BMJ Open is committed to open peer review. As part of this commitment we make the peer review history of every article we publish publicly available.

When an article is published we post the peer reviewers' comments and the authors' responses online. We also post the versions of the paper that were used during peer review. These are the versions that the peer review comments apply to.

The versions of the paper that follow are the versions that were submitted during the peer review process. They are not the versions of record or the final published versions. They should not be cited or distributed as the published version of this manuscript.

BMJ Open is an open access journal and the full, final, typeset and author-corrected version of record of the manuscript is available on our site with no access controls, subscription charges or pay-per-view fees (<u>http://bmjopen.bmj.com</u>).

If you have any questions on BMJ Open's open peer review process please email <u>editorial.bmjopen@bmj.com</u>

BMJ Open

The use of driving-impairing medicines by the population.

Journal:	BMJ Open
Manuscript ID	bmjopen-2017-017618
Article Type:	Research
Date Submitted by the Author:	03-May-2017
Complete List of Authors:	Gutierrez-Abejón, Eduardo; Gerencia Regional de Salud de Castilla y León, Technical Direction of Pharmaceutical Assistance Herrera-Gómez, Francisco; Universidad de Valladolid, Pharmacology and Therapeutics Criado-Espegel, Paloma; Gerencia Regional de Salud de Castilla y León, Technical Direction of Pharmaceutical Assistance Alvarez, F. Javier; University of Valladolid, Pharmacology & Therapeutics
Primary Subject Heading :	Global health
Secondary Subject Heading:	Public health
Keywords:	drug prescription, drug utilization, risk assessment, accident, traffic, automobile driving, driving-impairing medicines



BMJ Open

The use of driving-impairing medicines by the population.

Eduardo Gutierrez-Abejón¹, Francisco Herrera-Gómez^{2,3}, Paloma Criado-Espegel¹ and F. Javier Álvarez²

¹Technical Direction of Pharmaceutical Assistance, Gerencia Regional de Salud de Castilla y León, Valladolid, Spain.

²Pharmacology, Faculty of Medicine, University of Valladolid, Valladolid, Spain.

³Department of Nephrology, Complejo Asistencial de Zamora, Zamora, Spain.

Correspondence: Professor F. Javier Álvarez, Pharmacology, Faculty of Medicine, University of Valladolid, Ethics Review Board, Hospital Clínico Universitario de Valladolid, Valladolid, Spain. Tel: +34 983 423077; Fax: +34 983 423022; E-mail: <u>alvarez@med.uva.es</u>

Running head: Medicines and driving.

Keywords: Accidents, Traffic. Automobile Driving. Drug Prescriptions. Drug Utilization. Risk Assessment. Driving-impairing medicines.

Word count (excluding the title page, abstract, references, tables, and figures): 2624

Number of tables: 04

Number of figures: 02

ABSTRACT

OBJECTIVE

To assess the use of driving-impairing medicines (DIM) by the general population, with special reference to length of use and concomitant use.

DESIGN

Open cohort study.

SETTING

Year-2015 granted medicines consumption data recorded in the Castile & León medicines dispensation registry were consulted.

PARTICIPANTS

Medicines and DIM consumers from a Spanish population (Castile & León: 2.4 million inhabitants).

EXPOSURES

Medicines and DIM consumption. Patterns of use by age and gender, based on the length of use (acute: 1-7 days, sub-chronic: 8-29 days, and chronic use: \geq 30 days), were of interest. Estimations regarding the distribution of licensed drivers by age and gender were made in order to know the patterns of use of DIM.

RESULTS: DIM were consumed by 34.4% (95%CI 34.3-34.5) of the general population in 2015, more commonly with regularity (chronic use: 22.5% versus acute use: 5.3%), and more frequently by the elderly. On average, 2.3 DIM *per* person were dispensed, particularly to chronic users (2.8 DIM *per* person). Age and gender distribution differences were observed between the Castile & León medicines dispensation registry data and the drivers' license census data. Of all

BMJ Open

DIM dispensed, 83.8% were ATC group N medicines, which were prescribed to 29.2% of the population.

CONCLUSIONS: The use of DIM was frequent in the general population and among drivers in our region. Chronic use was common, but acute and sub-acute use should also be taken into account. The fact that ATC group N medicines were the most consumed highlights the need to improve dispensation tools.

Instation tools.

STRENGTHS AND LIMITATIONS OF THE STUDY

- This study explores the consumption of all driving-impairing medicines and patterns of use by age and gender corresponding to a European population in a comprehensive way.
- The Anatomical Therapeutic Chemical code (ATC) was used for improve transparency and reproducibility of our findings.
- The information covers all dispensed medicines by the public health system in Spain, but not hospital dispensed medications, nor over-the-counter medicines, some of which may be driving-impairing medicines.

BACKGROUND

Driving a motor vehicle is a multifaceted task and requires appropriate cognitive and psychomotor skills (e.g. alertness, concentration, reaction time, visual acuity).¹⁻² Medicines can adversely affect these driving-related skills and, consequently, be a hazard to traffic safety.³⁻⁴ There is increasing awareness that the implementation of appropriate measures to limit the consumption of alcohol and other substances ("illicit" drugs and medicines) while driving may impact on road accident occurrence.⁵ Nevertheless, it is as yet unknown how frequent the consumption of driving-impairing medicines (DIM) in the general population is, or how frequently several of these drugs are consumed concomitantly.⁶

On the one hand, most developed countries perform toxicological analyses on road accident casualties and fatalities, and the presence of illicit drugs and medicines (either used legally or illegally) are detected.^{7,8} On-road tests (at random or on target populations) are used ever more frequently worldwide: the on-site screening devices detect some groups of illicit drugs and some medicines in saliva (oral fluid), confirmation analyses being performed later.^{7,8} In other countries, blood analyses⁹ are carried out rather than screening on saliva. The information from these sources (data on casualties/fatalities and on-road test data) gives only a partial vision of the problem regarding medicines and driving.¹⁰

On the other hand, medicine regulatory agencies do attempt to provide appropriate information to the public concerning the problem: in the European Union, the summary of product characteristics and the package leaflets contain information on medicines that "affect the ability to drive and to use machines".^{11,12} Furthermore, there have been some attempts to categorize the

effects of medicines on driving¹³ and some countries, such as France¹⁴ and Spain,¹⁵ have introduced specific mandatory pictograms; or ancillary warning labels, as in the Netherlands¹⁶ and Australia.¹⁷

The Spanish Law of 2007 (Royal Decree 1345/2007)¹⁵ establishes that newly authorized medicines which may negatively affect fitness to drive or to handle dangerous machinery should include a warning symbol (pictogram) on the outside of the packaging. Since 2011, all medicines that can possibly affect fitness to drive and are commercially available in Spain have this pictogram on the packaging.¹⁸ As of January 2016, a total of 2013 medicinal drugs permitted in Spain had been reviewed, of which 402 (20%) include the pictogram on medicines and driving on the packaging.¹⁹

We have considered these medicines with the pictogram "medicines and driving" on the packaging in Spain as driving impairing medicines or, to be more exact, potentially impairing medicines on driving. In 2016, a national consensus on Medicines and Driving was reached in Spain: to know to what extent the population taking DIM is a priority and to decipher patterns of use for these drugs.²⁰ This does not apply only to motor vehicle drivers and professional drivers, but also the population at large, as well as all road users, including pedestrians and the ever more common cyclists. Thus, the presence of illicit drugs and medicines is also frequently found in pedestrians involved in fatal road accidents.²¹

In our opinion, medicine-screening on the road should not be viewed as punishment of patients, but rather a tool to better inform patients, as well as to prescribe and dispense better. The detection of some medicines in on-road tests has been the object of awareness-raising in public

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

BMJ Open

and health professionals, as well as the subject of campaigns to inform the general public, as in the UK.²² In some countries, for instance the UK.²³ Spain²⁴ or Norway.²⁵ on-road positive cases to medicines are not fined if they were used according to a physician's prescription. Again, it is necessary that the consumption of DIM and their patterns of use should be known in detail, given that there is a shortage of information about this.

Consequently, the aim of our study was to explore the use of DIM by the general population. The consumption of medicines with the pictogram "medicines and driving" was assessed on the basis of our dispensation registry, focusing on concomitant use of these drugs and on their length of use. In addition, estimations were compared with the drivers' license census in order to know the patterns of use among drivers.

METHODS

Study population: CONCYLIA database

Access was provided to the CONCYLIA database to assess the dispensation of granted medicines by the Spanish public health system in Castile & León during 2015.²⁶

Basically, the CONCYLIA database includes information on all medicine dispensations by the public health system, except those dispensed at hospitals, medical prescriptions dispensed through private medicine clinics, and those which do not require a medical prescription ('over the counter' medication). We assessed medicine dispensation per person using the patient identification number: for each person, any dispensation during 2015 was identified (medicinal

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

product, number of doses, data of dispensation, etc). For data protection, the final database provided by the health system was anonymized, and no personal identification was included.

