

# Seasonality of antidepressant prescriptions and sick leaves

Dietmar Winkler<sup>1</sup>, Berthold Reichardt<sup>2</sup>, Georg S. Kranz<sup>1,3,4</sup>, Lucie Bartova<sup>1</sup>, Siegfried Kasper<sup>1</sup>, Edda Pjrek<sup>1</sup>

<sup>1</sup> Department of Psychiatry and Psychotherapy, Medical University of Vienna, Austria

<sup>2</sup> Sickness Fund Burgenland (BGKK), Eisenstadt, Austria

<sup>3</sup> Department of Rehabilitation Sciences, The Hong Kong Polytechnic University, Hung Hom, Hong Kong, China

<sup>4</sup> Laboratory of Neuropsychology, The University of Hong Kong, Pokfulam, Hong Kong, China

Send all correspondence to:

Assoc. Prof. Priv.-Doz. Dr. Dietmar Winkler, M.D.  
Department of Psychiatry and Psychotherapy  
Medical University of Vienna  
Währinger Gürtel 18-20  
A-1090 Vienna

Phone: +43.1.40400.35470

Fax: +43.1.40400.30990

Email: dietmar.winkler@meduniwien.ac.at

Word count:	2188	Main section
	249	Abstract
33 references, 2 table, 6 figures		

*Submitted for publication in*

**Journal of Clinical Psychiatry**

July 2018

## **ABSTRACT**

**Objective:** The aim of the present study was to estimate the number of patients with a seasonal prescription pattern of antidepressants, which might be taken as a surrogate marker for medicated patients with seasonal affective disorder (SAD). Furthermore, we examined the time course of sick leaves for patients with seasonal and non-seasonal prescriptions of antidepressants.

**Methods:** A retrospective analysis of prescription data of all patients insured by the Sickness Fund Burgenland (BGKK) between 2005 and 2016 was performed. Patients with treatment initiation of an antidepressant in the last and first quarter of the year for at least two consecutive years were selected (SAD-med). Patients with continuation treatment in the third quarter and patients with initiation of antidepressant medication in the second and third quarter of the year were excluded.

**Results:** The mean yearly prescription rate for antidepressants was 9.6% in the insured population. 3.0% of patients treated with antidepressants and 0.9% of insured cases satisfied the definition of SAD-med. The mean number of yearly sick leave days was similar for SAD-med patients and those with non-seasonal prescriptions. Time series analysis showed that sick leaves in SAD-med were influenced by seasonal fluctuations for several years after the first antidepressant prescription.

**Conclusions:** Our study sheds light on antidepressant prescription and sick leave patterns in the general population. Compared to the prevalence of SAD, the estimated rate of SAD-med is substantial. Sick leaves appear to be closely linked to antidepressant prescriptions, and show a characteristic annual rhythm before and after the initial prescription.

## **KEYWORDS**

Seasonal affective disorder, fall-winter depression, sick leaves, antidepressant prescription

## **1. INTRODUCTION**

Seasonal affective disorder (SAD) is a subtype of mood disorder characterized by recurrent depressive episodes during fall and winter with subsequent remission or hypomanic/manic episodes during the next spring/summer period<sup>1</sup>. The prevalence of SAD according to DSM-IV/5 criteria<sup>2, 3</sup> in the general population has been estimated between 2.4% and 3.1% in temperate climates<sup>4-7</sup>. Light therapy is the first choice of treatment for SAD<sup>8</sup>. Antidepressant treatment is an alternative option for SAD and is often necessitated in patients with insufficient response to light therapy or logistical problems in adhering to this treatment<sup>9</sup>. Previous studies investigated the clinical usage of light therapy in psychiatric hospitals<sup>10</sup> and in office-based doctors<sup>11</sup>, but little is known about the actual use of psychopharmacologic medication in SAD in clinical practice.

The clinical diagnosis of SAD and most epidemiological and clinical studies on SAD rely on a history of seasonal mood episodes and are therefore subject to recall bias except in studies with a longitudinal design<sup>12-15</sup>. The retrospective approach of the diagnosis of SAD was criticized before<sup>16-18</sup>. Therefore, corroboration of the validity of SAD as a diagnostic entity by an alternative approach is an important line of research. Furthermore, there is a paucity of studies on the work-related consequences of SAD in comparison to non-seasonal depression. Sick leaves of SAD patients were studied before in clinical<sup>19</sup> and population-based samples<sup>4, 7</sup>, but these studies also relied on self-reports.

The aim of this study was to estimate the rate of seasonal prescriptions of antidepressants as surrogate marker for treated winter-type SAD in a large sample of the general population. Furthermore, we wanted to describe the pattern of sick leaves in patients with seasonal versus non-seasonal antidepressant prescriptions. We hypothesized a priori that sick leaves of patients with seasonal prescriptions would be accompanied with identifiable seasonality.

## **2. METHOD**

This study was approved by the Ethics Committee of the Medical University of Vienna (EC No. 1018/2017). A retrospective analysis of prescription data of all patients insured by the Sickness Fund Burgenland (BGKK) between 2005 and 2016 was performed. During this time frame a mean number of 195,135 persons (68.4% of the inhabitants of the Burgenland, one of the nine federal states of Austria) were insured by the BGKK. Patients with treatment

initiation of an antidepressant between the beginning (1st October) and the end (31st March) of the fall/winter season for at least two consecutive years were selected. Patients with continuation treatment during 1st July and 30th September and patients with initiation of antidepressant medication between 1st April and 30th September were excluded. This definition of seasonal prescriptions was termed SAD-med.

