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ORIGINAL RESEARCH

Predicting the duration of sickness absence spells due to back pain: a population-based study from Sweden

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ABSTRACT

Objectives We aimed to develop and validate a prediction model for the duration of sickness absence (SA) spells due to back pain (International Statistical Classification of Diseases and Related Health Problems 10th Revision: M54), using Swedish nationwide register microdata.

Methods Information on all new SA spells >14 days from 1 January 2010 to 30 June 2012 and on possible predictors were obtained. The duration of SA was predicted by using piecewise constant hazard models. Nine predictors were selected for the final model based on a priori decision and log-likelihood loss. The final model was estimated in a random sample of 70% of the SA spells and later validated in the remaining 30%.

Results Overall, 64 048 SA spells due to back pain were identified during the 2.5 years; 74% lasted ≤90 days, and 9% >365 days. The predictors included in the final model were age, sex, geographical region, employment status, multimorbidity, SA extent at the start of the spell, initiation of SA spell in primary healthcare and number of SA days and specialised outpatient healthcare visits from the preceding year. The overall c-statistic (0.547, 95% CI 0.542 to 0.552) suggested a low discriminatory capacity at the individual level. The c-statistic was 0.643 (95% CI 0.634 to 0.652) to predict >90 days spells, 0.686 (95% CI 0.676 to 0.697) to predict >180 spells and 0.753 (95% CI 0.740 to 0.766) to predict >365 days spells.

Conclusions The model discriminates SA spells >365 days from shorter SA spells with good discriminatory accuracy.

INTRODUCTION

Sickness absence (SA) due to poor health is a major public health problem.^{1–3} Musculoskeletal diagnoses (MSD) account for most of the SA spells in Europe, although SA due to mental disorders is growing.^{4,5} The costs of SA due to MSD incurred by individuals, employers and the society at large are high.^{1,2} Studies showed that low back pain, although not life-threatening, is a top contributing factor to the burden of MSD.⁶ Back pain is also a common reason for seeing a physician.^{7,8} So far, however, no consensus has been reached on prognosis and management of low back pain.⁹ Instead, many clinical practice guidelines for treatment and care of low back pain have been proposed.¹⁰ Regarding work incapacity, predictors of SA for back pain are well-known.^{11,12} The previous studies, however, have not focused on prediction of the duration

Key messages

What is already known about this subject?

- Predictors of sickness absence (SA) for back pain are well-known, but the previous studies have not focused on prediction of the duration of SA spells due to back pain.

What are the new findings?

- This study developed and validated a clinically implementable, pragmatic prediction model for the duration of SA spells >14 days due to back pain (M54) for the whole working-age population in Sweden.
- The model discriminates the long-term SA spells (>365 days) from shorter SA spells with good discriminatory accuracy.

How might this impact on policy or clinical practice in the foreseeable future?

- The prediction model can be applied in Sweden, and perhaps have applicability to other Nordic countries with similar welfare systems for social insurance.
- The results of this work are currently implemented in pilot tests in primary healthcare settings to help general practitioners to identify which of their sickness absent patients have high risk for long-term sickness absence.

of SA spells due to back pain.¹³ Effective prediction methods for the duration of SA spells due to low back pain can improve possibilities to identify patients at risk of long-term SA.

Several studies have developed and validated decision support tools to predict future work incapacity, whether ending in SA or not, in general or related to specific diagnoses.^{14–18} Most of these studies were limited by low sample sizes,^{19,20} specific study populations,^{17,19} short follow-up periods^{17,21} or lack of information on SA diagnosis.^{17–19} The grounds for the selection of predictors in those prediction models have varied from using theoretical frameworks, that is, the job-demand-control model,¹⁷ mathematical modelling of survey data,¹⁴ to selection based on literature.¹⁹ Only one previous study developed a model to predict duration of SA among those already on SA—with focus on SA due to stress-related diagnoses.²² Such prediction models are needed, especially in primary healthcare where such consultations are common among general



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practitioners.^{23 24} In this study, we developed and validated a prediction model for the duration of SA spells due to back pain with population data from Swedish nationwide registers.

