (C00-D48), 2): mental diagnoses (F00-F99, Z73), 3): other diagnoses (all remaining ICD codes), and 4): missing information. The outcome in the predictive model was defined as starting a new SA spell >14 days due to one of the following SA diagnoses (C00-D48, Z80, Z85, N61-N63, F00-F99, or Z73) during the time-window of 14 days before to 29 days after the BC diagnosis. This time window was based on the frequencies of start of new SA spells in the full cohort, in relation to diagnosis date (T₀). For some women there was a delay before the diagnosis was included in the Cancer Register (even if the women were informed) and for others, treatment did not start until weeks later. The reason for including “diagnoses related to BC” and “other cancer diagnoses” in the predictive model was that sometimes a broader category of cancer diagnoses is given in the medical certificate. Mental diagnoses were also included in the predictive model as that a cancer diagnosis might lead to anxiety or depression.[42]

The included sociodemographic, disease-related, and comorbidity covariates (listed in Table 1) were selected for the predictive model based on previous findings regarding factors influencing SA and RTW. Missing information on educational level was coded as elementary school. Cancer-stage groups were assigned using the TNM Classification of Malignant Tumours[40] and categorized as: T0N0M0+stage 0+I, stage II, stage III+IV, and missing all TNM (with no T, N, or M information), respectively. When T, N, or M information was missing in one or two of the categories or classified as 'X' (assessment not possible), the value was set to 0. If more than one tumour was registered, with different diagnosis dates in 2010, the most advanced tumour was selected. The main ICD–10 diagnoses for healthcare were coded by the treating physicians. Healthcare due to uncomplicated delivery (O80) or not related to morbidity (e.g., screening) was excluded.
Statistical analyses

The mean number of SA and of DP net days/year, respectively, were calculated for all women, using the BC diagnosis date ($T_0$) as reference, for the two years before $T_0$ and three years after $T_0$ ($Y_{-2}$ to $Y_{+3}$). This was done for all SA and DP as well as for the four SA/DP diagnostic categories mentioned above. The annual numbers and proportions of women with SA/DP due to the different diagnoses were also calculated. The denominator used in these calculations varied somewhat over the years due to the exclusion of women (turning 65 years, emigration, or death).

In the predictive model regarding risk of new SA related to time of diagnosis, 2954 women were included. For those analyses we excluded the 521 women (14.7%) already on SA or on DP for full-time or nearly full-time (75–100%) at $T_0$. Additionally, 61 women were excluded due to lack of covariate information, or because of extreme values on some of the continuous variables, e.g., number of healthcare visits or inpatient days.

The risk of a new SA spell due to BC or related diagnoses, other cancer diagnoses, or mental diagnoses was modelled using multivariable logistic regression\cite{45, 46} with a logistic model formulated as follows:

$$\log\{p(y_i = 1)/p(y_i = 0)\} = x_i' \beta$$

where $y_i$ denotes the SA status of individual $i$, and $x_i$ is a vector of observed covariates. Natural cubic splines\cite{47–49} were used to model potentially nonlinear effects of continuous covariates. The five variables that were modelled using splines were: age and number of previous: SA days, DP days, outpatient healthcare visits, and inpatient days, respectively, in the two pre-diagnostic years. An optimal threshold $c$ was selected, such that predicting $SA = 1$ whenever the fitted probability was above $c$, minimized the sum of false positive (FP) and false negative (FN) and maximized the proportion of correctly classified observations. Also, the receiver operating characteristic (ROC) was calculated.

Results

Sociodemographic and diagnostic covariates are presented in Table 1 for the entire cohort ($N = 3536$) as well as for the group included in the modelling ($n = 2954$). About 40% of the women in both groups were $\geq 56$ years. The compositions of women regarding distribution of characteristics in the two groups were fairly similar, except for no DP days during the two pre-diagnostic years: 82% among all vs. 94% in the model, as expected due to the inclusion criteria in the modelling group. The majority had the earliest stages of BC. In the two pre-diagnosis years ($Y_{-1}$ and $Y_{-2}$), the majority had no SA days (81% vs. 83%), about half (56% vs. 53%) had at least one visit in specialized outpatient healthcare while few (12% vs. 8%) had at least one inpatient day. At BC diagnosis, 11.3% of the women were already on SA and 17.5% on DP.