All dates of sick leaves during 2004 and 2016 were derived from the BGKK database for SAD-med patients and patients with non-seasonal prescriptions of antidepressants (non-SAD). A total of 167.121 sick leave periods were transformed using a special data converter programmed with C# and Microsoft Visual Studio 2017<sup>20</sup> resulting in tabular daily sick leave data relative to the first antidepressant prescription for each single patient.

Statistical analyses were performed using the R Project for Statistical Computing (version 3.4.3)<sup>21</sup> together with the packages gmodels<sup>22</sup>, forecast<sup>23</sup>, lmtest<sup>24</sup>, and ggplot2<sup>25</sup>.

After visual inspection, sick leave data of SAD-med and non-SAD patients from day -120 to day +2190 relative to the first antidepressant prescription were further aggregated to form time series by months. Friedman test was performed to test the null hypothesis of equal distribution of detrended data (after calculating first differences) with months as groups.

An autoregressive integrated moving average (ARIMA) model was fit to the time series both using the auto.arima method of R's forecast package (selection criteria being the Akaike information criterion [AIC], the Bayesian information criterion [BIC] and parsimony) and a stepwise manual selection process based on the autocorrelation (Acf) and partial autocorrelation (Pacf) functions of the time series and the residuals of the models as well as considering root mean square error (RMSE) and mean absolute scaled error (MASE).

The frequency of sick leaves was compared at four a priori defined 30 day time periods: half a year before the first antidepressant prescription (days -198 to -168), around the day of prescription (days -15 to 15), half a year (days 168 to 198) and one year (days 350 to 380) after the first prescription. For sick leave calculations only data of patients between 18 to 65 years were used and influences of shorter observation periods were accounted for. The level of statistical significance (two-tailed) was set to  $p \leq 0.05$ . The Bonferroni-Holm correction was applied to the p-values to correct for multiple testing where appropriate.

### 3. RESULTS

During the 12 years of analysis, a total of 58,138 patients (29.8%) received at least one prescription of an antidepressant. The mean yearly prescription rate of antidepressants was 9.6% of all insured persons (mean 10-year prescription rate: 21.8%). We found 1750 patients (3.0% of treated patients and 0.9% of all insured cases) satisfying the definition of SAD-med (mean 10-year prevalence 2.5% and 0.7%, respectively). Table 1 displays demographic variables and group differences. The rates by year for patients with at least one antidepressant prescription and SAD-med patients are presented in Figure 1. 79.0% of patients in the SAD-med group had seasonal prescriptions for 2 years, 15.7% for 3 years, 3.2% for 4 years, 0.9% for 5 years, and 1.2% for 6 to 12 years (Figure 2).

The mean number of sick leave days per year was  $17.5 \pm 30.1$  in all patients with at least one antidepressant prescription. A comparison of sick leave days in SAD-med and non-SAD (also subdivided by gender) is presented in table 2. Percentages of patients being on sick leave in the SAD-med and non-SAD group for each day 1 year before to 5 years after the initial antidepressant prescription are presented in Figure 4. Comparisons for sick leaves showed a significant difference between the time period -0.5 years before first prescription and the following predefined epochs (0, +0.5 and +1 year relative to prescription) in SAD-med ( $\chi^2 = 136.94$ ,  $df = 3$ ,  $p < 0.0001$ ) and non-SAD ( $\chi^2 = 4234.5$ ,  $df = 3$ ,  $p < 0.0001$ ). Post-hoc tests revealed that compared to -0.5 years the SAD-med group had higher sick leaves at 0 years ( $Z = -10.013$ ,  $p < 0.0001$ ) and at +1 year ( $Z = -3.023$ ,  $p = 0.015$ ), but not at +0.5 years ( $Z = -1.987$ ,  $p = 0.188$ ), while the non-SAD group exhibited higher sick leaves only at 0 years ( $Z = -8.211$ ,  $p < 0.0001$ ) and 0.5 years ( $Z = -11.515$ ,  $p < 0.0001$ ) compared to -0.5 years. SAD-med patients had significantly higher sick leave levels than non-SAD patients 1 year after the initial prescription ( $Z = -5.522$ ,  $p < 0.0001$ ), but group differences were not significant at -0.5, 0 and 0.5 years.

Friedman test for sick leave data in SAD-med patients using month as a factor was statistically significant ( $\chi^2 = 24.169$ ,  $df = 11$ ,  $p = 0.012$ ), but not in non-SAD patients ( $\chi^2 = 11.985$ ,  $df = 11$ ,  $p = 0.365$ ). An ARIMA (0, 1, 0)  $\times$  (1, 0, 0)<sub>12</sub> model was selected as the best model to describe monthly time series in SAD-med (RMSE = 0.132, MASE = 0.323). The seasonal AR term (estimate  $\pm$  standard error:  $0.550 \pm 0.126$ ) was statistically significant ( $p < 0.0001$ ). A Box-Ljung test was applied to the residuals of the ARIMA model and found to be not significant ( $\chi^2 = 24.926$ ,  $df = 24$ ,  $p = 0.410$ ). The best ARIMA model for non-SAD data

was (1, 1, 0) without any seasonal terms (Box-Ljung test:  $\chi^2 = 6.039$ ,  $df = 24$ ,  $p = 0.999$ ). Seasonal decomposition<sup>26</sup> of monthly time series data of sick leaves of SAD-med patients into seasonal fluctuations, trend and remainder is displayed in Figure 3.

