

BMJ Open Effect of reimbursement restriction policy on the use of benzodiazepines in the Netherlands: an interrupted time series analysis

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ABSTRACT

Objectives Use of benzodiazepines has health risks. Reimbursement was restricted in the Netherlands from January 2009 onwards with the goal to reduce chronic use and healthcare expenditures. The aim of this study is to assess the initial and long-term effects of this policy on benzodiazepine use.

Design Interrupted time series analysis, segmented regression models, Kaplan-Meier survival analysis and Cox proportional hazards analysis.

Setting A 10% random sample of benzodiazepine dispensings by outpatient pharmacies between January 2002 and August 2015 were obtained from the PHARMO database. This database covered a catchment area representing about 3.6 million residents in 2015.

Participants 2500800 benzodiazepine prescriptions from 128603 patients were included.

Intervention Reimbursement restriction policy from January 2009 onwards.

Outcome measures Changes in: the volume of dispensed prescriptions and doses, the incidence, prevalence of incidental, regular and chronic use and discontinuation rates of benzodiazepines.

Results The volume of dispensed prescriptions and doses decreased by 12.5% (95% CI 9.0% to 15.9%) and 15.1% (95% CI 11.4% to 17.3%) respectively in January 2009 compared with December 2008. A clear initial effect on the overall incidence (−14.7%; 95% CI −19.8% to 9.6%) and the prevalence of incidental (−17.8%; 95% CI −23.9% to 11.7%), regular (−20.0%; 95% CI −26.1% to 13.9%) and chronic (−16.0%; 95% CI −23.1% to 8.9%) use was observed. A statistically significant reduction in the monthly trend per 1000 medication users was observed for the overall incidence (−0.017; 95% CI −0.031 to 0.003) and the prevalence of incidental (−3.624; 95% CI −4.996 to 2.252) but not for regular (−0.304; 95% CI −1.204 to 0.596) and chronic (0.136; 95% CI −0.858 to 1.130) use. Patients who started treatment before policy had a slightly higher probability of discontinuation (HR=1.013; 95% CI 1.004 to 1.022).

Conclusions The reimbursement policy had a significant initial effect on the volume, incidence and prevalence of benzodiazepine use. In addition, there is a statistically significant reduction in the monthly trend of overall incidence and of the prevalence of incidental use. No statistically significant reduction in the monthly trend

Strengths and limitations of this study

- ▶ The strengths of this study are the large sample size, long follow-up, large number of time intervals and presence of data of all outpatient prescriptions, including those initiated in secondary care. This makes it possible to focus on short-term and long-term effects.
- ▶ Another strength is use of an interrupted time series design to evaluate the effect of this policy change. This analysis is the golden standard for evaluation of the effects of policy change.
- ▶ This study has several limitations. First, it is difficult to attribute the observed changes in benzodiazepine use to a single intervention, the reimbursement restriction policy, because many confounding factors change in time. However, to our knowledge there were no other health system changes that could be a rival explanation for the observed changes.
- ▶ Second, due to PHARMO regulations, only a 10% random sample of PHARMO database was obtained and analysed. Nevertheless, the statistical significance of our analysis shows that the size of the data set is sufficient.

of chronic use, the main purpose of the reimbursement restriction, could be demonstrated.

INTRODUCTION

Benzodiazepines (BZD) are widely used in the treatment of panic disorders, anxiety, insomnia and epilepsy. In addition to beneficial therapeutic effects, chronic use of BZDs and BZD-related drugs can have health risks. It is associated with rebound insomnia, cognitive impairment and Alzheimer's disease,^{1 2} increased injurious falls³ and (hip) fractures,^{4 5} traffic accidents⁶ and increased (overdose) mortality.^{7 8} In patients with substance abuse disorders it is associated with dependence, and risk of misuse and abuse.⁹ International clinical guidelines therefore state that the pros and cons need



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to be carefully weighed when first prescribing a BZD to an individual patient and to restrict use to a short period. Despite increased awareness regarding risks associated with chronic use and recommendations to limit use in guidelines, these BZDs are among the most used medicines in developed countries.¹⁰ Based on data of all health insurers the prevalence of BZD use in the Netherlands was 10.75% in 2007; 1 758 665 users filled 11 753 939 prescriptions.¹¹ Kurkoo *et al* estimated the prevalence of long-term use in the general population to be about 3%, based on a systematic review of register-based studies on BZD use published in 1993–2014.¹² The relative proportion of chronic use in adult BZD users ranges from 6% to 76%.¹² Chronic and excessive use became a public health issue and led to multiple campaigns to both reduce the prescription of these drugs and to stimulate discontinuation of chronic treatment.¹³