Proportions of women with SA and/or DP
During the year after the BC diagnosis date \((Y_{+1})\), 28\% of the women had no SA >14 days (Table 2), while 67\% had SA due to cancer; nearly half of those (32\%) for >180 days. In the second year \((Y_{+2})\), 35\% of the women had at least some SA, regardless of SA diagnosis. For cancer SA diagnoses, the corresponding proportion was 25\% during \(Y_{+2}\). During \(Y_{+3}\), the corresponding proportions were 25\% and 12\%, respectively. The proportions with SA due to mental diagnoses did not vary much between the studied years (3\textendash 4\%). For SA due to other and missing diagnoses, the corresponding proportions were 8\textendash 11\% all years, except for \(Y_{+1}\) when it was 6\%. The proportion of women with DP ranged from 15 to 18\% during all the five studied years. During \(Y_{+1}\), 15\% of the women had neither SA nor DP. Before BC diagnosis, i.e., during \(Y_{-2}\) and \(Y_{-1}\), that proportion was 73\%. During \(Y_{+2}\) and \(Y_{+3}\), the corresponding proportions were 52\% and 62\%, respectively.

### Mean SA and DP days/year

During \(Y_{+1}\), the mean number of SA days was 121.3. This was significantly higher than the numbers before diagnosis; (6.7 in \(Y_{-1}\) and 9.0 in \(Y_{-2}\)) (Figure 1). During \(Y_{+1}\), 108.8 of the SA days were due to cancer. That number was lower already in \(Y_{+2}\), i.e., 26.6 days. In \(Y_{+3}\), it was 14.0 days. Mean number of DP days/year was about 50 before BC diagnosis. Due to older women already on DP in the year before \(T_0\) becoming 65 years of age, that number decreased to about 40 days/year in \(Y_{+2}\) and \(Y_{+3}\).

### New SA spell

For the 3015 women who did not have an ongoing SA nor DP of the extent of 75\textendash 100\% at the time of BC diagnoses \((T_0)\) Table 3 shows the numbers and percentages of women who had a first new SA spell in relation to \(T_0\). The same is shown for three specific SA diagnostic groups, i.e., cancer, mental diagnoses, and the other diagnoses (including missing), respectively. Of these women, 51\% had a first new SA spell in relation to \(T_0\), that is, the period that was studied in the predictive modelling. Of these SA spells, 95\% were due to cancer. In the following 30-day period, i.e., from 30 to 59 days after \(T_0\), another 20\% had a first SA spell, of which 96\% were with cancer. A little less than one fifth of the women (18\%) had no new SA spells from \(T_0\) until end of follow-up three years later. Nine women were granted DP during the period. That is, about 80\% of those at risk of a new SA spell following BC diagnosis, had such a spell in the first year \((Y_{+1})\), and the majority of them (70\%) in the first three months.

### Predictive model

Out of the variables (see Table 1) included in the multivariable logistic regression model for the risk of having a new SA spell >14 days (due to BC or related diagnoses, other cancer diagnoses, or mental diagnoses) in connection with a first BC diagnosis \((T_0)\), the following variables were statistically significant (p<0.05): occupational sector, living area, age, cancer stage, educational level, and number of
previous net SA days (listed according to predictive strength from high to low). In Figure 2, the receiver operating characteristic (ROC) curve is illustrated. The dot indicates the coordinates (FP, FN) corresponding to the selected value of threshold $c$. The predictive model could correctly predict 67% out of 2954 women, given the optimal threshold 0.54. Results are summarized in Table 4 where the area under the curve (AUC)\cite{50,51} is also reported.