METHODS

A prospective cohort study using register microdata was conducted including all new 64 048 SA spells in Sweden >14 days that were initiated in the 2.5-year period of 1 January 2010 to 20 June 2012 with an International Statistical Classification of Diseases and Related Health Problems 10th Revision (ICD-10) diagnosis code 'M54', that is, dorsalgia/back pain.²⁵

Register linkage

The SA spells were identified by use of the Swedish Social Insurance Agency's Micro Data for Analysis of the Social Insurance (MiDAS) register, including information on all SA spells with benefits from the Agency. Information from MiDAS (dates, diagnoses and extent of SA and disability pension (DP) and employment status when the SA spell begun) was linked with the following four nationwide registers at individual level by the use of the unique personal identity number of all residents in Sweden²⁶:

- ▶ The Longitudinal Integration Database for Health Insurance and Labour Market Studies (Swedish abbreviation LISA) from Statistics Sweden, regarding information on sociodemographic factors (age, sex, country of birth, educational level, occupational sector, marital status, family composition, emigration year, geographical area and type of living area for the calendar year preceding the start date of the index SA spell).²⁷
- ▶ From the National Board of Health and Welfare three registers were used regarding information for 2009–2012; the Swedish inpatient and specialised outpatient registers²⁸ (dates and diagnoses of inpatient and specialised outpatient healthcare); the Prescribed Drug register²⁹ (dates and the Anatomical Therapeutic Chemical (ATC) codes for dispensed prescribed drugs); and the Cause of Death register (dates of deaths).³⁰

Social insurance system in Sweden

In Sweden, all residents aged 16 years or above, with an income from work or unemployment benefits are eligible for SA benefits, if the work capacity of the individual is reduced because of a disease or an injury.⁴ For most employed, the first 14 days of a SA spell are reimbursed by the employer, otherwise from the Social Insurance Agency. From day 8, a sickness certificate from a physician is required. For individuals aged 19–64 years, DP can be granted, if their work capacity is long-term or permanently reduced because of a disease or an injury. Both SA and DP can be granted for 100%, 75%, 50% or 25% of ordinary working hours. This means that people on part-time DP, for example, for 25%, at the same time can have a part-time SA spell.

Information on SA spells

The *duration* of each SA spell was calculated from the first until the last day of the SA spell, except when it ended with DP. In this case, the duration of the spell was set to 1000 days (n=1613). SA spells >1000 days were fixed at 1000 days (n=148) to control for extreme outliers in our analysis as has been done before.²²

Predictors

A decision was made that for clinical implementation a maximum of nine variables could be included in the model—it was assumed

that more variables would take too much time for the general practitioner to include in the algorithm during the consultation. Of these, three were to be included as they were routinely collected information in the medical records (sex, age, and geographical area) and the other six which were simple enough for the certifying physician to ask the patient about. Initially, the number of possible predictors was very large (130) as has been described in detail earlier.^{22 31 32} Many of these predictors were redundant or partially redundant and/or highly collinear as shown in an earlier study based on the same data sources.²²

There were very few missing values for the different possible predictors. Those with missing information on country of birth (0.05% of the SA spells) were coded as 'non-EU country' and missing information on educational level (0.3% of the SA spells) were coded as education at elementary level. As described earlier,²² several analyses were conducted with the different variables, to check the predictive value, especially comparing hierarchically related variables referring to specific diagnoses versus their non-diagnosis specific pair. The general variable performed at least as well, but often better than the specific variable. Hence, the general variables were kept for further analyses. The general variables had also an advantage because they could facilitate implementation of the model in clinical settings, as information for its application was to be provided by the patient during the consultation. Last, no difference in predictive value existed if the number of previous SA days were counted during 1–365 days or the 366–730 days period before the start date of the index SA spell. That was also the case regarding hospitalisation and specialised outpatient visits in the two different previous years. Therefore, we choose the most recent period to facilitate the implementation of the model. Furthermore, recall bias, when using information from the patient, was also likely to be smaller if only the recent year had to be considered. Hence, this selection process resulted 14 variables, which were independent predictors for the duration of the SA spell. They are listed below:

Possible predictors regarding the day when the SA spell started:

- ▶ Age: grouped as 18–34, 35–40, 42–50, 51–57 and 58–64 years.
- ▶ Sex: woman, man.
- ▶ Geographical regions: Northern, Middle, Western and Southern Sweden, and Stockholm/Gotland.
- ▶ Country of birth: Sweden, other Nordic, other EU25 and rest of the world.
- ▶ Family composition was described with a four-categories composite variable constrained from the following possible situations: living with or without a partner and living with or without children (aged <18 years).
- ▶ Educational level: categorised as elementary (≤ 9 years), high school (10–12 years) and university/college (>12 years).
- ▶ Employment status: in paid work, on unemployment benefit or at parental leave or student benefit.
- ▶ SA extent: 25%, 50%, 75% or full-time (100%).
- ▶ Partial DP: yes, no.
- ▶ Whether the SA spell was initiated in primary healthcare or not was assessed by having had any inpatient or specialised outpatient healthcare during the period of 4 days before and 8 days after the start of the SA spell and categorised as 'yes' or 'no'.