Target population

The population distribution covered by the public health system and the population distribution according to the population census match well: Castile & León had a population of 2,428,901 in December 2015,²⁷ 2,376,717 being covered by the public health system at that time (97.85% of the total).²⁸

As not all persons had a motor vehicle license, calculations were made regarding the distribution of licensed drivers by age and gender, using the Castile & León drivers' license census up to December 2015.²⁹ This was done as no information on medicine use by drivers is recorded in the CONCYLIA database. Therefore, results are presented regarding the general population based on the drivers' license census data (Table 1).

Driving-impairing medicines

As mentioned above, granted medicines in Spain with the pictogram "medicines and driving" were considered as DIM.

In the CONCYLIA database, each one of the medicines, based on the Anatomical Therapeutic Chemical code (ATC), is identified as having such a pictogram or not; this information was taken from the Spanish Medicine Agency, from information updated to February 1st 2016.¹⁹

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

Variables and ethical issues

The following variables describing the consumption of medicines and DIM in Castile & León in

the year 2015 were considered:

i) Yearly frequency of all medicine use;

ii) Yearly frequency of DIM use: acute (1-7 days), sub-acute (8-29 days) and chronic or regular

use (\geq 30 days);

iii) Yearly frequency of daily use of at least one DIM;

iv) Number and means of different DIM taken within 2015.

All analyses were made considering age and gender distribution.

Ethic Review Board approval was obtained (Reference number PI 16-387, approved on March 17th, 2016)

Statistical analysis

All values are given as percentages (frequencies) with a 95% confidence interval (95%CI) or as means \pm standard deviations (SD). For comparisons, the Student's *t* test was used for continuous variables and Pearson's chi-square test for categorical variables. Two-tailed P < 0.05 were considered significant. All statistical analyses were performed using the Statistical Package for Social Sciences (SPSS version 23.0.; SPSS Inc, Chicago, IL).

RESULTS

Descriptive mapping

A total of 48,858,588 medicines were dispensed in 2015. Nearly 3 out of 4 people took a medicinal product in 2015, more females than males (78.4% versus 68.8%, p < 0.05), and this increased in line with age (Table 1).

One out of three (34.4%, 95%CI 34.3-34.5) consumed a DIM in 2015, again more frequent among females (40.3% versus 28.3%, p < 0.05), and this also increased in line with age. A majority needed to use these medicines on a regular basis (chronic use: 22.5%), while the use for a few days or weeks accounted for, respectively, 5.3% and 6.6%, with similar patterns of use by age and gender (Table 1).

However, if the distribution is performed with respect to the drivers' license census, 25.4% (95%CI 25.3-25.43) of people took a DIM in 2015, more males than females (26.5% versus 23.7%, p < 0.05) and mostly regularly (chronic use: 15.3%, 95%CI 15.2-15.32; sub-acute use: 5.96%, 95%CI 5.92-5.99; acute use: 4.14%, 95%CI 4.11-4.18).

Figure 1 and Table 1 show those who used DIM in 2015, their distribution by age and gender, and regarding the drivers' license census. Age trends differ between the sexes: consumption dropped dramatically among female drivers from 60 years of age as compared to male drivers using less DIM over 75 years of age.

BMJ Open

At least one DIM was consumed daily by 5.6% (95%CI 5.52-5.58) of people, and by 3.7% (95%CI 3.67-3.73) of licensed drivers.

On average (Table 2), each person taking DIM took 2.3 medicines (2.1 according to the drivers' license census data). Acute and sub-acute consumers (97.5% and 69.1%, respectively) took just one DIM, while chronic consumers (71.5%) took 2 DIM or more (mean number of DIM use: 2.8). Trends between sexes were similar when the drivers' license census data was analyzed (Table 2).

Types of DIM consumed

Of the 10,862,138 DIM dispensed, 9,102,052 (83.8%) belong to the ATC classification group N (Nervous System), 1,176,864 (10.8%) to the group A (Alimentary Tract and Metabolism) and 160,631 (1.5%) to the group R (Respiratory System). ATC group N medicines were prescribed to 29.2% of the population (21.3% regarding the drivers' license census), group A medicines to 5.4% (4% for drivers), and group R medicines to 4% (2.3% for drivers). (Table 3)

Interestingly, ATC groups N, A and R were more frequently prescribed to females than males, all people considered. When considering licensed drivers, the trends showed no differences. Table 4 shows those DIM used for a few days or weeks and chronically.

DISCUSSION

A detailed description of the consumption of DIM in the general population from Castile & León in 2015 is provided. DIM were consumed by 34.4% (95%CI 34.3–34.5) of the general population, and more commonly on a regular basis (22.5%). However, the use for a few days (5.3%) or a few weeks (6.6%) should not be neglected. The consumption of DIM increased in line with age. Acute and sub-acute consumers took at least one DIM, and chronic users took nearly three. Of all DIM dispensed, 83.8% belong to the ATC classification group N (Nervous System), which were dispensed to 29.2% of the population. Similar trends were found regarding the distribution of licensed drivers by gender, but not by age.

The DRUID [1] project provides information on the prevalence of use in Europe of some types of medicines randomly detected in drivers. Of all positive matches (1.36%), benzodiazepines (0.9%), Z-drugs (0.12%), and opioids (0.35%) were confirmed. Furthermore, there is information for other developed countries on the consumption of alcohol, illicit drugs and certain medicines by people injured/killed in road traffic accidents.⁷⁻¹⁰ Although progress has been made in understanding this social and medical problem of driving under the effects of medicines, available data allows just a partial vision, as only a few groups of DIM (mainly psychotropic drugs) have been analyzed in blood and oral fluid specimens from drivers. There has also been an attempt to estimate DIM consumption based on dispensed medicines,⁶ or using driver consumption surveys.³⁰ Our study provides a detailed overview of all DIM used by the general population and, to our knowledge, this is the first work in this matter.

BMJ Open

One out of 3 used a DIM in 2015. Importantly, acute users represented a sizeable proportion of all drivers consuming DIM (5.3%). The effect of medications on driving is more relevant in the first days of use.^{31,32} Drivers consuming DIM for a few days might therefore be the most affected, particularly those taking more than one medicine, and this must be taken into account. In addition, multiple daily dosing is an important factor to consider,^{6,33} especially for drivers over 50 years old.³³

More than 2 DIM were dispensed (drivers and non-drivers), particularly to chronic users who took nearly three. In addition, approximately 6% of people consumed at least one DIM daily during the year 2015. Impairment on driving seems to diminish with chronic/stable DIM use,³² probably due to tolerance.³⁴ However, clinical explorations of fitness to drive under the effects of DIM should be performed.³⁴ Tolerance is a problem that has not been completely assessed and nor has what happens when more than one medicine is consumed. A higher prevalence of regular and daily use of DIM are not uncommon in Spain and other developed countries. So our results provide an epidemiological view of the current impact of medicine use patterns which highlight the importance of daily regimens, as well as for elderly acute users.

Our results show that some types of medicines are more prescribed than others. ATC group N medicines were prescribed with predilection, mostly to females. This is corroborated by the study of Ravera *et al.*.⁶ The finding of frequent DIM use is not surprising, as 20% of granted medicines (402 out of 2,013) in Spain are DIM (with the pictogram "medicines and driving" on the packaging). In addition, 83.8% of dispensed DIM in Castile & León were ATC group N medicines (178 out of 402). In this context, mandatory pictograms and warning labels contribute to awareness of DIM consumption risks for consumer engagement,^{35,36} the noticeability of these

medication warnings being a challenging task.³⁷ Furthermore, there are initiatives worldwide for refining information on the risk categorization of drugs^{1,13,38,39} that must be implemented in dispensing support tools (software)⁴⁰ for a better prescription/dispensation of DIM.

Our study provides in detail which DIM are consumed and how. We thus answer the objectives of the Spanish consensus on Medicines and Driving reached recently. Giving clearer information about the influence of medicines on driving in order to sensitize health professionals and the general population on the negative effects of DIM is a priority.²⁰ Our results stress the need to improve the communication of DIM risks, in line with recent requirements. Nevertheless, DIM risk communication is a complex clinical, methodological and epidemiological challenge, and the "boosters" (warning label methods, dispensation software, information campaigns, etc.) should be cautiously implemented in key steps, the main objective being to minimize road accidents. Again, detailed knowledge of the use of DIM is a priority.

This study has some limitations. The health system in Spain is public and free, and we used the data from a medicine dispensation registry, which implies that the information covers all dispensed medicines within such a system, but not hospital dispensed medications, nor over-the-counter medicines, some of which may have the Spanish pictogram. Data is presented regarding the general population, not only drivers. However, even pedestrians and cyclists could be involved in road traffic accidents, DIM use being a possible cause. In the CONCYLIA database, no information on medicine use by drivers is recorded, and calculations were made regarding the distribution of licensed drivers. This should be taken into account, as the distribution of drivers in other countries or regions could be different, especially because information is not available on the extent to which drivers with a license drove vehicles.