**Discussion**

In this longitudinal population-based cohort study of all women in Sweden aged 19–64 years with a first BC diagnosis in 2010, the proportion with SA >14 days increased substantially in the post-diagnostic 12 months. Nevertheless, most women were not on SA for extensive times during the first year after diagnosis date (e.g., >180 net days) and about 17% were already on SA/DP due to other diagnoses when diagnosed with BC. In the third year after BC diagnosis date ($Y_{+3}$), the mean number of SA and DP days was low and 25% of the women had no SA at all. The variation in proportion of women with DP was minor during the five years being studied. The elevated proportion of women on SA and the higher number of SA days after BC diagnosis, compared to the years before BC diagnosis, was in SA due to cancer throughout follow-up, e.g., not due to mental diagnoses. Of the 3015 women not already on SA/DP at time of BC diagnosis, 51% had a new SA spell and for 98% of them the SA was due to a cancer diagnosis, cancer-related diagnosis, or mental diagnosis. Using a predictive model including disease-related and socio-demographic factors, 67% of the women could be correctly classified into having or not having a new SA spell.

**Strengths and limitations**

Data from a total population provided us with a rare opportunity to study SA and DP in connection with BC in all women of working age. Other strengths are the longitudinal cohort design, that all women fulfilling the inclusion criteria of a first BC diagnosis in a whole country could be included, not a sample; also that extensive microdata on morbidity and sociodemographic variables from several high-quality registers could be linked at individual level\cite{52–54} and that data were not self-reported, avoiding recall bias. The large cohort also allowed sub-group analyses of specific SA/DP diagnoses, not only the BC diagnosis. The latter circumstance makes the results and model more useful in practice. High female employment frequency, complete coverage of the public SA/DP insurances, and no dropouts make the internal validity of the study very strong. A further strength is that we were able to exclude women no longer at risk of SA/DP due to death, turning 65, or emigration during follow-up. Findings can be generalized to women with BC in countries with comparable employment frequencies and coverage of SA/DP benefits.

The validity of SA and DP diagnoses is sometimes discussed but seldom investigated. To the best of our knowledge, there is only one such study, regarding SA diagnoses, which found the validity to be acceptable\cite{42}. The validity of DP diagnoses is likely to be even higher, since DP is only granted after a
long process of medical evaluation.[41] However, the stigma associated with mental diagnoses[55] might imply an underestimation of SA/DP due to such diagnoses. Other limitations of our study are that we had no information on SA spells \(\leq 14\) days and only information about the first and main SA diagnosis of a SA spell. Further, we did not have information on cancer treatment, which is a factor that might attenuate the association between cancer stage and SA.[14, 15]

We were able to replicate previous findings that the prevalence of SA was considerably higher in the year post-BC diagnosis than before diagnosis, using a population-based nationwide cohort.[13, 14] In a previous Swedish population-based study of all women aged 20–65 with a first BC diagnosis in 2005,[13] almost the same proportion of women as here presented (71% vs. 72%) had some SA during the first 12 months following BC diagnosis. In the cohort from 2005, however, a larger proportion of women, i.e., 19%, had SA already in the year before BC diagnosis, compared to 11% in our 2010 cohort. Also, proportions on DP were slightly higher in the pre-diagnostic year \((Y_{-1})\) in the 2005 cohort compared to the current cohort, 20.6% vs. 17.5%. In the following two years, the decline in SA was somewhat faster in our 2010 cohort. This difference might imply an impact of the stricter Swedish SA/DP regulations implemented in 2008 [56, 57]. However, the hypothesis should be studied further using other study designs.

The higher prevalence of SA after BC diagnosis, compared to the pre-diagnostic period, was due to cancer only, not due to mental nor to other somatic diagnoses in neither our cohort nor the 2005 cohort used in the previous Swedish study from our group,[13] i.e., SA due to mental diagnoses did not increase after the BC diagnosis. This is noteworthy, as studies have reported higher risks of anxiety and depression after BC diagnosis.[35, 36] One possible explanation is that mental disorders in women with BC are not recognized by sickness certifying physicians.[58] Another explanation is that such mental disorders did not reduce work capacity to such a high level that SA was required or that the women, if needed, soon had adequate treatment for mental disorders—more knowledge is needed on this. It is also possible that mental diagnoses was stated as a secondary SA diagnosis or becoming a main SA diagnosis later during the SA spell [59] and thus not captured by our data.