Worldwide different strategies have been used to reduce the use of BZDs.^{13–17} One of these strategies is to influence behaviour of patients, physicians and/or pharmacists by introduction of financial incentives, like pay for performance, copayments and restriction or termination of reimbursement.¹⁸ Since January 2009, BZDs are no longer reimbursed when used as anxiolytic, hypnotic or sedative in the Netherlands. The purpose of this policy change was to reduce chronic use and lower healthcare expenses. Coverage remained in case of epilepsy, palliative sedation and multiple psychiatric disorders, under the condition that the physician considered that no alternative treatment was suitable for the patient at hand. Before the introduction of this policy change, the Dutch Ministry of Healthcare calculated an annual net saving of €70 million for the years 2009–2013.¹⁹

Hoebert *et al* and Kollen *et al* studied the short-term effects of this reimbursement restriction in general practice. They showed that the policy change has led to a moderate decrease in BZD use in patients with newly diagnosed anxiety or sleeping disorder in general practice in 2009,²⁰ a decrease in the mean number of prescription days and a slight decrease in the number of long-term users in 2009 and 2010.²¹ However, the focus of their study was on newly diagnosed anxiety and sleeping disorders, general practice and the short-term effect of the policy change.

This study is an analysis of the reimbursement restriction in the Netherlands on the initial and long-term effects on the use of BZDs, in which the use is compared before and after the implementation of reimbursement restriction in January 2009.

METHODS

Setting

Data for this study were obtained from the PHARMO Outpatient Pharmacy Database (www.pharmo.nl). This database comprises general practitioner (GP) and specialist-prescribed medication dispensed by the outpatient pharmacy. The dispensing records include information

on type of product, date, strength, dosage regimen, quantity, route of administration and prescriber specialty. Records are coded according to the WHO Anatomical Therapeutic Chemical (ATC) Classification System. Outpatient pharmacy data cover a catchment area representing about 3.6 million residents in 2015; almost a quarter of the Dutch population. Patient information includes gender, year of birth and socioeconomic status. Each patient is identified with an anonymous unique patient identification code that enables the observation of patient drug use in time.

In the Netherlands the coverage of pharmaceutical care is regulated by the Health Insurance Act.²² The Ministry of Health, Welfare and Sport and the Healthcare Institute of the Netherlands decide which drugs fall under the mandatory health insurance package. Registered medicines have to be assessed before they can be included in the Medicines Reimbursement System (GVS). Medicines listed in the GVS are fully or partially reimbursed by health insurers. Once a year the Ministry of Health, Welfare and Sport evaluates and actualises the list in order to keep healthcare affordable.

This research was done without patient involvement. Patients were not invited to comment on the study design and were not consulted to develop patient-relevant outcomes or interpret the results. Patients were not invited to contribute to the writing or editing of this document for readability or accuracy.

Study population

All medication records of a random 10% selection of all patients included in PHARMO who received at least one prescription for an oral BZD in the time period between January 2002 and August 2015 were collected and included. Due to the exceeding of the maximum allowed quantity of records by PHARMO a 10% sample was made available for this study. In the Netherlands, the following oral BZDs were available and dispensed during the study period: anxiolytics (N05BA: diazepam, chlor-diazepoxide, oxazepam, clorazepate, lorazepam, bromazepam, clobazam, alprazolam), hypnotics and sedatives (N05CD: flurazepam, nitrazepam, flunitrazepam, lormetazepam, temazepam, midazolam and N05CF: zopiclone, zolpidem) and antiepileptics (N03AE01: clonazepam).

Outcome measures

To estimate the effect of the policy change on the use of BZDs on population and patient level, the following four outcome measures were defined: dispensing volume, incidence, intensity and discontinuation of BZD use.

Volume of dispensed BZDs: dispensed prescriptions and dispensed defined daily doses

First, to estimate the effect of the policy change on the total volume of BZD use, two outcome measures were chosen: the total number of dispensed prescriptions and the total amount of dispensed defined daily doses (DDD). The DDD is the assumed average maintenance dose per

day for a drug used for its main indication in adults defined by the WHO. The purpose of the ATC/DDD system is to serve as a tool for drug utilisation monitoring and research in order to improve quality of drug use.

The total number of dispensed prescriptions and the total amount of dispensed DDDs per month were calculated. To correct for changes in the size of the dynamic population over time, 10% of the total number of medication users present in de PHARMO-database was calculated per month. This figure is then used as denominator to calculate the monthly *dispensing rate* and *DDD rate* per 1000 medication users.

$$\text{dispensing rate} = \frac{\begin{array}{c} \text{[number of dispensed prescriptions} \\ \text{in 10\% sample]} \\ \text{per month} \end{array}}{10\% * \begin{array}{c} \text{[number of medication users present} \\ \text{in PHARMO database]} \\ \text{per month} \end{array}} * 1.000$$

$$\text{DDD rate} = \frac{\begin{array}{c} \text{[sum of dispensed DDD} \\ \text{in 10\% sample]} \\ \text{per month} \end{array}}{10\% * \begin{array}{c} \text{[number of medication users present} \\ \text{in PHARMO database]} \\ \text{per month} \end{array}} * 1.000$$

Incidence of BZD use

Second, the incidence of BZD use was assessed. Incidence of use was defined as the absence of BZD use within 3 months before a prescription was dispensed. The total number of medication users present per month in de PHARMO-database was used to calculate the incidence. The incidence therefore reflects the frequency of starting a BZD per month.