CONCLUSION

The use of driving-impairing medicines was frequent in the general population from Castile & León in 2015. Chronic use (30 days or more) was common, but acute use (1-7 days) and subacute use (8-29 days) must not be observed with indifference, because they might be the most relevant regarding the communication of DIM consumption risks. ATC group N medicines were the most prescribed and this fact stresses the need to ameliorate prescription and dispensation tools.

Contributors

F.J.A. conceived the study design. E.G.-A. conducted the study. E.G.-A., F.H.-G., P.C.-E. and F.J.A. analyzed the data, contributed to the interpretation of the results, and wrote the manuscript. All authors reviewed and approved the final version of the manuscript.

Funding

This work has been supported by the Dirección General de Tráfico (DGT) reference: SP IP2017-02120.

Competing Interests

No competing interests.

No additional data are available

<text>

REFERENCES

- Schulze H, Schumacher M, Urmeew R, Auerbach K, Alvarez FJ, Bernhoft IM, de Gier JJ, Hagenzieker M, Houwing S, Knoche A, Pilgerstorfer M, Zlender B. Driving under the influence of drugs, alcohol and medicines in europe — findings from the DRUID project. Lisbon: EMCDDA; 2012. <u>http://www.emcdda.europa.eu/publications/thematic-papers/druid</u> (Accessed 9 Jan 2017)
- Ramaekers JG. Drugs and Driving Research in Medicinal Drug Development. Trends Pharmacol Sci. 2017 Mar 7. pii: S0165-6147(17)30022-6.
- Ravera S, van Rein N, de Gier JJ, de Jong-van den Berg LT. Road traffic accidents and psychotropic medication use in The Netherlands: a case-control study. Br J Clin Pharmacol 2011; 72: 505-13.
- Orriols L, Delorme B, Gadegbeku B, Tricotel A, Contrand B, Laumon B, Salmi LR, Lagarde E; CESIR research group.. Prescription medicines and the risk of road traffic crashes: a French registry-based study. PLoS Med 2010; 16: e1000366.
- 5. WHO. Drug use and road safety: a policy brief. Geneva, Switzerland, World Health Organization, 2016. <u>http://www.who.int/iris/handle/10665/249533</u> (Accessed 9 Jan 2017)
- Ravera S, Hummel SA, Stolk P, Heerdink RE, de Jong-van den Berg LT, de Gier JJ. The use of driving impairing medicines: a European survey. Eur J Clin Pharmacol 2009; 65: 1139-47.
- Gómez-Talegón T, Fierro I, González-Luque JC, Colás M, López-Rivadulla M, Javier Álvarez F. Prevalence of psychoactive substances, alcohol, illicit drugs, and medicines, in Spanish drivers: a roadside study. Forensic Sci Int 2012; 223: 106-13.

8. Simonsen KW, Steentoft A, Hels T, Bernhoft IM, Rasmussen BS, Linnet K. Presence of

- psychoactive substances in oral fluid from randomly selected drivers in Denmark. Forensic Sci Int 2012; 221: 33-8. 9. Bezemer KD, Smink BE, van Maanen R, Verschraagen M, de Gier JJ. Prevalence of medicinal drugs in suspected impaired drivers and a comparison with the use in the general Dutch population. Forensic Sci Int 2014; 241: 203-11. 10. Rudisill TM, Zhao S, Abate MA, Coben JH, Zhu M. Trends in drug use among drivers killed in U.S. traffic crashes, 1999-2010. Accid Anal Prev 2014; 70: 178-87. 11. Council Directive 83/570/EEC of 26 October 1983 amending Directives 65/65/EEC, 75/318/EEC and 75/319/EEC on the approximation of provisions laid down by law, regulation or administrative action relating to proprietary medicinal products. OJ L 332, 28.11.1983, p. 1–10 (DA, DE, EL, EN, FR, IT, NL). http://data.europa.eu/eli/dir/1983/570/oj (Accessed 9 Jan 2017) 12. European Commission. Enterprise and Industry Directorate-GeneralA guideline on summary of product characteristics (SmPC) 2009. [online]. Available at http://ec.europa.eu/health/files/eudralex/vol-2/c/smpc guideline rev2 en.pdf and http://ec.europa.eu/health/documents/eudralex/vol-2/index en.htm (Accessed 9 Jan 2017) 13. de Gier JJ, Alvarez FJ, Mercier-Guyon C, Verstraete AG. Prescribing and dispensing guidelines for medicinal drugs affecting driving performance. In: Drugs, Driving and Traffic Safety, eds Verster JC, Pandi-Perumal SR, Ramaekers JG, de Gier JJ. Basel: Birkaeuser Verlag AG, 2009: 121-34.
 - 14. Ministère de la Santé et des Solidarités. Direction Generale de la Santé. Arrêté du 18 Juillet 2005 pris pour l'application de l'article R.5121-139 du code de la santé publique et relative à l'opposition d'un pictogramme sur le conditionnement extérieur de certain

BMJ Open

	médicaments et produitsJournal Officiel de la République Française. Août 2005
	(SAN/P0522726A) (Accessed 9 Jan 2017)
15	. Real Decreto 1345/2007, de 11 de octubre, por el que se regula el procedimiento de
	autorización, registro y condiciones de dispensación de los medicamentos de uso humano
	fabricados industrialmente. pp. 45652–45698. BOE de 7 de Noviembre de 2007.
	http://www.boe.es/diario_boe/txt.php?id=BOE-A-2007-19249 (Accessed 9 Jan 2017)
16	. Patrício Monteiro S. Driving-impairing Medicines and Traffic Safety: Patient's
	Perspectives (PhD thesis). University of Groningen, 2014.
17	. Jomaa I, Odisho M, Cheung JM, Wong K, Ellis JG, Smyth T, et al. Pharmacists'
	perceptions and communication of risk for alertness impairing medications. Res Social
	Adm Pharm. 2017 Jan 10. pii: S1551-7411(16)30377-1.
18	. Agencia Española de Medicamentos y Productos Sanitarios. Medicamentos y
	Conduccion. http://www.aemps.gob.es/industria/etiquetado/conduccion/home.htm
	(Accessed July 2016)
19	. Agencia Española de Medicamentos y Productos Sanitarios. Medicamentos y
	Conduccion: Listados de principios activos por grupos ATC* e incorporación del
	pictograma de la conducción
	http://www.aemps.gob.es/industria/etiquetado/conduccion/listadosPrincipios/home.htm
	(Accessed July 2016)
20	. Documento de consenso sobre medicamentos y conducción en España: información a la
	población general y papel de los profesionales sanitarios.
	http://www.msssi.gob.es/profesionales/saludPublica/prevPromocion/docs/Medicamentos_
	<pre>conduccion_DocConsenso.pdf (Accessed May 2016)</pre>

 Instituto Nacional de Toxicología y Ciencias Forenses. Víctimas mortales en accidentes de tráfico: memoria 2016. Ministerio de Justicia.

https://administraciondejusticia.gob.es/paj/PA_WebApp_SGNTJ_NPAJ/descarga/Memor ia%20Trafico%20INTCF202014.pdf?idFile=00359cf9-26d5-4d33-96ea-ec9703c78470

(Accessed May 2016).

22. GOV:UK. Drugs and driving: the law. Collection Drug driving.

https://www.gov.uk/drug-driving-law (Accessed 9 Jan 2017)

https://www.gov.uk/government/collections/drug-driving#table-of-drugs-and-limits

(Accessed 9 Jan 2017)

23. Department for Transport. Guidance for healthcare professionals on drug driving.

London: Department for Transport; 2014.