Regarding the predictive model, our accuracy of 67% is considered below recommended levels for clinical use.[38, 60] Nevertheless, the results from the model are promising regarding the possibility to further develop a predictive model.

Our results highlight the importance of communicating to women with BC as well as to employers of the fact that most women with BC return to work rather quickly, in order to promote optimal work adjustments as soon as possible, especially for groups with an elevated risk of SA/DP.

**Conclusions**

In this population-based prospective cohort study we found that although BC and BC treatment can have large impacts on work capacity, not all women diagnosed with BC had extensive SA/DP even in the first 12 months after diagnosis. Moreover, it is possible to give a good prediction of which women with BC
who are at high risk of new SA. Our predictive model should be further developed to assist the RTW measures for people with cancer diagnosis in the future.

**List Of Abbreviations**

Breast cancer (BC); sickness absence (SA); disability pension (DP); return to work (RTW); International Classification of Diseases 10th version (ICD–10); Longitudinal Integration Database for Health Insurance and Labor Market Studies (LISA); Micro-data for Analyses of Social Insurance (MiDAS); false positive (FP); false negative (FN); receiver operating characteristic (ROC); area under the curve (AUC)

**Declarations**

*Ethics approval*

The project was approved by the Regional Ethical Review Board of Stockholm, Sweden and was conducted in accordance with the Declaration of Helsinki.

*Consent for publication*

Not applicable

*Availability of data and materials*

The data used in this study is administered by the Division of Insurance Medicine, Karolinska Institutet, and cannot be made publically. According to the General Data Protection Regulation, the Swedish law SFS 2018:218, the Swedish Data Protection Act, the Swedish Ethical Review Act, and the Public Access to Information and Secrecy Act, these type of sensitive data can only be made available, after legal review, for researchers who meet the criteria for access to this type of sensitive and confidential data. Readers may contact Professor Kristina Alexanderson (kristina.alexanderson@ki.se) regarding the data.

*Competing interests*

The authors declare that they have no competing interests.

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*Authors’ contributions*

Contribution ship statement: KA, PK, EM-R, and EF were responsible for the study concept and design. KA obtained the research funding. PK, PF conducted the data management and analysis. PK drafted the manuscript. All authors were involved in result interpretation and manuscript revision. LC finalized the
results and manuscript, and was responsible for manuscript submission. All authors had full access to all of the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis.