Possible predictors based on the 365 days preceding the start date of the SA spell:

- ▶ The number of SA days: 0, 1–90, 91–180 or >180 days.
- ▶ The number of days spent in inpatient healthcare: 0, 1–2 or >2 days.

Table 1 Characteristics of all the new sickness absence (SA) spells due to back pain (M54) >14 days during a 2.5-year period

Characteristics	All	Training data	Validation data
Number of SA spells	64 048	44 833	19 215
Number of individuals	56 856	41 146	18 521
Duration of the SA spell	n (%)		
≤90 days	47 107 (73.5)	33 011 (73.6)	14 096 (73.3)
>90 and ≤180 days	6563 (10.2)	4563 (10.2)	2000 (10.4)
>180 and ≤365 days	4495 (7.0)	3126 (7.0)	1369 (7.1)
>365 and <1000 days	4122 (6.4)	2898 (6.5)	1224 (6.4)
≥1000 days	1761 (2.7)	1235 (2.8)	526 (2.7)
Situation at the end of the SA spell*	n (%)		
Death	142 (0.2)	99 (0.2)	43 (0.2)
Disability pension	1653 (2.6)	1160 (2.6)	493 (2.6)
The individual was >65 years of age†	339 (0.5)	236 (0.5)	103 (0.5)
Returning to work or other activity	61 882 (96.7)	43 316 (96.7)	18 576 (96.7)
Other reasons‡	32 (<0.1)	22 (<0.1)	10 (<0.1)
General descriptive statistics for the duration of the SA spells	Days		
Median	37	37	37
IQR	22–98	22–98	22–99
Mean (SD)	130.3±233.2	130.3±233.7	130.1±232.2

*As only one person emigrated, it is not being shown in the table.

†Sixty-five years is the general age for retirement in Sweden.

‡Shift to other social benefits such as parental leave or unemployment.

- ▶ The number of specialised outpatient visits: 0, 1–2 or >2 visits.
- ▶ Multimorbidity was defined, based on dispensed prescribed medication. At least one purchase for a minimum of three different ATC groups were considered as having multimorbidity.

Information about all these items was assumed to be easily retrieved during a patient-physician consultation. To achieve the a priori decided three routinely available and six patient-physician question predictors for the model, we selected the 6 variables with the highest predictive values to the final model from this set of 14 variables through additional analyses.

Statistical analysis

The dataset was randomly split into a training dataset (70% of the SA spells) for model development and a validation dataset (30% of the SA spells) for external validation as has been done before.²² Descriptive statistics are presented in table 1 for the characteristics of the SA spells due to back pain and for the distribution of the predictors in the training and the validation dataset. The duration of the SA spells was used as outcome in the analyses. Piecewise constant hazards regression models were fitted with 20 time intervals of empirical quantiles to predict the duration of the SA spells (see online supplementary tables 1-2).

As mentioned above, to facilitate implementation to clinical practice, three routinely available predictors, that is, age, sex and geographical region, were included in the model. Further predictors were selected based on the loss of the log-likelihood when they were removed one-by-one from the full model. Those six that resulted in the largest decreases in the log-likelihood were included in the final model, beside the three preselected

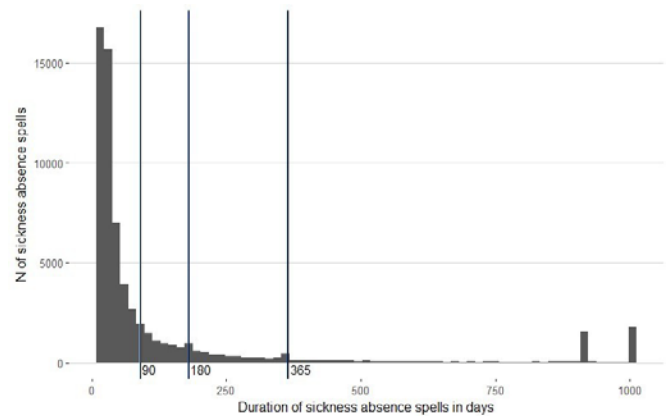


Figure 1 Histogram of the distribution of the sickness absence (SA) spells. Vertical lines indicate the cut-off at duration of 90, 180 and 365 days.

predictors. Both Akaike information criterion (AIC) and Bayesian information criterion (BIC) were used to compare the goodness-of-fit of the final model to the full model.