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/325275/he althcare-profs-drug-driving.pdf (Accessed 9 Jan 2017)

- 24. Álvarez FJ, González-Luque JC, Seguí-Gómez M. Drugs, Substance Use Disorder and Driving: Intervention of Health Professionals in the Treatment of Addictions. Adicciones 2015; 27: 161-7.
- 25. Norwegian Ministry of Transport and Communications. Driving under the influence of non-alcohol drugs-legal limits implemented in Norway. Oslo: Norwegian Government Security and Service Organization; 2014.

https://www.regjeringen.no/globalassets/upload/sd/vedlegg/brosjyrer/sd_ruspavirket_kjori ng_net.pdf (Accessed 9 Jan 2017)

26. CONCYLIA. Sistema de Información de Farmacia. Gerencia Regional de Salud de Castilla y León.

BMJ Open

27. Instituto Nacional de Estadística. Cifras oficiales de población de los municipios
españoles: Revisión del Padrón Municipal.
http://www.ine.es/dyngs/INEbase/es/operacion.htm?c=Estadistica_C&cid=125473617701
<u>1&menu=resultados&secc=1254736195458&idp=1254734710990</u> (Accessed 15 Jun
2016)
28. Tarjeta Sanitaria de SACYL. Recursos y Gestión Poblacional. Gerencia Regional de
Salud. 2015.
29. Ministerio del Interior. Dirección General de Tráfico. Estadísticas e Indicadores. Permisos
de conducción. http://www.dgt.es/es/seguridad-vial/estadisticas-e-indicadores/permisos-
conduccion/ (Accessed 15 Jun 2016).
30. Del Río MC, Alvarez FJ. Medication and fitness to drive. Pharmacoepidemiol Drug Saf
2003; 12(5): 389-94.
31. Barbone F, McMahon AD, Davey PG, Morris AD, Reid IC, McDevitt DG, MacDonald
TM. Association of road-traffic accidents with benzodiazepine use. Lancet 1998; 352:
1331-6.
32. Wilhelmi BG, Cohen SP. A framework for "driving under the influence of drugs" policy
for the opioid using driver. Pain Physician 2012; 15: ES215-30.
33. Monárrez-Espino J, Laflamme L, Elling B, Möller J. Number of medications and road
traffic crashes in senior Swedish drivers: a population-based matched case-control study.
Inj Prev 2014; 20(2): 81-7.
34. Schumacher MB, Jongen S, Knoche A, Petzke F, Vuurman EF, Vollrath M, Ramaekers
JG. Effect of chronic opioid therapy on actual driving performance in non-cancer pain
patients. Psychopharmacology (Berl) 2017; 234: 989-99.

- 35. Monteiro SP, Huiskes R, Van Dijk L, Van Weert JC, De Gier JJ. How effective are pictograms in communicating risk about driving-impairing medicines? Traffic Inj Prev 2013; 143: 299-308.
- 36. Emich B, van Dijk L, Monteiro SP, de Gier JJ. A study comparing the effectiveness of three warning labels on the package of driving-impairing medicines. Int J Clin Pharm 2014; 366: 1152-9.
- 37. Smyth T, Sheehan M, Siskind V, Mercier-Guyon C, Mallaret M. Consumer perceptions of medication warnings about driving: a comparison of French and Australian labels. Traffic Inj Prev 2013; 14: 557-64.
- 38. DRUID, 2012. DRUID Driving Under the Influence of Drugs, Alcohol and Medicines. Final Report. <u>www.druid-</u>
 project ov/DRUID/EN/Discomination/downloads and links/Final Report pdf2, blob=n

project.eu/DRUID/EN/Dissemination/downloads_and_links/Final_Report.pdf?_blob=pub licationFile (Accessed 9 Jan 2017)

- 39. International Council on Alcohol, Drugs and Traffic Safety. Utrecht: ICADTS; 2001.
- 40. Legrand SA, Boets S, Meesmann U, Verstraete AG. Medicines and driving: evaluation of training and software support for patient counselling by pharmacists. Int J Clin Pharm 2012; 34: 633-43.
- Emich B, van Dijk L, Monteiro SP, de Gier JJ. A study comparing the effectiveness of three warning labels on the package of driving-impairing medicines. Int J Clin Pharm 2014; 36: 1152-9.
- 42. Goyal RK, Rajan SS, Essien EJ, Sansgiry SS. Effectiveness of FDA's new over-thecounter acetaminophen warning label in improving consumer risk perception of liver damage. J Clin Pharm Ther 2012; 37: 681-5.

BMJ Open

2	
3	
1	
4	
5	
6	
7	
, Q	
0	
9	
10	
11	
12	
12	
13	
14	
15	
16	
47	
17	
18	
19	
20	
20	
21	
22	
23	
24	
2-7	
25	
26	
27	
28	
20	
29	
30	
31	
32	
22	
33	
34	
35	
36	
27	
37	
38	
39	
40	
41	
42	
43	
44	
15	
40	
46	
47	
48	
40	
49	
50	
51	
52	
52	
55	
54	
55	
56	
57	
ວ/ = -	
58	
59	
60	

43. McCarthy DM, Davis TC, King JP, Mullen RJ, Bailey SC, Serper M, Jacobson KI
Parker RM, Wolf MS. Take-Wait-Stop: a patient-centered strategy for writing PR
medication instructions. J Health Commun 2013; 18 Suppl 1: 40-8.

- 44. Houwing S, Hagenzieker M, Mathijssen RP, Legrand SA, Verstraete AG, Hels T, Bernhoft IM, Simonsen KW, Lillsunde P, Favretto D, Ferrara SD, Caplinskiene M, Movig KL, Brookhuis KA. Random and systematic errors in case-control studies calculating the injury risk of driving under the influence of psychoactive substances. Accid Anal Prev 2013; 52: 144-53.
- 45. Houwing S, Legrand SA, Mathijssen R, Hagenzieker M, Verstraete AG, Brookhuis K. Prevalence of psychoactive substances in dutch and belgian traffic. J Stud Alcohol Drugs 2012; 73(6): 951-60.
- 46. Brubacher JR, Chan H, Purssell E, Tuyp BJ, Ting DK, Mehrnoush V. Minor Injury Crashes: Prevalence of Driver-Related Risk Factors and Outcome. J Emerg Med 2017 Mar 7. pii: S0736-4679(17)30113-0.
- Rudisill TM, Zhao S, Abate MA, Coben JH, Zhu M. Trends in drug use among drivers killed in U.S. traffic crashes, 1999-2010. Accid Anal Prev 2014; 70: 178-87.
- 48. Orriols L, Luxcey A, Contrand B, Gadegbeku B, Delorme B, Tricotel A, Moore N, Salmi LR, Lagarde E. Road traffic crash risk associated with benzodiazepine and z-hypnotic use after implementation of a colour-graded pictogram: a responsibility study. Br J Clin Pharmacol 2016; 82(6): 1625-1635.

Table 1. Data on consumption of medicines according to CONCYLIA database and drivers' license census

	Population in Castile & León with health	Drivers' license census (December 2015)	All medicines % (95CI)	Medici "Me	nes with the p dicines and Dr % (95Cl)	Drivers using medicines with the pictogram "Medicines and Driving" %	
	insurance			Acute	Sub-acute	Chronic	(95CI)
	(December						
	2015)						
TOTAL	2 376 717	1 470 389	73.69	5.31	6.64	22.46	25.36
			(73.63–73.75)	(5.28–5.34)	(6.61–6.67)	(22.41–22.51)	(25.29–25.43)
Sex							
Male	1 168 591	887 357	68.78	5.06	5.58	17.69	
Fomalo	1 209 126	E 02 022	(08.70-08.80)	(5.02-5.10)	(5.54-5.62)	(17.02-17.70)	(20.37-20.55)
remale	1 208 120	565 052	(78.36–78.50)	(5.52–5.60)	(7.61–7.71)	(26.98–27.14)	(23.58–23.8)
Age range (Male/Female)							
0-4	45 405 / 42 504	-	76.59 / 74.86	15.61 / 15.31	2.81 / 2.37	0.52 / 0.43	-
5-9	50 925 / 48 078	-	71.22 / 69.60	6.82 / 7.05	1.57 / 1.34	2.73 / 1.50	-
10-14	49 439 / 47 220	-	67.24 / 65.53	3.47 / 3.86	1.48 / 1.49	6.92 / 3.12	-
15-19	48 620 / 46 904	9 282 / 5 586	62.49/68.16	3.17 / 4.18	2.88 / 4.71	7.06 / 5.33	2.50 / 1.69
20-24	54 724 / 53 382	43 294 / 35 387	53.00 / 67.78	3.48 / 4.77	3.81 / 6.75 🗸	4.51 / 5.87	9.34 / 11.53
25-29	62 787 / 61 247	55 831 / 50 618	47.83 / 65.06	3.51 / 5.06	4.15 / 7.36	4.81 / 7.23	11.09 / 16.24
30-34	75 089 / 71 664	69 810 / 61 387	46.72 / 66.41	3.51 / 5.27	4.62 / 7.87	6.02 / 9.44	13.16 / 19.34
35-39	90 372 / 87 031	86 841 / 75 838	50.20 / 68.35	3.95 / 5.20	5.25 / 8.40	7.84 / 12.28	16.38 / 22.56
40-44	92 686 / 89 879	89 294 / 76 277	54.63 / 69.28	4.10 / 5.27	5.72 / 8.89	10.12 / 16.23	19.20 / 25.79
45-49	93 082 / 91 643	90 151 / 74 310	59.23 / 72.04	4.32 / 5.47	5.99 / 9.55	12.90 / 20.68	22.48 / 28.95
50-54	93 252 / 90 618	90 450 / 67 282	66.50 / 77.55	4.87 / 5.83	6.47 / 9.86	16.37 / 25.63	26.88 / 30.67
55-59	87 280 / 84 212	85 820 / 56 346	74.43 / 82.71	5.09 / 5.83	7.04 / 9.60	20.38 / 31.45	31.96 / 31.37