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**Tables**

Table 1. Characteristics of the study cohort and the sub-cohort for modelling
| Covariates                      | The whole cohort | The cohort used for modelling |
|---------------------------------|-----------------|------------------------------|
|                                 | Number | (%) | Number | (%) |
| **All**                         | 3536   | 100 | 2954   | 100 |
| **Age group**                   |        |     |        |     |
| 18-35                           | 122    | 3.5 | 111    | 3.8 |
| 36-45                           | 665    | 18.8| 610    | 20.7|
| 46-50                           | 583    | 16.5| 515    | 17.4|
| 51-55                           | 630    | 17.8| 525    | 17.8|
| 56-60                           | 844    | 23.9| 664    | 22.5|
| 61-63                           | 692    | 19.6| 529    | 17.9|
| **Country of birth**            |        |     |        |     |
| Sweden                          | 2950   | 83.4| 2509   | 84.9|
| Other country                   | 586    | 16.6| 445    | 15.1|
| **Educational level**           |        |     |        |     |
| Elementary school (≤9 years)    | 551    | 15.6| 378    | 12.8|
| High school (10-12)             | 1549   | 43.8| 1265   | 42.8|
| College/University (>12)        | 1436   | 40.6| 1311   | 44.4|
| **Geographic living area**      |        |     |        |     |
| North                           | 425    | 12.0| 344    | 11.7|
| Middle                          | 486    | 13.7| 403    | 13.6|
| Stockholm                       | 823    | 23.3| 710    | 24.0|
| West                            | 934    | 26.4| 767    | 26.0|
| South                           | 868    | 24.6| 730    | 24.7|
| **Type of living area**         |        |     |        |     |
| Larger cities                   | 1376   | 38.9| 1172   | 39.7|
| Medium cities                   | 1252   | 35.4| 1051   | 35.6|
| More rural areas                | 908    | 25.7| 731    | 24.8|
| **Family composition**          |        |     |        |     |
| Married/cohab., no child at home| 1042   | 29.5| 852    | 28.8|
| Married/cohab., child at home   | 1235   | 34.9| 1129   | 38.2|
| Single, no child at home        | 885    | 25.0| 656    | 22.2|
| Single, child at home           | 374    | 10.6| 317    | 10.7|
| **Marital status**              |        |     |        |     |
| Unmarried, divorced, widow      | 1523   | 43.1| 1228   | 41.6|
| Married, registered partnership | 2013   | 56.9| 1726   | 58.4|
| **Occupational sector**         |        |     |        |     |
| Not in paid work/no information | 810    | 22.9| 379    | 12.8|
| Public                          | 1409   | 39.9| 1330   | 45.0|
| Private                         | 1317   | 37.3| 1245   | 42.2|
| **Cancer stage**                |        |     |        |     |
| Missing all T, N, M             | 25     | 0.7 |        |     |
| TON0M0 + Stage 0+I              | 2120   | 60.0| 1799   | 60.9|
| Stage II                        | 1162   | 32.9| 990    | 33.5|
| Stage III+IV                    | 229    | 6.5 | 165    | 5.6 |
| **Previous SA, net days**       |        |     |        |     |
| No previous SA                  | 2870   | 81.2| 2444   | 82.7|
| 0.25-90                         | 485    | 13.7| 417    | 14.1|
| ≥90                             | 181    | 5.1 | 93     | 3.2 |
| **Previous SA, diagnoses**      |        |     |        |     |
| Mental diagnoses                | 161    | 4.6 | 124    | 4.2 |
| Previous DP net days¹ | 0      | 2908   | 82.2  | 2771   | 93.8  |
|-----------------------|--------|--------|-------|--------|-------|
| 0.25-365              | 180    | 5.1    | 153   | 5.2    | 365   |
| ≥365                  | 448    | 12.7   | 30    | 1.0    | 12.7  |
| Previous DP diagnoses¹ |        |        |       |        |       |
| Mental diagnoses      | 181    | 5.1    | 47    | 1.6    |       |
| Other diagnoses       | 466    | 13.2   | 137   | 4.6    |       |
| Previous visits in specialized outpatient care¹ |        |        |       |        |       |
| 0                     | 1549   | 43.8   | 1390  | 47.1   |       |
| 1-2 visits            | 1039   | 29.4   | 898   | 30.4   |       |
| ≥3 visits             | 948    | 26.8   | 666   | 22.6   |       |
| Previous visits in outpatient care, diagnoses¹ |        |        |       |        |       |
| Mental diagnoses      | 166    | 4.7    | 72    | 2.4    |       |
| Other diagnoses       | 1825   | 51.6   | 1441  | 48.8   |       |
| Previous inpatient care, days¹ |        |        |       |        |       |
| 0                     | 3123   | 88.3   | 2709  | 91.7   |       |
| 1-14 days             | 354    | 10.0   | 235   | 8.0    |       |
| ≥14 days              | 59     | 1.7    | 10    | 0.3    |       |
| Previous inpatient care, diagnoses¹ |        |        |       |        |       |
| Mental diagnoses      | 32     | 0.9    | 11    | 0.4    |       |
| Other diagnoses       | 391    | 11.1   | 238   | 8.1    |       |

The table included sociodemographic factors, cancer stage, and previous sickness absence (SA), disability pension (DP), and healthcare (n and %) for the cohort of all women in Sweden <65 years with a first breast cancer diagnosis in 2010 as well as for those included in the logistic regression used to build a predictive model.

¹ Previous = in the period 730 – 15 days before the BC diagnosis date

² The first 14 days of SA spells are not included.