Quantile-quantile plot of the survival probabilities at observed time of event T were used for calibration. The quantiles of the predicted probability that an SA will be longer than its observed length $P(T > T_{\text{obs}})$ was compared with the quantiles of a uniform distribution to visually test the hypothesis that $P(T > T_{\text{obs}})$ is uniformly distributed, that is, quantile-quantile plots were used to assess if the model is correctly specified. The overall discriminatory capacity was assessed using the c-statistic.^{33 34} The CIs for the overall c-statistics were obtained by using bootstrap resampling ($n=1000$). The observed versus predicted survival probabilities at predefined SA durations (90, 180 and 365) were plotted. Receiver operating characteristics (ROC) curves and their corresponding c-statistics with 95% CI, calculated by the DeLong approach,³⁵ were used to evaluate discriminatory accuracy for predicting duration >90, >180 and >365 days, respectively.

R V.3.4.3³⁶ was used for statistical analysis and graphics (packages ‘pch’, ‘pROC’, ‘e1071’, ‘Hmisc’, ‘ggplot2’ and ‘pch’),³⁷ and SAS V.9.4 for the data management.

RESULTS

Characteristics of the 64 048 SA spells are shown in table 1. Overall, 73.5% of the spells lasted ≤90 days, while 9.2% ($n=5883$) became >365 days (table 1 and figure 1). The median duration of the SA spells was 37 days (the IQR 22–98 days). Most of the spells (96.7%) ended due to return to work or to other activity (eg, studies or parental leave), while 2.6% of the SA spells continued as DP (table 1).

The distribution of the baseline predictors is presented in table 2. Employment status, multimorbidity, SA extent at the start of the spell, whether the SA spell was initialised in primary healthcare as well as the gross number of SA days and number of specialised outpatient clinic visits during the 365 days preceding the start date of the SA spell, were the six selected predictors that had the largest log-likelihoods loss, and therefore were included in the final model together with the three routinely available variables: age, sex, and geographical region (table 3).

Both the AIC (453 078 vs 453 089) and BIC (458 664 vs 456 737) were lower for the final compared with the full model. The log-likelihood was only slightly higher for the final than for the full model (−226 119 and −225 904 with 420 and 640 free

Table 2 Tabulation of possible baseline predictors for the training and test data

Predictor	Training data (n of spells, %)	Test data (n of spells, %)
Age (years)		
18–30	5421 (12.1)	2339 (12.2)
31–40	9626 (21.5)	4074 (21.2)
41–50	13 145 (29.3)	5624 (29.3)
51–57	8762 (19.5)	3811 (19.8)
58–64	7876 (17.6)	3367 (17.5)
Sex		
Women	24 791 (55.3)	10 647 (55.4)
Men	20 042 (44.7)	8568 (44.6)
Geographical region		
Northern	6306 (14.1)	2688 (14.0)
Middle	6651 (14.8)	2741 (14.3)
Stockholm/Gotland	8750 (19.5)	3716 (19.3)
Western	13 264 (29.6)	5776 (30.1)
Southern	9862 (22.0)	4294 (22.3)
Educational level		
Elementary (≤9)	8779 (19.6)	3783 (19.7)
High school (10–12)	9110 (20.3)	3856 (20.1)
University/college (>12)	26 944 (60.0)	11 576 (60.2)
Family composition		
Living with partner and children	18 015 (40.2)	7927 (41.3)
Single with children at home	4776 (10.7)	1946 (10.1)
Living with partner without children	7302 (16.3)	3112 (16.2)
Single without children	14 740 (32.9)	6230 (32.4)
Country of birth		
Sweden	34 379 (76.7)	14 730 (76.7)
Other Nordic	1551 (3.5)	680 (3.5)
Other EU25	1262 (2.8)	525 (2.8)
Rest of the world	7641 (17.0)	3260 (17.0)
Occupational status		
Employed	41 711 (93.0)	17 831 (92.8)
Unemployed	2463 (5.5)	1097 (5.7)
Parental leave/student	659 (1.5)	287 (1.5)
Part-time disability pension at start of the SA spell		
Yes	2781 (6.2)	1144 (6.0)
No	42 052 (93.8)	18 071 (94.0)
SA extent at the start of the SA spell		
100%	36 491 (81.4)	15 698 (81.7)
75%	1154 (25.7)	476 (2.5)
50%	5943 (13.2)	2504 (13.0)
25%	1245 (27.8)	537 (2.8)
Multimorbidity*		
Yes	11 440 (25.5)	4861 (25.3)
No	33 393 (74.5)	14 354 (74.7)
SA days in the 365 days before the SA spell		
0	31 161 (70.5)	13 412 (69.8)
>0 and ≤90	10 612 (23.7)	4516 (23.5)
>90 and ≤180	1718 (3.8)	704 (3.7)
>180	1342 (3.0)	583 (3.0)
Number of days in inpatient care during the 365 days before the SA spell†		
0	40 935 (91.3)	17 586 (91.5)
1–2	16 463 (3.7)	711 (3.7)
>2	2255 (5.1)	918 (4.8)
Number of visits in specialised outpatient care during the 365 days before the SA spell		

continued

Table 2 continued

Predictor	Training data (n of spells, %)	Test data (n of spells, %)
0	24 072 (53.7)	10 370 (54.0)
1–2	13 795 (30.8)	5821 (30.3)
>2	6966 (15.5)	3024 (15.7)
The SA spell was initiated in primary healthcare		
Yes	38 795 (85.4)	16 508 (85.9)
No	6525 (14.6)	2707 (14.1)

*Any inpatient or specialised outpatient healthcare during the period of 4 days before and 8 days after the start of the SA spell.