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

BMJ Open

2	
3 4	60
5	65
6	70
7 8	75
9	80
10	85
12	90
$\begin{array}{c} 13\\ 14\\ 15\\ 16\\ 7\\ 8\\ 9\\ 0\\ 1\\ 2\\ 2\\ 3\\ 4\\ 5\\ 6\\ 7\\ 8\\ 9\\ 0\\ 1\\ 2\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 9\\ 0\\ 1\\ 2\\ 3\\ 3\\ 3\\ 3\\ 9\\ 0\\ 1\\ 2\\ 3\\ 3\\ 3\\ 3\\ 9\\ 0\\ 1\\ 2\\ 3\\ 3\\ 3\\ 3\\ 9\\ 0\\ 1\\ 2\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\ 3\\$	

60-64	72 448 / 69 337	71 450 / 36 255	82.67 / 87.96	5.57 / 5.98	7.34 / 9.87	25.65 / 37.38	38.03 / 27.84
65-69	65 430 / 66 777	62 572 / 23 964	89.09 / 91.10	5.88 / 5.90	7.97 / 9.69	31.21 / 43.95	43.09 / 21.37
70-74	56 526 / 61 968	51 161 / 12 390	93.79 / 94.39	5.86 / 5.55	8.01/9.14	38.05 / 52.02	46.98 / 13.34
75-79	45 154 / 56 939	35 993 / 5 000	93.34 / 93.00	5.76 / 4.74	7.89 / 8.21	44.10 / 58.52	46.04 / 6.28
80-84	44 543 / 62 354	28 304 / 1 941	95.81/95.85	5.41/4.21	7.38 / 7.21	51.24 / 64.90	40.69 / 2.38
85-89	27 547 / 46 335	14 160 / 429	99.79 / 98.09	4.98 / 3.66	7.63 / 6.57	56.91 / 69.24	35.74 / 0.74
90 and more	13 282 / 30 034	2 944 / 22	99.90 / 98.51	4.59 / 3.39	8.20 / 6.37	58.92 / 68.08	15.89 / 0.06
Abbreviations: 95CI, confidence interval.							

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

Frequency	Number of DIM	Patients under treatment % (95Cl)			Drivers under treatment % (95Cl)			
of use		Males	Females	Total	Males	Females	Total	
Acute	1	97.82	97.17	97.48	98.09	97.36	97.81	
		(97.71-97.94)	(97.05-97.3)	(97.39-97.56)	(97.95-98.22)	(97.16-97.57)	(97.69-97.93)	
	2	2.16	2.80	2.50	1.89	2.6	2.16	
		(2.04-2.28)	(2.67-2.92)	(2.41-2.58)	(1.75-2.03)	(2.39-2.8)	(2.04-2.28)	
	3 or more	0.02	0.03	0.03	0.02	0.04	0.03	
		(0.01-0.03)	(0.02-0.05)	(0.02-0.03)	(0.01-0.04)	(0.01-0.06)	(0.02-0.04)	
	Mean (±SD)	1.02	1.03	1.03	1.02	1.03	1.02	
		1.02-1.02)	(1.03-1.03)	(1.03-1.03)	(1.02 - 1.02)	(1.03-1.03)	(1.02 - 1.02)	
Sub-acute	1	71.59	67.39	69.13	71.75	68.59	70.41	
		(71.25-71.94)	(67.08-67.69)	(68.90-69.35)	(71.36-72.15)	(68.12-69.06)	(70.1-70.71)	
	2	24.71	27.35	26.26	24.53	26.32	25.29	
		(24.38-25.04)	(27.06-27.63)	(26.04-26.47)	(24.15-24.9)	(25.87-26.77)	(25-25.58)	
	3 or more	3.69	5.27	4.62	3.72	5.09	4.3	
		(3.55-3.84)	(5.12-5.41)	(4.51-4.72)	(3.56-3.89)	(4.86-5.31)	(4.17-4.44)	
	Mean (±SD)	1.32	1.38	1.36	1.32	1.37	1.34	
		(1.32-1.32)	(1.38-1.38)	(1.36-1.36)	(1.32-1.32)	(1.36-1.38)	(1.34-1.34)	
Chronic	1	31.36	23.84	26.76	30.91	25.58	29.07	
		(31.04-31.67)	(23.65-24.05)	(26.58-26.93)	(30.54-31.28)	(25.12-26.03)	(28.78-29.36)	
	2	28.34	26.96	27.49	28.33	28.22	28.29	
		(28.03-28.64)	(26.74-27.17)	(27.31-27.67)	(27.97-28.69)	(27.75-28.68)	(28-28.58)	
	3 or more	40.31	49.20	45.75	40.76	46.21	42.64	
		(39.97-40.64)	(48.95-49.44)	(45.56-45.95)	(40.36-41.15)	(45.69-46.72)	(42.33-42.95)	
	Mean (±SD)	2.63	2.96	2.83	2.65	2.86	2.72	
		(2.62-2.64)	(2.95-2.97)	(2.82-2.84)	(2.64-2.66)	(2.85-2.87)	(2.71-2.73)	
Total	Mean (±SD)	2.08	2.39	2.27	2.1	2.15	2.12	
		(2.07-2.09)	(2.38-2.40)	(2.27-2.27)	(2.09-2.11)	(2.14-2.16)	(2.11-2.13)	

Table 2. Frequency of the consumption of driving-impairing medicines

Abbreviations: 95%CI, 95% confidence interval; DIM, Driving-impairing medicines.

ATC Groups	Pa	atients under treatmei	nt	Drivers under treatment			
		% (95CI)			% (95CI)		
	Males	Females	Total	Males	Females	Total	
Α	5.3 (5.26-5.34)	5.5 (5.46-5.54)	5.4 (5.37-5.43)	5.13 (5.08-5.17)	2.24 (2.21-2.28)	3.98 (3.95-4.02)	
С	0.36 (0.35-0.37)	0.26 (0.25-0.26)	0.31 (0.3-0.31)	0.32 (0.31-0.33)	0.04 (0.04-0.05)	0.21 (0.2-0.22)	
D	0.12 (0.12-0.13)	0.11 (0.1-0.11)	0.11 (0.11-0.12)	0.07 (0.07-0.08)	0.08 (0.07-0.09)	0.08 (0.07-0.08)	
G	0.6 (0.59-0.62)	1 (0.99-1.02)	0.81 (0.79-0.82)	0.52 (0.5-0.53)	0.49 (0.47-0.51)	0.51 (0.49-0.52)	
١	0.02 (0.02-0.03)	0.03 (0.03-0.04)	0.03 (0.03-0.03)	0.03 (0.02-0.03)	0.03 (0.02-0.03)	0.03 (0.02-0.03)	
L	0.51 (0.49-0.52)	0.07 (0.06-0.07)	0.28 (0.28-0.29)	0.37 (0.36-0.38)	0.05 (0.04-0.05)	0.24 (0.23-0.25)	
М	0.7 (0.69-0.72)	1.11 (1.09-1.13)	0.91 (0.9-0.92)	0.78 (0.76-0.8)	1.02 (0.99-1.04)	0.87 (0.86-0.89)	
Ν	22.45 (22.37-22.52)	35.68 (35.6-35.77)	29.17 (29.12-29.23)	21.51 (21.42-21.6)	21.02 (20.91-21.12)	21.31 (21.25-21.38)	
Р	0.08 (0.07-0.08)	0.24 (0.23-0.25)	0.16 (0.15-0.16)	0.08 (0.07-0.09)	0.19 (0.18-0.2)	0.12 (0.12-0.13)	
R	3.62 (3.59-3.65)	4.38 (4.34-4.41)	4 (3.98-4.03)	2.51 (2.48-2.54)	1.98 (1.94-2.02)	2.3 (2.27-2.32)	
S	0.41 (0.4-0.42)	0.42 (0.41-0.43)	0.41 (0.41-0.42)	0.33 (0.32-0.35)	0.1 (0.09-0.1)	0.24 (0.23-0.25)	
V	0 (0-0)	0 (0-0)	0 (0-0)	0 (0-0)	0 (0-0)	0 (0-0)	

utic Chemical Ci, .ar system; D, Dermato, sex hormones and insulins; .usculo-skeletal system; N, Nervous sys, .y organs; V, Various. Abbreviations: ATC, Anatomical Therapeutic Chemical Classification System; A, Alimentary tract and metabolism; B, Blood and blood forming organs; C, Cardiovascular system; D, Dermatologicals; G, Genito-urinary system and sex hormones; H, Systemic hormonal preparations, excluding sex hormones and insulins; J, Antiinfectives for systemic use; L, Antineoplastic and immunomodulating agents; M, Musculo-skeletal system; N, Nervous system; P, Antiparasitic products, insecticides and repellents; R, Respiratory system; S, Sensory organs; V, Various.