Intensity of prevalent BZD use: incidental, regular and chronic use

Third, the intensity of BZD use was assessed among prevalent users. Prevalent use was defined as the presence of at least one dispensed prescription in a calendar year. For each prevalent BZP user, the BZD use was classified as 'incidental', 'regular' and 'chronic' based on the number of days of use within 365 days after the first dispensed prescription in a calendar year; that is, 1–30 days, 31–180 days and more than 180 days. The intensity of each prescription was calculated using the dispensed quantity and the dose regimen. In case of missing data, the duration of prescriptions was set on 30 days, which is the maximum dispensed quantity for BZDs in the Netherlands.

Discontinuation of BZD

Fourth, the treatment discontinuation of all prevalent BZD users was assessed. Discontinuation was defined as the absence of a BZD prescription within 91 days after the theoretical end date of the last BZD prescription. Kaplan-Meier survival curves were constructed to visualise the

Table 1 Possible effects of policy change in general

Effect type	Level effect	Delta slope (trend)
Initial and long-term effects	Decrease	Negative
Initial effect only	Decrease	No change
Initial effect and long-term opposite effects	Decrease	Positive
Long-term effect only	No change	Negative
No effect	No change	No change

difference in time to discontinuation between patients starting BZD use in the years 2004–2008 (before policy change) and 2009–2013 (after policy change).

Data analysis

The primary statistical model for the effect of the policy change on the dispensing rate, DDD rate, incidence and prevalence of BZD use was segmented regression, which is commonly used for estimating the effects of interventions in interrupted time series (ITS) studies.^{23–25} The Cochrane Collaboration Guideline on ITS was used, applying a segmented regression model.²⁶ Because the announcement of the policy change in September 2008 could have influenced the number of filled prescriptions due to anticipation, the data of the last 3 months before, and the first month after the intervention were excluded from the models. For methodological reasons, it is not possible to calculate the prevalence of initial, regular and chronic use for the years 2014 and 2015. A least squares regression line of predicted values is fitted in each segment, with the assumption of a linear trend over time. The goodness of fit of the models is provided with the R² value.

In an ITS model, policy effects can be measured as a change in the slopes before and after policy, indicating the monthly trends, and the change in level after the implementation of the policy. Based on this analysis, [table 1](#) gives a summary of possible effects from statistical analysis.

Cox proportional hazards analysis was applied to estimate the strength of the association between the 5-year period before and after policy and the risk of discontinuation, and was expressed as an HR with 95% CIs. Observations were censored at the 3 months before and at the 1 month after the policy change. Age, sex and socioeconomic status were considered potential confounders. Confounders resulting in a change of more than 10% in the risk estimate were included in the final model.

A sensitivity analysis was conducted to assess the influence of our definition of BZD discontinuation by expanding the 91-day period to 182 and 365 days and by defining incident as not having dispensed a BZD in the 182 and 365 days before.

SAS Enterprise Guide V.7.1 (SAS Institute) and FoxPro V.6.0 (Microsoft, Redmond, Washington) were used for

data extraction and analysis. Statistical analyses were performed in SPSS V.24 (SPSS).

RESULTS

Sample characteristics

A total of 2500800 filled prescriptions for oral BZD for 128603 unique patients were identified and included. Absolute sample characteristics per year are shown in [table 2](#). Due to the dynamic character of the size of this database throughout the years the absolute number of patients and prescriptions fluctuates. The number of female BZD users is about twice as large as the number of male users. This ratio does not change substantially over time. The proportional use between the different age groups slowly shifts over time to an increase in the percentage of users over 65 years.

Effect analysis

The effect of the policy on the volume; dispensing rate and on the dispensed DDD rate

[Figure 1A](#) shows the *dispensing rate* trend for BZDs, with a distinction between raw and estimated trends and upper and lower confidence limits ($R^2=0.75$). At the beginning of the study the estimated *dispensing rate* was 114.4 (95% CI 102.9 to 125.8) per 1000 medication users per month. The estimated *dispensing rate* decreased by 12.5% (95% CI 9.0% to 15.9%) from 107.2 to 93.8 in January 2009 just after the reimbursement restriction, which shows that the level of the filled prescription rate changed significantly. In contrast, prepolicy trend (-0.085 , $p=0.002$) was not significantly different from the postpolicy trend (-0.063 , $p=0.029$); the monthly trend change after policy was 0.022 (95% CI -0.056 to 0.100) per 1000 medication users. Therefore, a long-term effect was lacking, so the policy had only an initial effect on the dispensing rate.