†Excluding codes O80 and Z00-Z99 except Z73.0 at least three different types of medication (different ATC codes) at least once.

ATC, Anatomical Therapeutic Chemical; SA, sickness absence.

parameters, respectively). The probability integral transform value of the model was basically uniform, indicating a good calibration and specification of the final model. The overall c-statistic was rather low, 0.547 (95% CI 0.542 to 0.552) suggesting a low discriminatory capacity at the individual level. Binary predictions of risk of SA duration showed reasonable performance for SA spells >90, >180 days, and good performance for SA spells >356 days. The c-statistics were 0.643 (95% CI 0.634 to 0.652), 0.686 (95% CI 0.676 to 0.697) and 0.753 (95% CI 0.740 to 0.766), respectively (figure 2).

DISCUSSION

We developed and validated a prediction model for the duration of SA spells >14 days due to back pain (M54) to be easily implementable in primary healthcare. Analyses were based on all SA spells due to M54 in Sweden during a period of 2.5 years. Age, sex, geographical region, employment status, multimorbidity, SA extent at the start of the spell, whether the SA was initialised in primary healthcare settings as well as the number of SA days and visits in specialised outpatient healthcare during the preceding year were included into the final model as predictors. The overall discriminatory capacity to predict SA duration at an

Table 3 Decrease in likelihood of the 14 predictors of the duration of SA spells due to M54

Predictors	Loss of log-likelihood for the variable	P value	100×pseudo-R ²
Employment status	1037	<0.001	0.46
Gross SA days in the 365 previous days before start of the SA spell	542	<0.001	0.24
Age	375	<0.001	0.17
SA extent at the start of the SA spell	350	<0.001	0.15
Multimorbidity*	108	<0.001	0.05
Geographical region	96	<0.001	0.04
The SA spell was initialised at primary healthcare	74	<0.001	0.03
Number of specialised outpatient care visits in the 365 previous days	68	<0.001	0.03
Country of birth	65	<0.001	0.03
Partial disability pension at start of SA spell	49	<0.001	0.02
Family status	47	0.004	0.02
Educational level	37	0.001	0.02
Sex	32	<0.001	0.01
Number of days spent in inpatient care in the 365 previous days	13	0.961	0.01

*Defined as at least three different types of medication (different ATC codes) at least once. ATC, Anatomical Therapeutic Chemical; SA, sickness absence.