ATC Group	Frequency of use	Pa	ntients under treatn % (95CI)	ler treatment Drive % 5CD			ers under treatment % (95Cl)	
		Males	Females	Total	Females	Males	Total	
Α	Acute	0.68 (0.61-0.74)	1.53 (1.44-1.62)	1.12 (1.06-1.18)	0.57 (0.5-0.64)	2.22 (1.96-2.47)	0.94 (0.86-1.02)	
	Sub-acute	17.7 (17.4 - 18)	29.63 (29.28-29.98)	23.87 (23.64-24.11)	17.69 (17.34-18.04)	53.18 (52.32-54.03)	25.62 (25.27-25.97)	
	Chronic	81.62 (81.32-81.93)	68.84 (68.49-69.19)	75.01 (74.77-75.24)	81.74 (81.38-82.09)	44.6 (43.75-45.46)	73.44 (73.09-73.8)	
N	Acute	18.06 (17.92-18.21)	12.79 (12.69-12.89)	14.79 (14.7-14.87)	18.02 (17.85-18.2)	16.55 (16.34-16.75)	17.45 (17.31-17.58)	
	Sub-acute	21.6 (21.44-21.76)	18.8 (18.68-18.91)	19.86 (19.76-19.95)	23.53 (23.34-23.72)	25.72 (25.48-25.97)	24.39 (24.24-24.54)	
	Chronic	60.33 (60.15-60.52)	68.41 (68.27-68.55)	65.36 (65.24-65.47)	58.45 (58.23-58.67)	57.73 (57.45-58.01)	58.17 (57.99-58.34)	
R	Acute	71.93 (71.5-72.36)	74.38 (74.01-74.76)	73.29 (73.01-73.58)	70.21 (69.61-70.81)	78.92 (78.17-79.66)	73.18 (72.71-73.65)	
	Sub-acute	22.63 (22.24-23.03)	21.93 (21.57-22.28)	22.24 (21.98-22.5)	23.86 (23.3-24.42)	18.49 (17.78-19.2)	22.03 (21.58-22.47)	
	Chronic	5.43 (5.22-5.65)	3.69 (3.53-3.85)	4.46 (4.33-4.6)	5.93 (5.62-6.24)	2.59 (2.3-2.88)	4.79 (4.56-5.02)	

Table 4. Frequency of the consumption of driving-impairing medicines A, N and R ATC group.

Abbreviations: ATC, Anatomical Therapeutic Chemical Classification System; A, Alimentary tract and metabolism; N, Nervous system; R, Respiratory system



Figure 1. Frequency of medicine consumption in Castile & León in 2015.

BMJ Open

Use of driving-impairing medicines by the population: a population-based registry study.

Journal:	BMJ Open
Manuscript ID	bmjopen-2017-017618.R1
Article Type:	Research
Date Submitted by the Author:	04-Aug-2017
Complete List of Authors:	Gutierrez-Abejón, Eduardo; Gerencia Regional de Salud de Castilla y leon, Technical Direction of Pharmaceutical Assistance Herrera-Gómez, Francisco; Universidad de Valladolid, Pharmacology and Therapeutics Criado-Espegel, Paloma; Gerencia Regional de Salud de Castilla y León, Technical Direction of Pharmaceutical Assistance Alvarez, F. Javier; University of Valladolid, Pharmacology & Therapeutics
Primary Subject Heading :	Global health
Secondary Subject Heading:	Public health
Keywords:	drug prescription, drug utilization, risk assessment, accident, traffic, automobile driving, driving-impairing medicines



Reviewed version

Use of driving-impairing medicines by the population: A population-based registry study

Eduardo Gutierrez-Abejón¹, Francisco Herrera-Gómez^{2,3}, Paloma Criado-Espegel¹ and F. Javier Álvarez^{2,4}

¹Technical Direction of Pharmaceutical Assistance, Gerencia Regional de Salud de Castilla y León, Valladolid, Spain.

²Pharmacology, Faculty of Medicine, University of Valladolid, Valladolid, Spain.

³Department of Nephrology, Complejo Asistencial de Zamora, Zamora, Spain.

⁴ CEIC/CEIm, Hospital Clínico Universitario de Valladolid, Valladolid, Spain

Correspondence: Professor F. Javier Álvarez, Pharmacology, Faculty of Medicine, University of Valladolid, CEIC/CEIm, Hospital Clínico Universitario de Valladolid, Valladolid, Spain. Tel: +34 983 423077; Fax: +34 983 423022; E-mail: <u>alvarez@med.uva.es</u> ORCID: 0000-0002-7566-5678

Running head: Medicines and driving.

Keywords: Accidents, Traffic. Automobile Driving. Drug Prescriptions. Drug Utilization. Risk Assessment. Driving-Impairing Medicines

Word count (excluding the title page, abstract, references, tables, and figures): 3139

Number of tables: 04

Number of figures: 02

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

ABSTRACT

OBJECTIVE

To assess the use of driving-impairing medicines (DIM) in the general population with special reference to length of use and concomitant use.

DESIGN

Population-based registry study.

SETTING

The year 2015 granted medicines consumption data recorded in the Castile and León (Spain) medicine dispensation registry was consulted.

PARTICIPANTS

Medicines and DIM consumers from a Spanish population (Castile & León: 2.4 million inhabitants).

EXPOSURE

Medicines and DIM consumption. Patterns of use by age and gender based on the length of use (acute: 1-7 days, sub-chronic: 8-29 days, and chronic use: \geq 30 days) were of interest. Estimations regarding the distribution of licensed drivers by age and gender were employed to determine the patterns of use of DIM.

RESULTS:

DIM were consumed by 34.4% (95%CI 34.3-34.5) of the general population in 2015, more commonly with regularity (chronic use: 22.5% versus acute use: 5.3%), and more frequently by the elderly. On average, 2.3 DIM *per* person were dispensed, particularly to chronic users (2.8 DIM *per* person). Age and gender distribution differences were observed between the Castile and León medicine dispensation registry data and the drivers' license census data. Of all DIM

BMJ Open

dispensed, 83.8% were in the Anatomical Therapeutic Chemical code (ATC) group nervous system medicines (N), which were prescribed to 29.2% of the population.

CONCLUSIONS:

The use of DIM was frequent in the general population. Chronic use was common, but acute and sub-acute use should also be considered. This finding highlights the need to make patients, health professionals, health providers, medicine regulatory agencies and policy-makers at large aware of the role DIM play in traffic safety.

in trans .

STRENGTHS AND LIMITATIONS OF THE STUDY

- This study explores the consumption of all driving-impairing medicines and patterns of use by age and gender corresponding to a European population.
- This highlights the need to make patients, health professionals, health providers, medicine regulatory agencies and policy-makers at large aware of the role of DIM in traffic safety.
- The information provided covers all dispensed medicines by the public health system in Spain but does not cover hospital dispensed medications or over-the-counter medicines, a portion of which may be DIM. Furthermore, no information is available on alcohol use or when the medicines were taken in relation to driving.



BACKGROUND

Driving a motor vehicle is a multifaceted task and requires appropriate cognitive and psychomotor skills (e.g., alertness, concentration, reaction time, visual acuity).¹⁻² Medicines can adversely affect these driving-related skills and, consequently, be a hazard to traffic safety.³⁻⁵ There is increasing awareness that implementation of appropriate measures to limit the consumption of alcohol and other substances ("illicit" drugs and medicines) while driving may have an impact on road accident occurrences.⁶ Nevertheless, to date, it is unknown how frequent the consumption of driving-impairing medicines (DIM) in the general population is, or how frequently several of these drugs are consumed concomitantly.⁷

Conversely, most developed countries perform toxicological analyses on road accident casualties and fatalities, and the presence of illicit drugs and medicines (either used legally or illegally) are detected.^{8,9} On-road tests (at random or on target populations) are used ever more frequently worldwide: the on-site screening devices detect some groups of illicit drugs and certain medicines in saliva (oral fluid), with confirmation analyses being performed later.^{8,9} In other countries, blood analyses¹⁰ are performed, rather than screening of saliva. The information from these sources (data on casualties/fatalities and on-road test data) gives only a partial vision of the problem regarding medicines and driving.¹¹

However, medicine regulatory agencies do attempt to provide appropriate information to the public concerning the problem: in the European Union, the summary of product characteristics and the package leaflets contain information on medicines that "affect the ability to drive and to use machines".^{12,13} Furthermore, there have been several attempts to categorize the effects of

medicines on driving¹⁴⁻¹⁶ and several countries, such as France¹⁷ and Spain,¹⁸ have introduced specific mandatory pictograms or ancillary warning labels, as in the Netherlands¹⁹ and Australia.²⁰

The Spanish Law of 2007 (Royal Decree 1345/2007)¹⁸ established the rule that newly authorized medicines that may negatively affect fitness to drive or to handle dangerous machinery should include a warning symbol (pictogram) on the outside of the packaging. Since 2011, all medicines that could possibly affect fitness to drive and are commercially available in Spain have this pictogram on the packaging.²¹ As of January 2016, a total of 2013 medicinal drugs permitted in Spain had been reviewed, of which 402 (20%) included the pictogram on medicines and driving on the packaging.²² This pictogram is well-regarded by the population.²³

We have considered these medicines with the pictogram "medicines and driving" on the packaging in Spain as driving impairing medicines or, to be more exact, potentially impairing medicines on driving. In 2016, a national consensus on Medicines and Driving was reached in Spain to determine the extent of the population taking DIM as a priority and to decipher patterns of use for these drugs.²⁴ This did not apply only to motor vehicle drivers and professional drivers but also the population at large, as well as to all road users, including pedestrians and the ever more common cyclists. Thus, the presence of illicit drugs and medicines was also frequently found in pedestrians involved in fatal road accidents.²⁵

The detection of some medicines in on-road tests has been the object of awareness-raising in public and health professionals, as well as the subject of campaigns to inform the general public, as in the UK.²⁶ In some countries, for instance, the UK,²⁷ Spain²⁸ or Norway,²⁹ on-road positive

BMJ Open

cases to medicines are not fined if they were used according to a physician's prescription. Again, it is necessary that the consumption of DIM and their patterns of use should be known in detail, given that there is a shortage of information about this.