³ That is, those at risk for a new SA spell

Table 2. Annual sickness absence (SA) or disability pension (DP) net days in women (number and percentage).
| SA/DP/ diagnoses | Mean net days per women | Y₂ n (%) | Y₁ n (%) | Y₁+1 n (%) | Y₁+2 n (%) | Y₁+3 n (%) |
|------------------|------------------------|---------|---------|-----------|-----------|-----------|
| SA¹               |                        |         |         |           |           |           |
| All              | 0                      | 3115 (88.4) | 3134 (88.7) | 978 (27.7) | 2282 (65.3) | 2401 (75.2) |
| >0-30            | 198 (5.6)              | 206 (5.8)  | 485 (13.7) | 422 (12.1) | 324 (10.2)  |
| >30-90           | 98 (2.8)               | 105 (3.0)  | 440 (12.4) | 297 (8.5)  | 165 (5.2)   |
| >90-180          | 55 (1.6)               | 50 (1.4)   | 366 (10.4) | 213 (6.1)  | 133 (4.2)   |
| >180             | 56 (1.6)               | 39 (1.1)   | 1267 (35.8) | 278 (8.0)  | 168 (5.3)   |
| Cancer²          | 0                      | 3509 (99.6) | 3506 (99.2) | 1165 (32.9) | 2610 (74.7) | 2801 (87.8) |
| >0-30            | ≤8                     | 19 (0.5)   | 486 (13.7) | 306 (8.8)  | 124 (3.9)   |
| >30-90           | ≤8                     | ≤8         | 408 (11.5) | 224 (6.4)  | 89 (2.8)    |
| >90-180          | ≤8                     | ≤8         | 337 (9.5)  | 151 (4.3)  | 75 (2.4)    |
| >180             | ≤8                     | ≤8         | 1140 (32.2) | 201 (5.8)  | 102 (3.2)   |
| Mental³          | 0                      | 3423 (97.2) | 3435 (97.2) | 3430 (97.0) | 3354 (96.0) | 3074 (96.3) |
| >0-30            | 42 (1.2)               | 48 (1.4)   | 25 (0.7)   | 44 (1.3)   | 50 (1.6)    |
| >30-90           | 22 (0.6)               | 30 (0.8)   | 22 (0.6)   | 43 (1.2)   | 21 (0.7)    |
| >90-180          | 16 (0.5)               | 11 (0.3)   | 15 (0.4)   | 19 (0.5)   | 21 (0.7)    |
| >180             | 19 (0.5)               | 10 (0.3)   | 44 (1.2)   | 32 (0.9)   | 25 (0.8)    |
| Other⁴           | 0                      | 3219 (91.4) | 3246 (91.9) | 3331 (94.2) | 3208 (91.9) | 2848 (89.3) |
| >0-30            | 158 (4.5)              | 153 (4.3)  | 85 (2.4)   | 151 (4.3)  | 205 (6.4)   |
| >30-90           | 74 (2.1)               | 73 (2.1)   | 29 (0.8)   | 62 (1.8)   | 63 (2.0)    |
| >90-180          | 37 (1.1)               | 36 (1.0)   | 16 (0.5)   | 30 (0.9)   | 35 (1.1)    |
| >180             | 34 (1.0)               | 26 (0.7)   | 75 (2.1)   | 41 (1.2)   | 40 (1.3)    |
| DP               | All                    | >0       | 613 (17.4) | 619 (17.5) | 606 (17.1) | 553 (15.8) | 492 (15.4) |
|                  | Cancer                 | >0       | ≤8         | ≤8         | 12 (0.3)   | 13 (0.4)   | 21 (0.7)   |
|                  | Mental                 | >0       | 171 (4.9)  | 170 (4.8)  | 160 (4.5)  | 148 (4.2)  | 135 (4.2)  |
|                  | Other²                 | >0       | 445 (12.6) | 449 (12.7) | 436 (12.3) | 393 (11.3) | 338 (10.6) |
| No SA/DP         | >0                     | 2578 (73.2) | 2579 (73.0) | 543 (15.4) | 1812 (51.9) | 1990 (62.4) |
| Not included⁵    | 14                     | ≤8         | 0         | 44         | 345        |