[Figure 1B](#) presents the monthly *DDD rate*, with a distinction between raw and estimated trends and upper and lower confidence limits ($R^2=0.92$). At the beginning of the study the estimated *DDD rate* was 2274 (95% CI 2072 to 2476) per month per 1000 medication users. The *DDD rate* decreased significantly by 15.1% (95% CI 11.4% to 17.3%) from 1811 to 1538 in January 2009, just after the reimbursement restriction. The prepolicy trend (-5.85 , $p<0.001$) was significantly different from the postpolicy trend (-2.82 , $p<0.001$); the monthly trend change after policy was 3.033 (95% CI 1.592 to 4.474) per 1000 medication users. Therefore, the policy had, according to our definition, an initial effect and a long-term opposite effect on the *DDD rate*.

The effect of the policy on incidence of BZD use

[Figure 2A](#) shows the monthly incidence of BZD use, with a distinction between raw and estimated trends and upper and lower confidence limits ($R^2=0.77$). At the beginning of the study the estimated incidence in April 2002 was 12.9 (95% CI 11.1 to 14.7) patients per month per 1000 medication users. The estimated incidence decreased

significantly by 14.7% (95% CI 9.6% to 19.8%) or from 12.0 to 10.24 per 1000 medication users in January 2009, just after the reimbursement restriction. The trend after policy (-0.028 , $p<0.001$) was significantly different from the trend before policy change (-0.011 , $p=0.019$); the monthly trend change after policy was -0.017 (95% CI -0.031 to 0.003) per 1000 medication users. Therefore, the policy change had an initial and long-term effect on the incidence of BZD use.

The effect of the policy on the intensity of BZD use: prevalence of incidental, regular and chronic use

[Figure 2B](#) shows the trend and level effect of the prevalence of incidental ($R^2=0.97$), regular ($R^2=0.98$) and chronic ($R^2=0.98$) use, with a distinction between raw and estimated trends and upper and lower confidence limits. In 2002, the estimated prevalence per 1000 medication users was 78.1 (95% CI 73.7 to 82.5) for incidental, 59.9 (95% CI 56.8 to 63.1) for regular and 60.2 (95% CI 57.4 to 63.1) for chronic use. Before policy there was an estimated annual reduction of -0.700 ($p=0.018$) for regular and -1.052 ($p=0.004$) for long-term use. The estimated prepolicy trend shows an increase in the prevalence of incidental use from 2002 to 2008 (0.888, $p=0.039$). The absolute-level effects for incidental use are -15.0 (95% CI -20.2 to 9.9), for regular use -11.1 (95% CI -14.5 to 7.7) and for chronic use -8.4 (95% CI -12.1 to 4.7) per 1000 medication users. The relative-level effects are shown in [table 3](#). The postpolicy trend for incidental use (-2.736 , $p=0.002$) was significantly different from the trend before policy (0.888, $p=0.039$); the yearly trend change after policy was -3.624 (95% CI -4.996 to 2.252) per 1000 medication users. Therefore, the policy had an initial and long-term effect on the prevalence of incidental use. The postpolicy trends for regular (-1.003 , $p=0.034$) and chronic (-0.915 , $p=0.069$) use were not significantly different from the prepolicy trends; the trend changes after policy were -0.304 (95% CI -1.204 to 0.596) for regular and 0.136 (95% CI -0.858 to 1.130) for chronic use. Therefore, the policy change had only an initial effect on the prevalence of regular and chronic use.

[Table 3](#) summarises the effects of the policy on the volume (dispensing rate and dispensed DDD rate), the incidence and the intensity (prevalence of incidental, regular and chronic use) of BZD use, as previously described.

The effect of the reimbursement restriction on the discontinuation of BZD treatment

The Kaplan-Meier curves for the 5-year period before and after policy change, illustrating the time to discontinuation after the start date of the first filled BZD prescription, showed no visual difference (see online supplementary figure 1).