#### 4. DISCUSSION

To our best knowledge this is the first study to provide comprehensive information on seasonal prescription patterns of antidepressants and associated sick leaves.

The prescription rate of antidepressants over the study period of 12 years was nearly 30% of the total population. On the other hand the adherence to antidepressants seems to be quite low: 15397 of 58138 patients (26.5%) only had a single prescription of an antidepressant with no re-prescriptions (Figure 6), which is in line with previously published data<sup>27</sup>. These early drop-outs are problematic insofar as treatment guidelines for major depression and anxiety disorders recommend treatment periods of 6 months or longer<sup>28, 29</sup>. Only 45.6% of our patients had 6 or more prescriptions, allowing us to infer that the majority of patients might not be treated according to the guidelines. Different reasons on part of the health system, the medical doctors, and the patients might be implicated in this phenomenon.

The 10-year prevalence of SAD according to the DSM-5<sup>3</sup> in the general population of Austria has been reported to be as high as 2.5%<sup>7</sup>. Our findings showed seasonal prescriptions as surrogate marker for treated SAD (SAD-med) in 0.74%. If SAD-med were a perfect match with the DSM-5 definition of SAD, this would indicate that nearly 30% of SAD patients would receive antidepressant treatment at some time point. However, the number of years with seasonal treatment (i. e. the stability of the definition) was low in SAD-med patients, which might be due to the generally low adherence to medication or to antidepressants being only second line for SAD patients beside other treatments (e. g. light therapy).

Between 2004 and 2016 the mean number of sick leaves in the Austrian working population was 12.7 days per year<sup>30</sup>. Within the observation time frame of -1 year before prescription to 5 years afterwards patients with antidepressant prescriptions in our sample had 37.8% more sick leave days. This figure strongly depends on the reference period and restriction of the time frame of observation around the prescription date results in far higher numbers.

Subgroup analysis of sick leave levels showed that males had higher sick leave days per year than females in our sample as is also observed in the general population of Austria<sup>30</sup>. A statistically significant difference between SAD-med and non-SAD was found, but the values of the two groups are numerically quite similar. This leads us to conclude that in terms of sick leaves SAD-med patients are no less sick than patients of the non-SAD group. Data from the literature already suggested higher sick leave levels of SAD patients than in subjects without the syndrome<sup>4, 7</sup>. Our analysis provides strong evidence for seasonality in sick leaves in SAD-med, while this feature is lacking in the non-SAD group. This is particularly important from a socioeconomic point of view, because the predictable and recurrent nature of SAD makes it especially well-suited for preventative treatment<sup>31, 32</sup>.

We could assume that the use of antidepressants leads to a decrease of sick leaves, but such a simple relationship cannot be derived from our data. The sick leave curve over time (figure 4) has a particular shape with a sharp increase during the last weeks before initial prescription and a more gradual decline afterwards. Nevertheless, return to baseline levels is slow and takes at least one year in non-SAD patients (in the SAD-med group sick leave levels peak again after one year). We can hypothesize that it is the increasing dynamics of the illness that leads to patients being signed off sick by their medical doctors shortly before or after the prescription of an antidepressant. A similar observation of a close link between sick leaves and antidepressant prescriptions has already been made by Gasse et al.<sup>33</sup> in a Danish cohort study. However, in contrast to their study our present analysis had an even larger sample and we were also able to include short-term sick leaves. The peak of sick leaves at initial prescription is produced by synchronization of the prescription date as day zero. The gradual decline over time after the initial prescription in the non-SAD group is not only due to clinical improvement of the patients but also due to patients having recurrent episodes at different time points and coming out of sync, whereas SAD-med patients are by definition synchronized with the seasons within the first two years.

Our approach might be limited by including patients with seasonal prescriptions for indications other than SAD (e. g. seasonal psychosocial stress). Moreover, SAD patients who only received treatment with an antidepressant once, would be found in the non-SAD group in this study. Finally, SAD patients with continuation treatment of antidepressants during the summer period were not detectable by our definition of SAD-med.

In this large study population, we were able to show that antidepressant prescriptions are regularly preceded and followed by increases of sick leaves, which lead to overall higher sick leave levels compared to the general population. In patients with seasonal recurrence of antidepressant treatment (3% of patients with an antidepressant prescription) also the sick leaves show a recurrence during the next years. Patients with seasonally recurring prescriptions have similar sick leave levels than patients with no seasonality. What we cannot learn from our study is what happens to patients with seasonal prescriptions, when they stop to take antidepressants. How many remit or develop another disorder? How many have non-pharmacological treatments? It will be for future studies to longitudinally investigate the treatment paths of those patients.