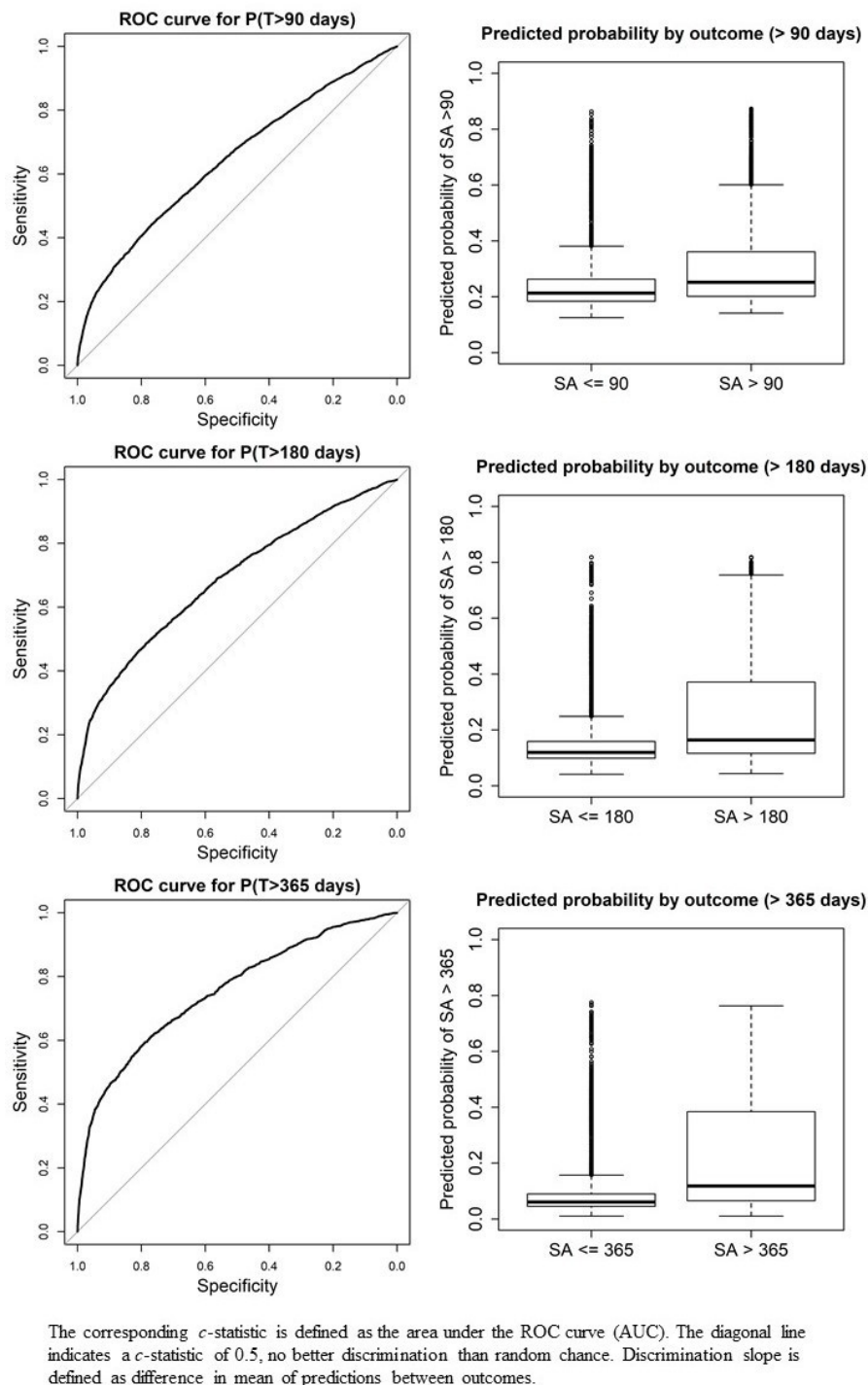


Figure 2 Receiver operating characteristics (ROC) curves and box plots of predicting duration of sickness absence (SA) for outcomes of >90, >180 and >365 days.

individual level was poor, but the prediction model was good at discriminating SA>365 days from shorter SA spells.

To the best of our knowledge, this is the first time that a prediction model has been developed to predict the duration of SA spells due to back pain. To be able to predict the risk that an ongoing SA spell due to back pain will be long, is an issue of great clinical importance, especially in primary healthcare. To date, to the best of our knowledge, the previous two studies using prediction models instead aimed at predicting which people, among those yet not on SA, will become long-term sickness absent due

to back pain or due to any diagnoses,^{19,38} hence not being comparable to this study with prediction of duration of SA spells.

Several studies have identified previous (long-term) SA as a strong risk factor for future SA¹⁷⁻¹⁹; we also found this to be a predictor and included it in our model. We also accounted for other factors such as age, sex, educational level and family composition based on their well-known importance for SA.^{19,32} As our major aim was to develop a minimalistic model that is easily implementable in primary healthcare settings with a reasonable model performance, we tested many predictors available in the

registers and those which were used in the final model were such that they could be assessed with great accuracy during the short time of a patient-physician consultation. Our model predicted long-term SA spells, that is, >365 days with good discriminatory accuracy, which may be useful to identify the patients on SA due to back pain who are at high risk of long-term SA or even DP, in order to take preventive actions.

The duration of SA spells is an extremely complex phenomenon, influenced by many factors at different structural levels, especially SA spells due to back pain.³² One can argue that if prediction of SA duration would be easy, a model would have been developed long ago. This model is a first step and we hope others will use other types of data to pursue development of such models.

The major strength of our work was its population-based design with a full coverage from nationwide registers,³⁹ as all SA spells >14 days that had been initialised in Sweden during the 2.5 years could be included in the study. Moreover, the use of high-quality register-based information^{26 28 40} (not self-reports) on the possible predictors, no dropouts, the complete and very large study group, meant that our data were not hampered by recall bias or selection bias and made it possible to validate our model for the whole population. However, it also limited our possibility to include some predictors that have been shown to be of importance for occurrence of SA but which are usually obtained through surveys, such as: self-rated health, sleep problems, body mass index, smoking or social support.^{17 18} Other studies might be able to investigate if such factors can increase the discriminative capacity of the model.