Cox proportional hazards analysis was used to compare the risk of discontinuation between patients starting BZDs before (2004–2008) and after (2009–2013) the reimbursement restriction. A small but significant difference

Table 2 Sample characteristics

	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015 (August)
Characteristics of dispensed prescriptions														
Dispensed prescriptions	118215	150613	163114	192420	200067	201962	213090	182760	189483	193409	195596	193845	187278	118948
Dispensed DDDs	1 473297	3 146900	3 338348	3 882 785	3 953726	3 915009	3 854070	3 001910	3 187987	3 028009	3 008484	2 954879	2 845929	1 789140
Mean dispensed DDD	12.46	20.89	20.47	20.18	19.76	19.38	18.09	16.43	16.82	15.66	15.38	15.24	15.2	15.04
DDD/dispensed prescription	6.40	6.52	6.49	6.57	6.61	6.50	6.96	6.87	7.18	7.48	7.85	7.90	7.99	6.79
Patient characteristics														
Overall patients (n)	18467	23083	25131	29297	30277	31054	30610	26612	26389	25859	24904	24525	23453	17530
Sex														
Male (%)	33.7	33.9	34.1	34.2	33.5	33.4	33.1	33.3	33.2	33.4	33.1	33.4	33.6	32.1
Female (%)	66.3	66.1	65.9	65.8	66.5	66.6	66.9	66.7	66.8	66.6	66.9	66.6	66.4	67.9
Age														
<18 years (%)	0.4	0.5	0.5	0.6	0.7	0.7	0.7	0.7	0.7	0.7	0.7	0.7	0.7	0.7
18–44 years (%)	25.9	25.9	25.8	25.4	24.8	24.5	24.5	23.6	23.5	22.9	22.3	21.6	21.2	18.7
45–65 years (%)	37.9	38.3	38.7	38.7	38.6	38.9	38.5	38.5	38.6	38.4	37.9	37.9	38.0	37.0
66–75 years (%)	16.7	16.3	16.1	16.0	16.4	16.6	16.6	16.3	16.5	17.1	17.6	18.4	18.6	20.2
>75 years (%)	19.2	19.1	18.8	19.3	19.5	19.4	19.8	20.8	20.6	20.8	21.6	21.4	21.5	23.4

DDD, defined daily dose.