## **STATEMENT OF INTEREST**

Dr. Kasper received grants/research support, consulting fees and/or honoraria within the last three years from Angelini, AOP Orphan Pharmaceuticals AG, AstraZeneca, Eli Lilly, Janssen, KRKA-Pharma, Lundbeck, Neuraxpharm, Pfizer, Pierre Fabre, Schwabe and Servier. Dr. Winkler has received lecture fees from Angelini, Lundbeck, and Pfizer. Dr. Kranz received travel grants from Roche, Pfizer and AOP Orphan Pharmaceuticals AG. The other authors report no financial or other relationship possibly relevant to the subject of this article.

## REFERENCES

1. Rosenthal NE, Sack DA, Gillin JC, et al. Seasonal affective disorder. A description of the syndrome and preliminary findings with light therapy. *Archives of General Psychiatry* 1984 Jan;41(1):72-80.
2. American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorder*, 4th edition, text revision (DSM-IV-TR). Washington, DC: American Psychiatric Press; 2000.
3. American Psychiatric Association. *Diagnostic and statistical manual of mental disorders*. 5th edition. Washington, DC: American Psychiatric Publishing; 2013.
4. Mersch PP, Middendorp HM, Bouhuys AL, Beersma DG, van den Hoofdakker RH. The prevalence of seasonal affective disorder in The Netherlands: a prospective and retrospective study of seasonal mood variation in the general population. *Biological Psychiatry* 1999 Apr 15;45(8):1013-1022.
5. Levitt AJ, Boyle MH. The impact of latitude on the prevalence of seasonal depression. *Canadian Journal of Psychiatry* 2002 May;47(4):361-367.
6. Michalak EE, Wilkinson C, Dowrick C, Wilkinson G. Seasonal affective disorder: prevalence, detection and current treatment in North Wales. *British Journal of Psychiatry* 2001 Jul;179:31-34.
7. Pjrek E, Baldinger-Melich P, Spies M, Papageorgiou K, Kasper S, Winkler D. Epidemiology and socioeconomic impact of seasonal affective disorder in Austria. *European Psychiatry* 2016 Feb;32:28-33.
8. Pail G, Huf W, Pjrek E, et al. Bright-light therapy in the treatment of mood disorders. *Neuropsychobiology* 2011;64(3):152-162.
9. Pjrek E, Winkler D, Stastny J, Konstantinidis A, Heiden A, Kasper S. Bright light therapy in seasonal affective disorder--does it suffice? *European Neuropsychopharmacology* 2004 Aug;14(4):347-351.

10. Fischer R, Kasper S, Pjrek E, Winkler D. On the application of light therapy in German-speaking countries. *European Archives of Psychiatry and Clinical Neuroscience* 2012 Sep;262(6):501-505.
11. Winkler-Pjrek E, Spies M, Baldinger-Melich P, Perkmann L, Kasper S, Winkler D. Use of Light Therapy by Office-Based Physicians. *Neuropsychobiology* 2016;74(4):182-187.
12. Clery-Melin ML, Gorwood P, Friedman S, Even C. Stability of the diagnosis of seasonal affective disorder in a long-term prospective study. *Journal of Affective Disorders* 2018 Feb;227:353-357.
13. Harmatz MG, Well AD, Overtree CE, Kawamura KY, Rosal M, Ockene IS. Seasonal variation of depression and other moods: a longitudinal approach. *Journal of Biological Rhythms* 2000 Aug;15(4):344-350.
14. Sakamoto K, Nakadaira S, Kamo K, Kamo T, Takahashi K. A longitudinal follow-up study of seasonal affective disorder. *American Journal of Psychiatry* 1995 Jun;152(6):862-868.
15. Wicki W, Angst J, Merikangas KR. The Zurich Study. XIV. Epidemiology of seasonal depression. *European Archives of Psychiatry and Clinical Neuroscience* 1992;241(5):301-306.
16. Traffanstedt MK, Mehta S, LoBello SG. Major depression with seasonal variation: Is it a valid construct? *Clinical Psychological Science* 2016;4(5):825-834.
17. Winkler D, Pjrek E, Spies M, et al. Has the existence of seasonal affective disorder been disproven? *Journal of Affective Disorders* 2017 Jan 15;208:54-55.
18. Hansen V, Skre I, Lund E. What is this thing called "SAD"? A critique of the concept of Seasonal Affective Disorder. *Epidemiologia e psichiatria sociale* 2008 Apr-Jun;17(2):120-127.
19. Pjrek E, Willeit M, Praschak-Rieder N, et al. Treatment of seasonal affective disorder with duloxetine: an open-label study. *Pharmacopsychiatry* 2008 May;41(3):100-105.