Another limitation of the study is that MiDAS includes only the main SA diagnosis of each SA spell, that is, we had no information on whether the physician assessed also other diagnoses to have contributed to the work incapacity. However, we were able to include information on multimorbidity based on dispensed prescribed medication and we included at least one purchase for a minimum of three different ATC groups; the number that had the highest predictive value. In the preliminary analyses, we also included information on multimorbidity from diagnosis-specific previous SA, DP, hospitalisation and specialised outpatient visits, however, information on dispensed prescribed medication contributed more to the model. Also, our results concern SA spells initiated in 2010–2012, and the predictive models need to be updated to examine possible needs of changing predictors when new data are available.

Furthermore, SA diagnosis was only registered at a 3-digit level for the diagnosis code (ICD-10) in MiDAS. As M54 is a large diagnosis group varying from unspecified pain conditions to inflammations and from radiculopathies to sciatica, information at a 4-digit level might have led to a possibility to make better predictions.

Our model was internally validated under the conditions applying to the Swedish healthcare and social insurance system, and therefore, can be applied in Sweden, and perhaps have applicability to other Nordic countries with similar systems. However, further research is needed to assess the performance in other social insurance settings as generalisability might be low in countries with different welfare systems and lower employment frequency than in the Nordic countries. The results of this work are currently implemented in pilot tests in primary healthcare settings to help general practitioners to identify which of their sickness absent patients have high risk for long-term SA.

CONCLUSIONS

We developed and validated a clinically implementable, pragmatic prediction model for the duration of SA spells >14 days due to back pain for the whole working-age population in Sweden. The model discriminates the long-term SA spells (>365 days) from shorter SA spells with good discriminatory accuracy.

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Contributors KA and EF contributed to the study conception and design. Material preparation, data collection and analysis were performed by KA, EF, KG, GA, MB and PF. The first draft of the manuscript was written by AR and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript.

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Competing interests None declared.

Patient consent for publication Not required.

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Data availability statement No data are available.

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REFERENCES

- 1 Bevan S, Quadrello T, McGee R, et al. *Fit for work? musculoskeletal disorders in the European workforce*. London: The Work Foundation, 2009.
- 2 High-Level Forum. *Sickness, disability and work: keeping on track in the economic downturn – background paper*. Stockholm, Sweden: Organisation for Economic Co-operation and Development, 2009.
- 3 OECD. *Sickness, disability and work: breaking the barriers*, 2010.
- 4 Försäkringskassan. *Social insurance in figures 2018*. Social Security Administration, Office of Policy, Office of Research, Evaluation, and Statistics, 2018.
- 5 Wittchen HU, Jacobi F, Rehm J, et al. The size and burden of mental disorders and other disorders of the brain in Europe 2010. *Eur Neuropsychopharmacol* 2011;21:655–79.
- 6 Lim SS, Vos T, Flaxman AD, et al. A comparative risk assessment of burden of disease and injury attributable to 67 risk factors and risk factor clusters in 21 regions, 1990–2010: a systematic analysis for the global burden of disease study 2010. *The Lancet* 2012;380:2224–60.
- 7 Edwards J, Hayden J, Asbridge M, et al. Prevalence of low back pain in emergency settings: a systematic review and meta-analysis. *BMC Musculoskelet Disord* 2017;18:143.
- 8 Meucci RD, Fassa AG, Faria NM. Prevalence of chronic low back pain: systematic review. *Rev Saúde Pública* 2015;49.
- 9 da C Menezes Costa L, Maher CG, Hancock MJ, et al. The prognosis of acute and persistent low-back pain: a meta-analysis. *CMAJ* 2012;184:E613–24.
- 10 O'Connell NE, Cook CE, Wand BM, et al. Clinical guidelines for low back pain: a critical review of consensus and inconsistencies across three major guidelines. *Best Pract Res Clin Rheumatol* 2016;30:968–80.
- 11 Wynne-Jones G, Cowen J, Jordan JL, et al. Absence from work and return to work in people with back pain: a systematic review and meta-analysis. *Occup Environ Med* 2014;71:448–56.
- 12 Hallegraeff JM, Krijnen WP, van der Schans CP, et al. Expectations about recovery from acute non-specific low back pain predict absence from usual work due to chronic low back pain: a systematic review. *J Physiother* 2012;58:165–72.
- 13 Lötters F, Burdorf A. Prognostic factors for duration of sickness absence due to musculoskeletal disorders. *Clin J Pain* 2006;22:212–21.
- 14 Airaksinen J, Jokela M, Virtanen M, et al. Development and validation of a risk prediction model for work disability: multicohort study. *Sci Rep* 2017;7:13578.
- 15 Wingbermühle RW, van Trijffel E, Nelissen PM, et al. Few promising multivariable prognostic models exist for recovery of people with non-specific neck pain in musculoskeletal primary care: a systematic review. *J Physiother* 2018;64:16–23.
- 16 Rose S. Machine learning for prediction in electronic health data. *JAMA Netw Open* 2018;1:e181404.