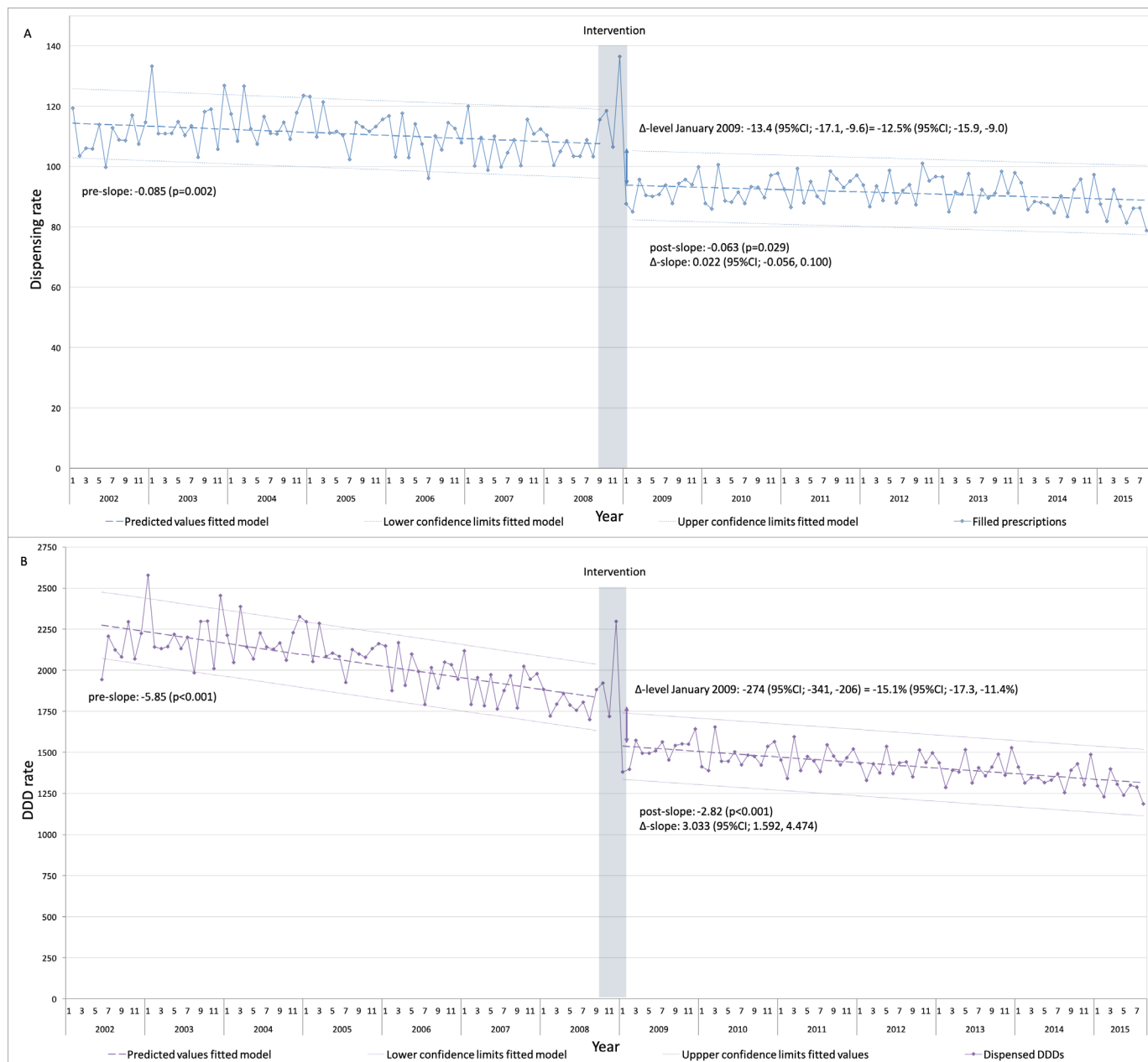


Figure 1 The volume of benzodiazepine (BZD) use over time. (A) The dispensing rate, the total dispensed prescriptions per month per 1000 medication users. (B) The defined daily dose (DDD) rate, the total DDDs dispensed per month per 1000 medication users.

in discontinuation rate was observed (HR=1.013; 95% CI 1.004 to 1.022), indicating that patients who started treatment had a higher chance of discontinuation in the 5-year period after the policy change. Adjusting for age, sex and socioeconomic status as potential confounders did not result in a change of more than 10% of the HR, so they were not included in the final model.

DISCUSSION

This study is the first study which evaluates the initial as well as the long-term effects of the reimbursement restriction policy on BZD use in the Netherlands. Just after the policy change the total number of dispensed

prescriptions as well as dispensed DDD dropped significantly. This effect persisted in the years after the policy change. On the patient level the reimbursement policy had a significant initial effect on the prevalence of both incidental, regular and chronic use. This effect was most prominent in regular use. Only in case of incidental use, there was a stronger monthly decrease in the number of users after the reimbursement restriction policy, indicating a long-term effect. A long-term effect on chronic use is lacking. The policy change had a significant initial effect on the overall incidence of BZD use. The monthly decrease in incidence is greater after policy, which is a strong indication for a long-term effect. When we compare