20. Microsoft Corporation. Microsoft Visual Studio 2017. Redmond, WA, USA; 2017.
21. R Core Team. R: A language and environment for statistical computing. R Foundation for Statistical Computing. <https://www.R-project.org/>. Vienna, Austria; 2017.
22. Warnes GR, Bolker B, Lumley T, Johnson RC. gmodels: Various R Programming Tools for Model Fitting. R package version 2.16.2. <https://CRAN.R-project.org/package=gmodels>; 2015.
23. Hyndman R. forecast: Forecasting functions for time series and linear models. R package version 8.2. <http://pkg.robjhyndman.com/forecast>; 2017.
24. Zeileis A, Hothorn T. Diagnostic Checking in Regression Relationships. R News 2(3), 7-10. <https://CRAN.R-project.org/doc/Rnews/>; 2002.
25. Wickham H. ggplot2: Elegant Graphics for Data Analysis. <https://ggplot2.tidyverse.org/>. New York: Springer; 2009.
26. Cleveland RB, W. S. Cleveland WS, McRae JE, Terpenning I. STL: A Seasonal-Trend Decomposition Procedure Based on Loess. Journal of Official Statistics 1990;6:3-73.
27. Hinteregger M, Füzi J, Reichardt B. Tracking antidepressant therapy patterns of an Austrian cohort. WONCA Europe Conference. Vienna; 2012.
28. Bandelow B, Sher L, Bunevicius R, et al. Guidelines for the pharmacological treatment of anxiety disorders, obsessive-compulsive disorder and posttraumatic stress disorder in primary care. International Journal of Psychiatry in Clinical Practice 2012 Jun;16(2):77-84.
29. Bauer M, Severus E, Kohler S, et al. World Federation of Societies of Biological Psychiatry (WFSBP) guidelines for biological treatment of unipolar depressive disorders. part 2: maintenance treatment of major depressive disorder-update 2015. World Journal of Biological Psychiatry 2015 Feb;16(2):76-95.
30. Statistik Austria. Krankenstandsfälle und -tage nach Geschlecht seit 1965. <https://www.statistik.at>; 2016.

31. Nussbaumer-Streit B, Pjrek E, Kien C, et al. Implementing prevention of seasonal affective disorder from patients' and physicians' perspective – a qualitative study. *BMC Psychiatry* 2018; submitted for publication.
32. Nussbaumer-Streit B, Winkler D, Spies M, Kasper S, Pjrek E. Prevention of seasonal affective disorder in daily clinical practice: results of a survey in German-speaking countries. *BMC Psychiatry* 2017 Jul 11;17(1):247.
33. Gasse C, Petersen L, Chollet J, Saragoussi D. Pattern and predictors of sick leave among users of antidepressants: a Danish retrospective register-based cohort study. *Journal of Affective Disorders* 2013 Dec;151(3):959-966.

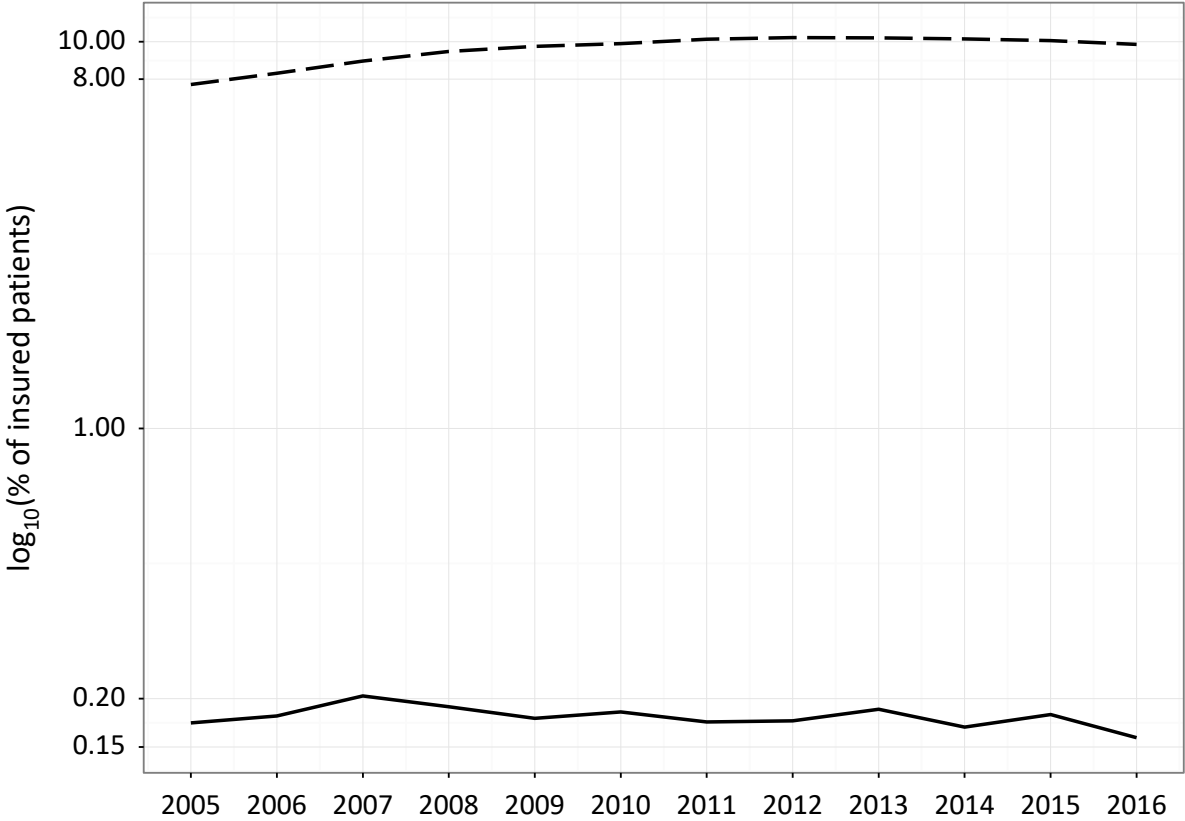
**TABLE 1.** Demographic variables of patients in the sample.