Fitness to drive evaluations have been applied in most developed countries^{30,31}, although the procedures differ markedly. Across the European Union, there is a minimum common regulation under Council Directive 439/EEC³⁰. Within the context of a fitness to drive evaluation, an issue to be considered is medication use (prescribed and over-the-counter) by the driver, although this should always be assessed under the complex relation between disease-medication, particularly among aged people who frequently suffer from several diseases and are poly-medicated.^{1,14-16}

Consequently, the aim of our study was to explore the use of DIM by the general population. The consumption of medicines with the pictogram "medicines and driving" was assessed on the basis of our dispensation registry, focusing on concomitant use of these drugs and on their length of use. In addition, estimations were compared with the drivers' license census to determine the patterns of use among drivers.

METHODS

Study population: CONCYLIA database

Access was provided to the CONCYLIA database to assess the dispensation of granted medicines by the Spanish public health system in Castile and León during 2015.³²

Basically, the CONCYLIA database includes information on all medicine dispensations by the public health system, except those dispensed at hospitals, medical prescriptions dispensed through private medicine clinics, and those that do not require a medical prescription ('over the counter' medications).

We assessed medicine dispensation per person using the patient identification number; that is, for each person, any dispensation during 2015 was identified (e.g., medicinal product, number of doses, and data of dispensation). For data protection, the final database provided by the health system was anonymized, and no personal identification was included.

Target population

The population distribution covered by the public health system and the population distribution according to the population census matched well: Castile and León had a population of 2,428,901 in December 2015^{33} , and 2,376,717 were covered by the public health system at that time (97.85% of the total).³⁴

The current target population of the study was the general population at large. However, not all persons had a motor vehicle license or drove motor vehicles. Due to the lack of information on driving recorded in the CONCYLIA database, weighting was performed to adjust consumption of DIM of the general population to licensed drivers by age and gender based on the Castile and León drivers' license census data up to December 2015.³⁵ Therefore, the results are presented in regard to the general population and/or as estimates of the driver population based on weighting to the drivers' license census data (Table 1).

DIM

As mentioned above, granted medicines in Spain with the pictogram "medicines and driving" were considered as DIM.

In the CONCYLIA database, based on the Anatomical Therapeutic Chemical code (ATC), each one of the medicines is identified as having such a pictogram or not. This information was taken from the Spanish Medicine Agency, updated to February 1, 2016.²²

Variables and ethical issues

The following variables describing the consumption of medicines and DIM in Castile and León in the year 2015 were considered:

i) Yearly frequency of all medicine use;

ii) Yearly frequency of DIM use: acute (1-7 days), sub-acute (8-29 days) and chronic or regular

use (\geq 30 days);

iii) Yearly frequency of daily use of at least one DIM;

iv) Number and means of different DIM taken within 2015.

All analyses were made considering age and gender distributions.

Ethics Review Board approval was obtained (Reference number PI 16-387, approved on March 17th, 2016)

Statistical analysis

All values are given as percentages (frequencies) with a 95% confidence interval (95%CI) or as the mean \pm standard deviations (SD). For comparisons, Student's *t* test was used for continuous variables and Pearson's chi-squared test for categorical variables. A two-tailed P < 0.05 was considered to be significant. All statistical analyses were performed using the Statistical Package for Social Sciences (SPSS version 23.0.; SPSS Inc, Chicago, IL).

RESULTS

Descriptive mapping

A total of 48,858,588 medicines were dispensed in 2015. Approximately 3 of 4 people took a medicinal product in 2015, with more females than males taking the product (78.4% versus 68.8%, p < 0.05), and this fraction increased with age (Table 1).

One of three (34.4%, 95%CI 34.3-34.5) consumed DIM in 2015, again more frequently among females (40.3% versus 28.3%, p < 0.05), and this also increased with age. A majority needed to use these medicines on a regular basis (chronic use: 22.5%), while the use for a few days or weeks accounted for 5.3% and 6.6%, respectively, with similar patterns of use by age and gender (Table 1).

However, if the distribution is performed with respect to the drivers' license census, 25.4% (95%CI 25.3-25.43) of people took DIM in 2015, with more males than females (26.5% versus

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

BMJ Open

23.7%, *p* < 0.05) and mostly took DIM regularly (chronic use: 15.3%, 95%CI 15.2-15.32; sub-acute use: 5.96%, 95%CI 5.92-5.99; acute use: 4.14%, 95%CI 4.11-4.18).

Figure 1 and Table 1 show those who used DIM in 2015, their distribution by age and gender, and relation to the drivers' license census. Age trends differed between the sexes with consumption dropping dramatically among female drivers from 60 years of age and male drivers using less DIM over 75 years of age.

At least one DIM was consumed daily by 5.6% (95%CI 5.52-5.58) of people, and by 3.7% (95%CI 3.67-3.73) of licensed drivers.

On average (Table 2), each person taking DIM took 2.3 medicines (2.1 according to the drivers' license census data). Acute and sub-acute consumers (97.5% and 69.1%, respectively) took only one DIM, while chronic consumers (71.5%) took 2 DIM or more (mean number of DIM use: 2.8). Trends between sexes were similar when the drivers' license census data were analysed (Table 2).

Types of DIM consumed

Of the 10,862,138 DIM dispensed, 9,102,052 (83.8%) belonged to the ATC classification group N (Nervous System), 1,176,864 (10.8%) to group A (Alimentary Tract and Metabolism) and 160,631 (1.5%) to group R (Respiratory System). ATC group N medicines were prescribed to 29.2% of the population (21.3% regarding the drivers' license census), group A medicines to 5.4% (4% for drivers), and group R medicines to 4% (2.3% for drivers) (Table 3).

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

Interestingly, ATC groups N, A and R were more frequently prescribed to females than males, all people considered. When considering licensed drivers, the trends showed no differences. Table 4 shows DIM used for several days or weeks and chronically.

DISCUSSION

A detailed description of the consumption of DIM in the general population from Castile and León in 2015 is provided. DIM were consumed by 34.4% (95%CI 34.3–34.5) of the general population and more commonly on a regular basis (22.5%). However, the use for several days (5.3%) or a few weeks (6.6%) should not be neglected. The consumption of DIM increased in line with age. Acute and sub-acute consumers took at least one DIM and chronic users took nearly three. Of all DIM dispensed, 83.8% belong to the ATC classification group N (Nervous System), which were dispensed to 29.2% of the population. Similar trends were found regarding the distribution of licensed drivers by gender but not by age.

The DRUID project¹ provides information on the prevalence of use in Europe of some types of medicines randomly detected in drivers³⁶. Of all positive matches (1.36%), benzodiazepines (0.9%), Z-drugs (0.12%), and opioids (0.35%) were frequently confirmed. Furthermore, there is information for other developed countries on the consumption of alcohol, illicit drugs and certain medicines by people injured/killed in road traffic accidents.⁸⁻¹¹ The DRUID study conducted on injured (seriously injured or killed) people in nine European countries did not produce a clear picture of the use of medicines (and illicit drugs), but combined use of alcohol with medicines (and/or illicit drugs) was shown to be much more common in drivers who had accidents than in

Page 13 of 37

BMJ Open

the driving population.³⁶ Although progress has been made in understanding this social and medical problem of driving under the effects of medicines, the available data enable only a partial view of the problem, as only several groups of DIM (mainly psychotropic drugs) have been analysed in blood and oral fluid specimens from drivers. There has also been an attempt to estimate DIM consumption based on dispensed medicines,⁷ or using driver consumption surveys.³⁷ Our study provides a detailed overview of all DIM used by the general population, and to the best of our knowledge, this study is the first on this matter.