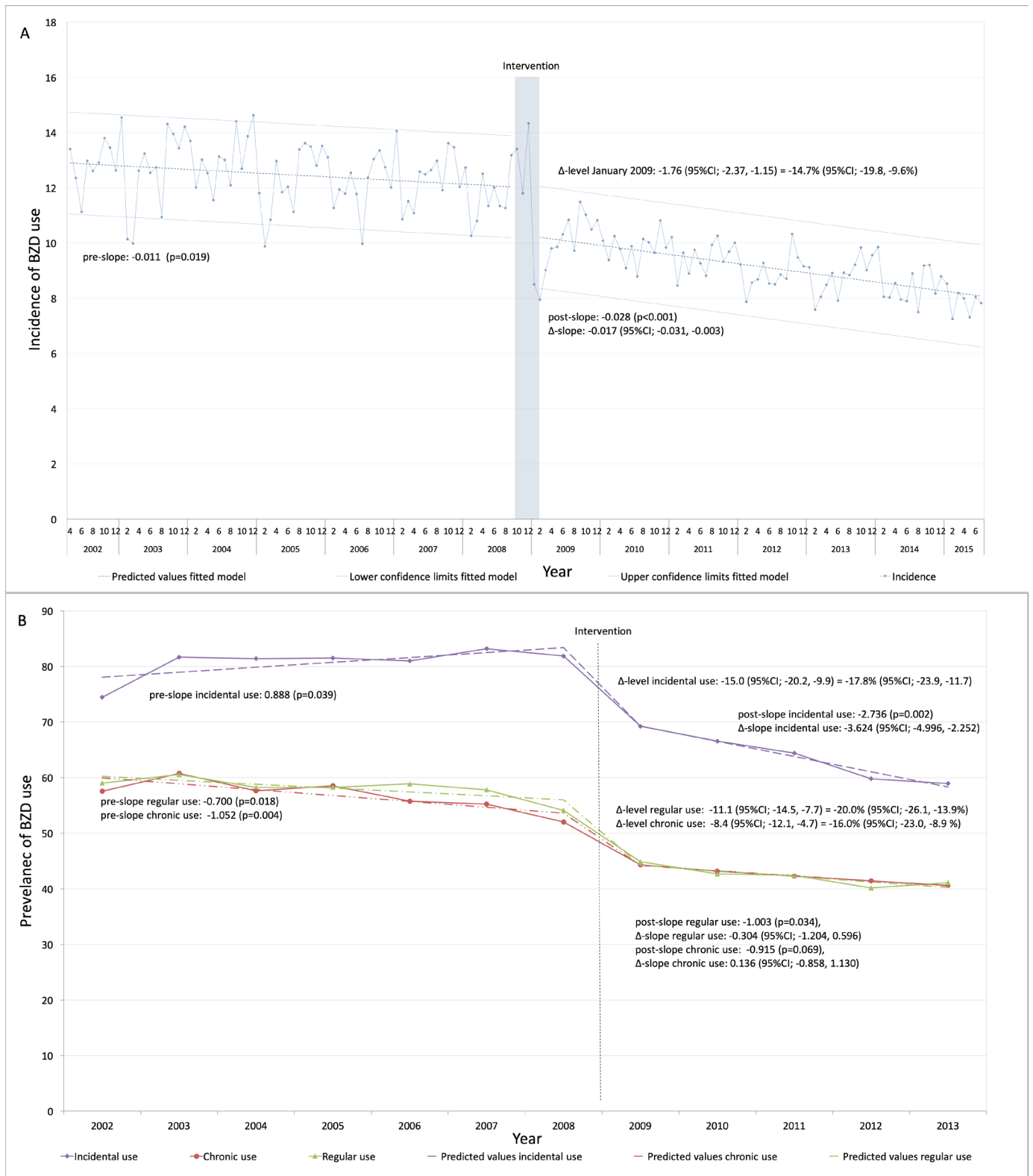


Figure 2 Incidence and intensity of benzodiazepine (BZD) use over time. (A) Incidence per month per 1000 medication users. (B) Prevalence of incidental, regular and chronic use of BZD per calendar year per 1000 medication users.

discontinuation of treatment in the 5-year period before and after the restriction policy the reimbursement had a statistical significant effect (HR 1013). This effect is small and does not appear to be of clinical significance.

Absence of a long-term effect on chronic use could be caused by the proportion of patients with the indication of epilepsy and multiple psychiatric disorders, where reimbursement is maintained. Moreover, BZD dependency

Table 3 Summary of effects of reimbursement restriction of BZDs on the volume, incidence and prevalence of benzodiazepine use

	Effects of policy change	Relative-level effect (95% CI)	Delta slope (trend) (95% CI)
Dispensing rate	Initial effect only	-12.5% (-15.9% to 9.0%)	0.022 (-0.056 to 0.100)
Dispensed DDD rate	Initial effect and long-term opposite effect	-15.1% (-17.3% to 11.4%)	3.033 (1.592 to 4.474)
Incidence	Initial and long-term effects	-14.7% (-19.8% to 9.6%)	-0.017 (-0.031 to 0.003)
Prevalence incidental use	Initial and long-term effects	-17.8% (-23.9% to 11.7%)	-3.624 (-4.996 to 2.252)
Prevalence regular use	Initial effect only	-20.0% (-26.1% to 13.9%)	-0.304 (-1.204 to 0.596)
Prevalence chronic use	Initial effect only	-16.0% (-23.0% to 8.9%)	0.136 (-0.858 to 1.130)

BZD, benzodiazepine; DDD, defined daily dose.

plays a bigger role in this subgroup, which might have an impact on discontinuation rates. Unfortunately, we have no data about indications and reimbursement on the individual patient level. Another mechanism we observe in daily practice can also play a part; prescribers can declare that a patient has multiple psychiatric disorders, although it is not true, only to keep the drug reimbursed. This could occur more frequently in chronic use of BZDs.

Another possible explanation for this phenomenon could be the difference in financial threshold in case of incidental use. The price of BZDs is relatively low, about €0.5–€2 for a month, compared with the pharmacist's service fee for dispensing (€6). The pharmacist's fee for dispensing a first prescription (€12) is about twice as high as for a subsequent prescription, which means that the financial threshold to start exceeds the threshold to continue BZD use. This could be an explanation for the fact that the long-term impact on the incidence and the prevalence of incidental use is positive and lacking in case of both regular and chronic use. Another argument is that it might be more difficult for physicians to stop treatment of individual patients than to prevent initiation, especially when it comes to addictive medication. Medication prescribing is a complex process where behaviour and dynamics regarding initiation are often less complicated than in case of medication cessation.