	<b>SAD-med</b>	<b>non-SAD</b>	
<b>N</b>	1750 (3.0%)	56388 (97.0%)	
<b>Sex</b>			
Females	1116 (63.8%)	37487 (66.5%)	$\chi^2 = 5.583, df = 1, p = 0.018$
Males	634 (36.2%)	18901 (33.5%)	
<b>Age</b> (years, $\mu \pm SD$ )	53.5 $\pm$ 17.8	55.4 $\pm$ 19.0	$t = -4.339, df = 1874.3, p < 0.0001$

**TABLE 2.** Number of sick leave days per year of SAD-med and non-SAD patients (subgroup of those between 18-65 years old) starting 1 year before initial antidepressant prescription to 5 years afterwards.

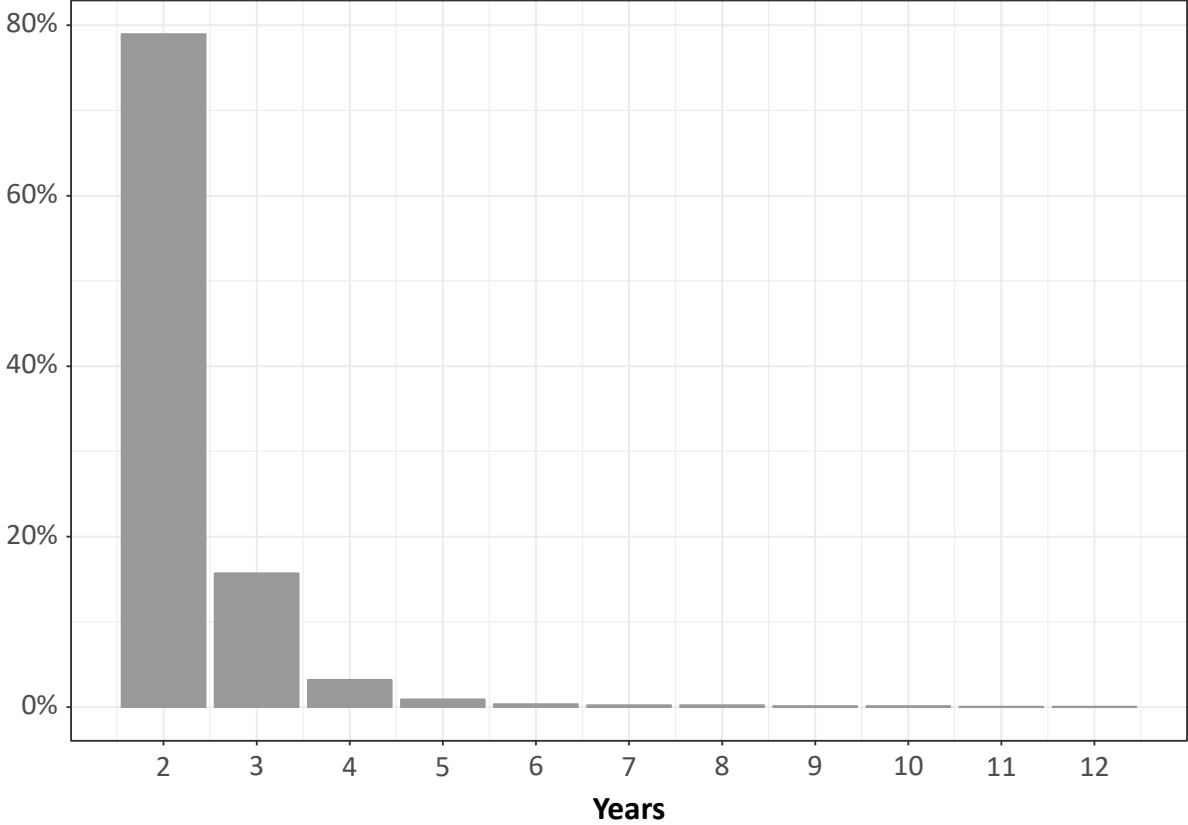
<b>Sick leave days / year (<math>\mu \pm SD</math>)</b>			
	<b>SAD-med</b>	<b>non-SAD</b>	
Total group	17.67 $\pm$ 28.09	17.45 $\pm$ 30.17	Z = -3.070, p < 0.002
Females	15.95 $\pm$ 27.31	16.05 $\pm$ 28.51	Z = -1.729, p = 0.084
Males	20.51 $\pm$ 29.14	20.07 $\pm$ 32.89	Z = -2.529, p = 0.011

**FIGURE 1.** Percentage of SAD-med patients (N=1750; solid line; see method section for definition) and percentage of patients with at least one antidepressant prescription (N=58138; dashed line) by year in all persons insured by the Sickness Fund Burgenland (BGKK) between 2005 and 2016.