- 17 Notenbomer A, van Rhenen W, Groothoff JW, *et al.* Predicting long-term sickness absence among employees with frequent sickness absence. *Int Arch Occup Environ Health* 2019;92:501–11.
- 18 Airaksinen J, Jokela M, Virtanen M, *et al.* Prediction of long-term absence due to sickness in employees: development and validation of a multifactorial risk score in two cohort studies. *Scand J Work Environ Health* 2018;44:274–82.
- 19 Roelen C, Thorsen S, Heymans M, *et al.* Development and validation of a prediction model for long-term sickness absence based on occupational health survey variables. *Disabil Rehabil* 2018;40:168–75.
- 20 Pind R. Testing a new 10-Item scale (Pind's LBP test) for prediction of sick leave lasting more than three days or more than two weeks after a general practitioner visit for acute low back pain. *Spine* 2014;39:E581–6.
- 21 Bosman LC, Dijkstra L, Oling CI, *et al.* Prediction models to identify workers at risk of sick leave due to low-back pain in the Dutch construction industry. *Scand J Work Environ Health* 2018;44:156–62.
- 22 Gémes K, Frumento P, Almondo G, *et al.* A prediction model for duration of sickness absence due to stress-related disorders. *J Affect Disord* 2019;250:9–15.
- 23 Turner JA, Shortreed SM, Saunders KW, *et al.* Optimizing prediction of back pain outcomes. *Pain* 2013;154:1391–401.
- 24 Bardin LD, King P, Maher CG. Diagnostic triage for low back pain: a practical approach for primary care. *Med J Aust* 2017;206:268–73.
- 25 World Health Organization. *ICD-10, the ICD-10 classification of mental and behavioural disorders: diagnostic criteria for research version: 2016*. Geneva: World Health Organization, 1993.
- 26 Ludvigsson JF, Otterblad-Olausson P, Pettersson BU, *et al.* The Swedish personal identity number: possibilities and pitfalls in healthcare and medical research. *Eur J Epidemiol* 2009;24:659–67.
- 27 Statistics Sweden. *Integrated database for labour market research*. Statistics Sweden, 2009.
- 28 Ludvigsson JF, Andersson E, Ekblom A, *et al.* External review and validation of the Swedish national inpatient register. *BMC Public Health* 2011;11:450.
- 29 Wettermark B, Hammar N, Fored CM, *et al.* The new Swedish Prescribed Drug Register--opportunities for pharmacoepidemiological research and experience from the first six months. *Pharmacoepidemiol Drug Saf* 2007;16:726–35.
- 30 Brooke HL, Talbäck M, Hörnblad J, *et al.* The Swedish cause of death register. *Eur J Epidemiol* 2017;32:765–73.
- 31 Allebeck P, Mastekaasa A. Chapter 5. Risk factors for sick leave - general studies. *Scand J Public Health* 2004;32:49–108.
- 32 Hansson T, Jensen I. Chapter 6. sickness absence due to back and neck disorders. *Scand J Public Health* 2004;32:109–51.
- 33 Uno H, Cai T, Pencina MJ, *et al.* On the C-statistics for evaluating overall adequacy of risk prediction procedures with censored survival data. *Stat Med* 2011;30:1105–17.
- 34 Harell F. *Regression modeling strategies: with applications to linear models, logistic regression, and survival analysis*. New York: Springer, 2001.
- 35 Robin X, Turck N, Hainard A, *et al.* pROC: an open-source package for R and S+ to analyze and compare ROC curves. *BMC Bioinformatics* 2011;12:77.
- 36 RCoreTeam. *R: a language and environment for statistical computing*. Vienna, Austria: R Foundation for Statistical Computing, 2017.
- 37 Frumento P. Pch: Piecewise constant hazards models for censored and truncated data: R package version 1.2, 2016. Available: <http://CRAN.Rproject.org/package=pch>
- 38 Pedersen J, Gerds TA, Bjoerner JB, *et al.* Prediction of future labour market outcome in a cohort of long-term sick- listed Danes. *BMC Public Health* 2014;14:494.
- 39 Försäkringskassan. *MIDAS Sjukpenning och rehabiliteringspenning (The MIDAS register. Sickness absence benefits)(In Swedish)*, 2011.
- 40 Ludvigsson JF, Almqvist C, Bonamy A-KE, *et al.* Registers of the Swedish total population and their use in medical research. *Eur J Epidemiol* 2016;31:125–36.