In addition to reduce chronic use, another aim of the reimbursement restriction was to reduce drug expenditures. Unfortunately, in the PHARMO Outpatient Pharmacy Database, data on costs and coverage status of the dispensed BZD were missing. In 2009, the year after the policy change, about 32% of dispensed prescriptions were covered by health insurance and €79 million was spent on BZDs, of which €56 million was paid by the users themselves.²⁷ While health insurers paid only 32% of all BZDs in 2009, in 2017 that share grew to almost half.²⁸ The absolute number of reimbursed prescriptions increased annually from 3350 million in 2009 to 5033 million in 2017. As a result, the annual net saving of €70 million for the years 2009–2013, calculated by the Dutch Ministry of Healthcare in 2008, has not been achieved. This is a simple way of estimating the intended financial effects

on an annual basis. Only an in-depth (health-economic) analysis of health gain and loss, healthcare utilisation and substitution to other (pharmaco)therapies, like antidepressants or psychotherapy, can reveal the true budget and health impact of the reimbursement restriction policy. The International Task Force on Benzodiazepines suggests that warnings against the negative effects of BZD might be influenced by the pharmaceutical industries in order to promote the use of newer patented alternatives like the serotonin reuptake inhibitors for anxiety.²⁹ These approaches are beyond the scope of this article.

The strengths of this study are the large sample size, the long follow-up, the large amount of time frames and the presence of data of all outpatient prescriptions, including those initiated in secondary care. This makes it possible to focus on short-term and long-term effects. Another strength is use of an ITS design to evaluate the effect of this policy change. This analysis is the golden standard for evaluation of the effects of policy change.

This study has several limitations. First, it is difficult to attribute the observed changes in BZD use to a single intervention, the reimbursement restriction policy, because many confounding factors change in time. However, to our knowledge there were no other health system changes that could be a rival explanation for the observed changes. Second, due to PHARMO regulations, only a 10% random sample of PHARMO database was obtained and analysed. Nevertheless, the statistical significance of our analysis shows that the size of the data set is sufficient. Finally, in this study dispensed prescriptions were used as an approximation of BZD use. This may not always be accurate as in fact not all dispensed prescriptions have led to their consumption.

Not only Dutch policymakers have chosen to apply financial incentives to reduce BZD use. In Italy, exclusion from reimbursement in 1994 was not enough to reduce the use of BZDs. Despite the reimbursement policy the total BZDs sold from 1989 to 1999 showed a 53% increase in Italy.³⁰ Although a French pay-for-performance strategy had a modest effect on the achievement of GPs regarding BZD indicators,¹⁶ it did not lead to the intended decrease of the use of BZDs in 2012.³¹ In a US policy, exclusion

from the Medicare Part D formulary in 2006 led to an immediate and sustained drop of 5% in number of BZD prescriptions filled by seniors.³²

Compared with these policies the effect of the Dutch reimbursement restriction policy is high. In contrast, compared with the effect of the New York State Regulatory action in 1989 that required triplicate forms for BZD prescribing which created a higher bureaucratic threshold for BZD prescribing which resulted in a 60% reduction in BZD use,³³ the effect of the reimbursement restriction in the Netherlands is modest.

A decrease in the use of BZDs does not automatically lead to a reduction of the negative events with which they are associated in the literature. For example, Wagner *et al* showed that this policy, despite this substantial reduction in the use of BZDs, did not lead to a decrease in hip fractures.¹⁵ So, substantial reductions in the use of BZD do not necessary lead to expected clinical benefits.

Not only in case of BZDs are introducing financial incentives used to influence the use of prescription drugs. Green *et al* reviewed the effects of reimbursement restrictions on the use of several drug classes.³⁴ Due to substantial differences in objectives, context and the presence of alternatives, it is difficult to generalise the results and to compare with the outcomes of this study. Green *et al* stated that policy design needs to be based on research quantifying the harm and benefit profiles of target and alternative drugs to avoid unwanted health system and health effects.

When policy changes are so profound and affect so many people, it is the responsibility of the authorities to make a sound assessment and share it with society. This information provides essential insights into 'what to do' and 'what not to do' in drug reimbursement policy. This allows policymakers to learn from previous initiatives and from initiatives abroad, and enables them to deploy reimbursement policy in an evidence-based manner. However, success of reimbursement policy is difficult to predict and highly dependent on purpose and context and a result of the complex interaction between social, cultural, economic and systemic factors. Neglecting this complex dynamic can cause failure of copied successful policies.

Despite these limitations, this study provides us important insights into the effects of the policy on the use of BZDs in the Netherlands and gives direction for follow-up research. Does this policy lead to intended or unintended substitution; the shift to other medicines or therapies or substance abuse? And, are there differences between prescribers, different BZDs, age groups and socioeconomic status? It is also important to investigate whether the decrease in the use of BZDs actually leads to a reduction of the associated adverse events like addiction, fall incidents, hip fractures and the impact on sickness absence and social costs in the Netherlands.

This is the first study with an ITS design on the initial and long-term effects of the 2009 reimbursement restriction policy on BZD use in the Netherlands. The policy change has a significant initial effect on the volume and

incidence of BZD use. In addition to an initial effect, there is a long-term effect in case of incidental use and overall incidence. Although there is a significant initial effect, the long-term effect on the chronic use of BZD, the main purpose of the reimbursement restriction, is lacking.

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