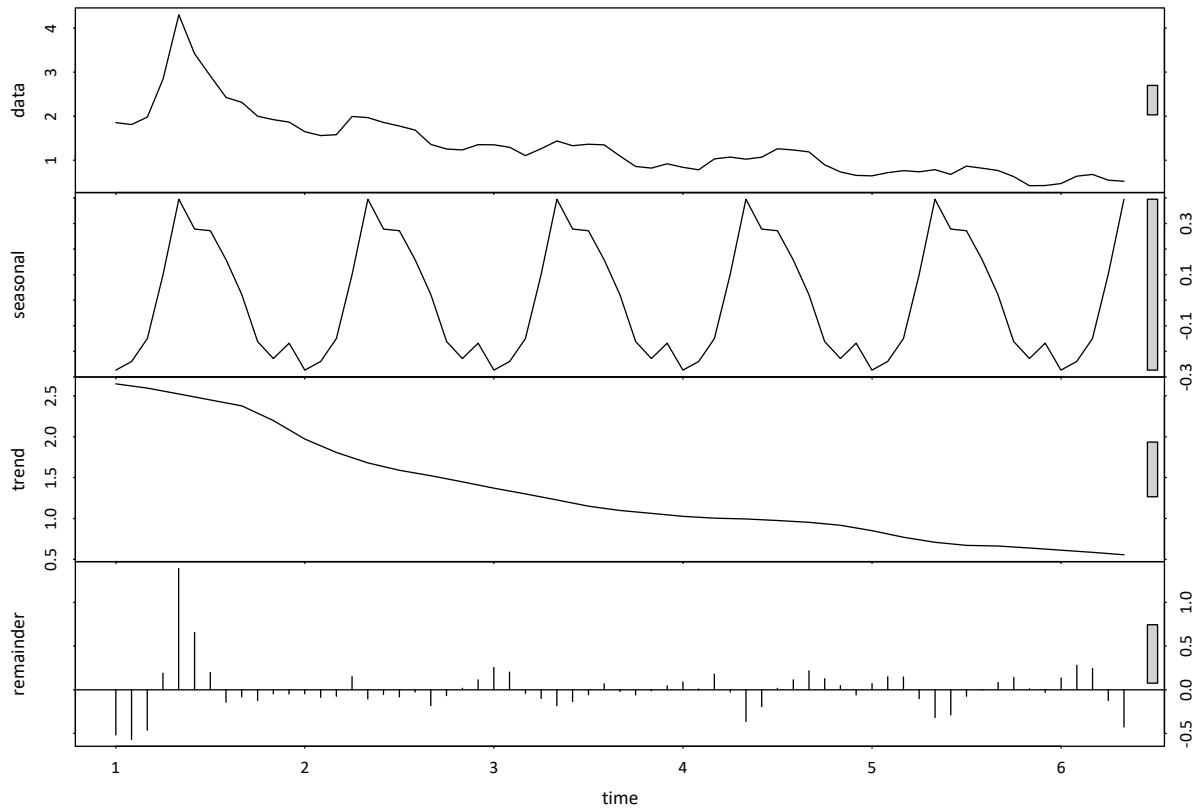




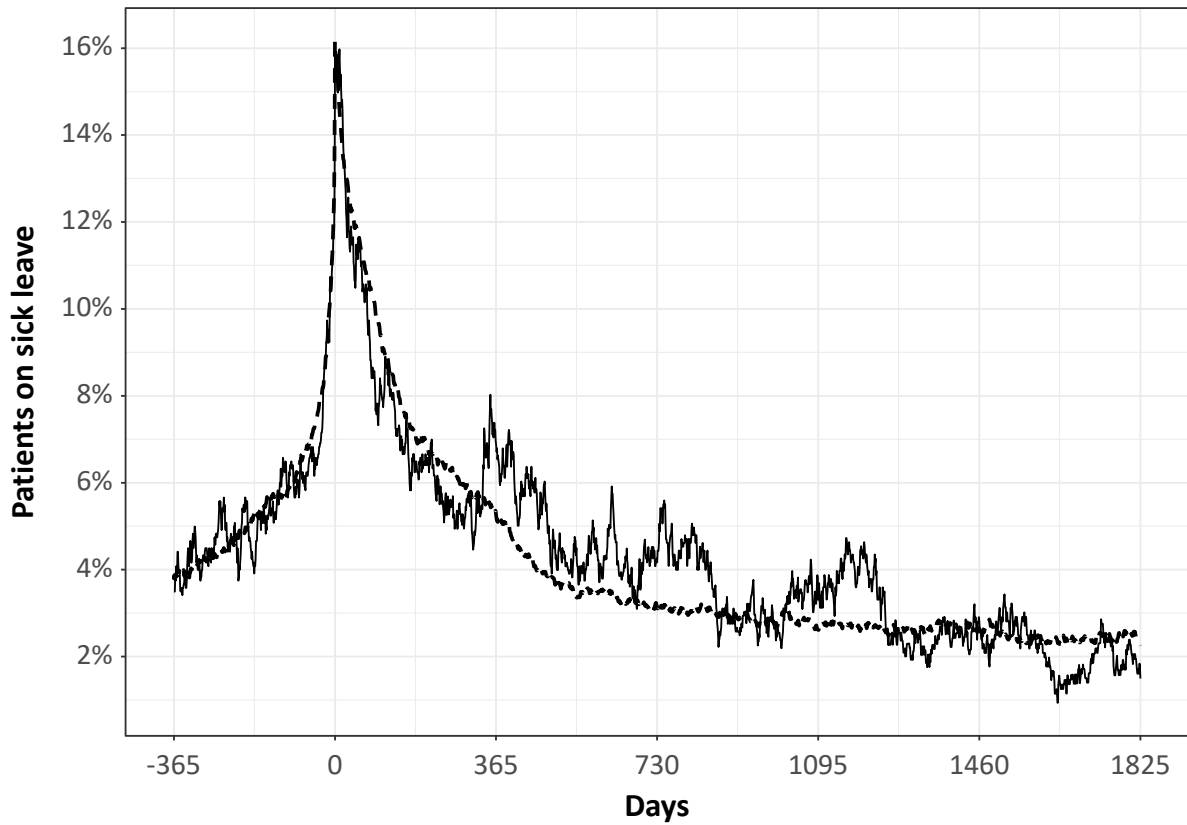
**FIGURE 2.** Percentage of SAD-med patients fulfilling the definition of SAD-med by number of years.