Reasons for not being included the specific year

| Reason                                      | 2010   |
|---------------------------------------------|--------|
| >65 years of age                            | 248    |
| Death ≤65 year of age                       | 43     |
| Not living in Sweden and ≤65 years          | 14     |

Included in the table were all women in Sweden <65 years with a first breast cancer diagnosis in 2010 (N=3536), during the two years before and three years after the breast cancer diagnosis date, presented for all SA/DP as well as by three categories of SA/DP diagnoses. Also, the number of women not included in the respective year are presented by reason for not being included.

¹ The first 14 days of SA spells are excluded.

² ICD-codes: C00-D48, Z80, Z85, N61-N63

³ ICD-codes: F00-F99, Z73
In the group “Other diagnoses”, also SA/DP with missing information on diagnosis were included.

Women who turned 65, died, or emigrated were included up to and including the year of the event.

Table 3. Number and percentages of women with a new sickness absence (SA) spell (>14 days).

| Days relative to \(T_0\) | Number (% of all women, column %) | Cancer (row%) | Mental (row%) | Other diagnoses (row%) |
|--------------------------|-----------------------------------|---------------|---------------|------------------------|
| No new SA spell before \(T_0\) to end of follow-up \(^3,4\) | 536 (17.8) | - | - | - |
| 14 days before \(T_0\) to 29 days after \(T_0\) | 1535 (50.9) | 95 | 3 | 2 |
| 30-59 days after \(T_0\) | 599 (19.9) | 96 | 1 | 3 |
| 60-89 days after \(T_0\) | 152 (5.0) | 93 | 3 | 4 |
| 90-119 days after \(T_0\) | 78 (2.6) | 92 | 1 | 6 |
| 120-179 days after \(T_0\) | 48 (1.6) | 90 | 0 | 10 |
| 180-364 days after \(T_0\) | 19 (0.6) | 68 | 0 | 32 |
| 365-729 days after \(T_0\) | 21 (0.7) | 57 | 14 | 29 |
| ≥730 days after \(T_0\) \(^4\) | 27 (0.9) | 18 | 15 | 67 |

The included individuals were 3015 women <65 years related to date of a first breast cancer diagnosis in 2010 (\(T_0\)), during the following three years; all SA and diagnosis-specific SA (cancer, mental, or others).

\(^1\) Women already (nearly) full-time (75-100%) SA or disability pension (DP) at \(T_0\) were not included.

\(^2\) ICD-codes: C00-D48, Z80, Z85, N61-N63

\(^3\) Nine of these women were granted DP during follow-up

Table 4. Values of the predictive model.

| Values | |
|--------|-------|
| Threshold (\(c\)) | 0.54 |
| % False positive | 15 |
| % False negative | 18 |
| % Correct | 67 |
| \textit{Area under the curve (AUC)} | 0.73 |

The values included in the table were: optimal threshold (\(c\)), proportion of false positive (FP) and false negative (FN), proportion of correctly classified observations, and area under the curve (AUC) in the analyses of women in Sweden <65 years with a first breast cancer diagnosis in 2010 (\(N=2954\)).

\(^1\) Women not at risk of a new SA spell at date of breast cancer diagnosis were not included (that is, those already on (nearly) full-time SA or DP (75-100%).
Figure 1

Figure 1. Mean annual number of sickness absence (SA) and disability pension (DP) net days by diagnosis. Included in the figure: all women in Sweden <65 years, with a first breast cancer diagnosis in 2010 (N=3536), in the two years before and three years after date of diagnosis (T0), respectively. Included in the denominator each year: women <65, alive, and living in Sweden.
Figure 2

Figure 2. The receiver operating characteristic (ROC) curve. Proportion of false positive/negative at different values of c in the cohort of women in Sweden <65 years with a first breast cancer diagnosis in 2010. The dot corresponds to the optimal choice of threshold c.