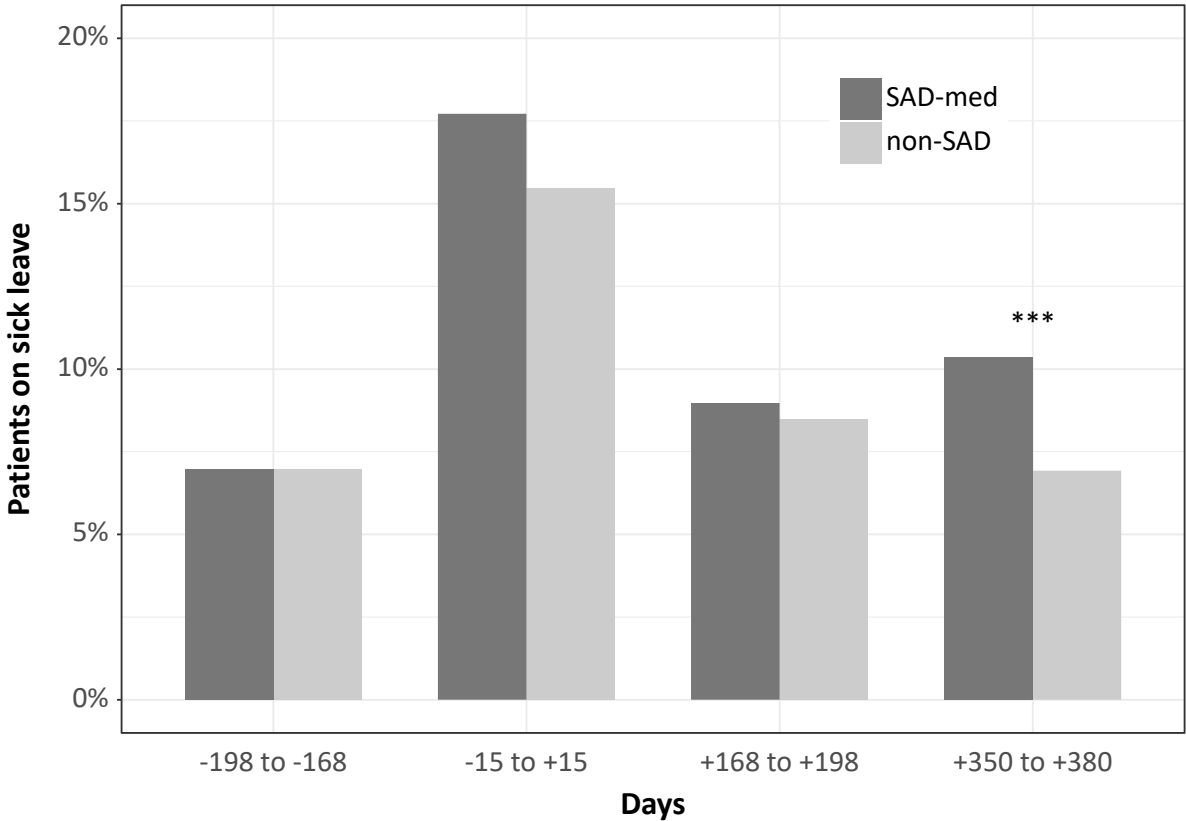
**FIGURE 3.** Seasonal decomposition of monthly time series data by loess<sup>26</sup> for sick leaves of SAD-med patients. Shown is the original time series in the top row, followed by the extracted seasonal pattern, the trend of the time series and the remainder in the following rows. Time series starts 3 months before the initial antidepressant prescription.



**FIGURE 4.** Percentage of patients on sick leave starting from 1 year (day -365) prior to the first prescription of an antidepressant (day 0) to 5 years (day 1825) afterwards. SAD-med patients are presented with a solid line and non-SAD patients with a dashed line. A distinctive seasonal pattern can readily be identified in SAD-med patients.



**FIGURE 5.** Percentage of SAD-med and non-SAD patients on sick leave during 4 à priori defined 30 day time periods: 1. half a year before antidepressant prescription (day -198 to -168), 2. around the first antidepressant prescription (day -15 to +15), 3. half a year later (day +168 to +198), and 4. one year later (day +350 to +380). \*\*\*  $p < 0.0001$  between SAD-med and non-SAD group.



**FIGURE 6.** Patients insured by the Sickness Fund Burgenland (BGKK) between 2005 and 2016 and with at least one antidepressant prescription (N=58138). Given is the percentage of patients with a certain number of prescriptions (1 to 19 or >20).