The combined use of DIM with alcohol is well-known to have marked effects on psychomotor performance.^{1,5,14-16} Furthermore, the risk of being seriously injured or killed while driving with these psychoactive substances was highly increased with multiple use and the risk increased severely with combined use with alcohol.^{1,36} Avoiding use of alcohol is a priority for safe driving,^{1,6} but particularly for those who take medicines, either acutely or regularly.

One of 3 used DIM in 2015. Importantly, acute users represented a sizeable proportion of all drivers consuming DIM (5.3%). The effect of medications on driving is more relevant in the first days of use.^{38,39} Drivers consuming DIM for few days might therefore be the most affected, particularly those taking more than one medicine, and this must be taken into account. In addition, multiple daily dosing is an important factor to consider,^{7,40} especially for drivers over 50 years old.⁴⁰

More than 2 DIM were dispensed (drivers and non-drivers), particularly to chronic users who took nearly three. In addition, approximately 6% of people consumed at least one DIM daily during the year 2015. Impairment of driving seems to diminish with chronic/stable DIM use,³⁹

probably due to tolerance.⁴¹ However, clinical explorations of fitness to drive under the effects of DIM should be performed.⁴¹ Tolerance is a problem that has not been completely assessed and what occurs when more than one medicine is consumed has also not been analysed. A higher prevalence of regular and daily use of DIM are not uncommon in Spain and other developed countries. Therefore our results provide an epidemiological view of the current impact of medicine use patterns that highlight the importance of daily regimens, as well as the importance for elderly acute users.

Our results showed that several types of medicines are prescribed more often than others. ATC group N medicines were prescribed with predilection, mostly to females. This finding was corroborated by the study of Ravera *et al.*⁷ The finding of frequent DIM use was not surprising, as 20% of the granted medicines (402 of 2,013) in Spain are DIM (with the pictogram "medicines and driving" on the packaging). In addition, 83.8% of dispensed DIM in Castile and León were ATC group N medicines (178 of 402). In this context, mandatory pictograms and warning labels contribute to awareness of DIM consumption risks for consumer engagement,^{42,43} with the noticeability of these medication warnings a challenging task.⁴⁴ Furthermore, there are initiatives worldwide for refining information on the risk categorization of drugs^{1,14-16,45,46} that must be implemented in dispensing support tools (software)⁴⁷ for a better prescription/dispensation of DIM.

Our study showed that DIM use by the population is frequent, even in young people/children, who are not motorized vehicle drivers: however, all of us are road users (pedestrians). Medicinal products authorized for use in children do not have the pictogram for medicines and driving in Spain; however, medicines that could be used by the population, including young people, include

BMJ Open

it. Although the topic of medicines and driving has focused on motorized vehicles, their use by cyclists and pedestrians²⁵ is a field of growing interest, especially involving road accidents.

Our study was based in a region of Spain. Current information from the CONCILYA medicines dispensation registry shows that medication use in Castile and León does not differ from other areas in Spain (as measured in Defined Daily Doses [DDD] *per* 1000 inhabitants-day)^{48,49} and are in line with those reported in other countries.⁷ Recently, Eurostat reported on medicine use in the European Union.⁵⁰ In the European health interview survey, conducted between 2013 and 2015, people were asked about self-reported medicine use. Our data by gender and age range agree well with these results, although figures from the Eurostat refer to medicine use in the two weeks prior to the survey, and the current data were based on any medicine dispensed in 2015. Therefore, although considered with caution due to possible country variations, the figures from the present study could be generalized to other developed countries.

Our study provides detailed information on which DIM are consumed and how. We thus fulfilled the objectives of the Spanish consensus on Medicines and Driving reached recently. Giving clearer information about the influence of medicines on driving to sensitize health professionals and the general population on the negative effects of DIM is a priority.²⁴ Our results stress the need to improve the communication of DIM risks, in line with recent requirements. Nevertheless, DIM risk communication is a complex clinical, methodological and epidemiological challenge, and the "boosters" (warning label methods, dispensation software, information campaigns, etc.) should be cautiously implemented in key steps, with the main objective to minimize road accidents. Again, detailed knowledge of the use of DIM is a priority.

This study has several limitations. The health system in Spain is public and free, and we used the data from a medicine dispensation registry, which implies that the information covers all dispensed medicines within such a system but not hospital dispensed medications or over-thecounter medicines, several of which may not have the Spanish pictogram. Data are presented regarding the general population, not only drivers, because even pedestrians and cyclists could be involved in road traffic accidents with DIM being a possible cause.²⁵ In the CONCYLIA database, no information on medicine use by drivers is recorded, and weighting was performed to adjust the consumption of DIM among licensed drivers by age and gender based on the Castile & León drivers' license census data. This should be taken into account, as the distribution of drivers in other countries or regions could be different, and especially because information is not available on the extent to which drivers with a license drove vehicles. Furthermore, we do not have information on patterns of alcohol use or on driving patterns. Importantly, the effect of drugs on driver behaviour (and crash risk) depends on when the drug was taken in relation to driving. Our study showed that a high percentage of drivers are taking DIM and are frequently taking several DIM. However, we do not have information about when the drivers took the medications; for example, they may have taken them at a time when their driving was unlikely to be impaired (i.e., before bed).

CONCLUSIONS

The use of DIM was frequent in the general population based on the findings of Castile and León in 2015. Chronic use (30 days or more) was common, but acute use (1-7 days) and sub-acute use (8-29 days) must not be overlooked because they might be the most relevant regarding DIM consumption and risks. ATC group N medicines were the most frequently prescribed.

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

BMJ Open

There is a need worldwide to improve interventions in the field of medicines and driving.^{1,6} Interventions have been suggested for such populations as the general public (information, awareness of risks for DIM use on driving),^{1,45} for health professionals (e.g., risk communications to the patients, categorization systems, fitness to drive evaluation),^{15,16} for health (prescribing and dispensation software tools),^{16,47} systems for health provider authorities/medicinal regulatory agencies for improving medicinal product labelling systems and rmation leance, inserted patient information leaflets,^{23,42-44} and for road safety policy-makers.^{1,6,45}

Contributors

F.J.A. conceived the study design. E.G.-A. conducted the study. E.G.-A., F.H.-G., P.C.-E. and F.J.A. analysed the data, contributed to the interpretation of the results, and wrote the manuscript. All authors reviewed and approved the final version of the manuscript.

Funding

t This work was supported by the Dirección General de Tráfico (DGT) reference: SP IP2017-02120.

Competing Interests

No competing interests.

Data sharing statement

No additional data are available

BMJ Open

REFERENCES

- Schulze H, Schumacher M, Urmeew R, Auerbach K, Alvarez FJ, Bernhoft IM, de Gier JJ, Hagenzieker M, Houwing S, Knoche A, Pilgerstorfer M, Zlender B. Driving under the influence of drugs, alcohol and medicines in europe — findings from the DRUID project. Lisbon, Portugal: EMCDDA. 2012. <u>http://www.emcdda.europa.eu/publications/thematicpapers/druid</u> (Accessed 9 January 2017)
- Ramaekers JG. Drugs and Driving Research in Medicinal Drug Development. Trends Pharmacol Sci 2017; 38: 319-21.
- Ravera S, van Rein N, de Gier JJ, de Jong-van den Berg LT. Road traffic accidents and psychotropic medication use in The Netherlands: a case-control study. Br J Clin Pharmacol 2011; 72: 505-13.
- Orriols L, Delorme B, Gadegbeku B, Tricotel A, Contrand B, Laumon B, Salmi LR, Lagarde E; CESIR research group. Prescription medicines and the risk of road traffic crashes: a French registry-based study. PLoS Med 2010; 16: e1000366.
- Berghaus G, Sticht G, Grellner W, Lenz D, Naumann TH, WiesenmüllerS. Meta-analysis of empirical studies concerning the effects of medicines and illegal drugs including pharmacokinetics on safe driving. DRUID project Deliverable 1.1.2b. Cologne, Germany: BAST. 2010. <u>http://www.druid-project.eu/Druid/EN/deliverales-</u> <u>list/downloads/Deliverable_1_1_2_B.pdf?__blob=publicationFile&v=1</u> (Accessed 17 July 2017).
- WHO. Drug use and road safety: a policy brief. Geneva, Switzerland: World Health Organization. 2016. <u>http://www.who.int/iris/handle/10665/249533</u> (Accessed 9 January 2017).

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

- 7. Ravera S, Hummel SA, Stolk P, Heerdink RE, de Jong-van den Berg LT, de Gier JJ. The use of driving impairing medicines: a European survey. Eur J Clin Pharmacol 2009: 65: 1139-47. 8. Gómez-Talegón T, Fierro I, González-Luque JC, Colás M, López-Rivadulla M, Javier Álvarez F. Prevalence of psychoactive substances, alcohol, illicit drugs, and medicines, in Spanish drivers: a roadside study. Forensic Sci Int 2012; 223: 106-13. 9. Simonsen KW, Steentoft A, Hels T, Bernhoft IM, Rasmussen BS, Linnet K. Presence of psychoactive substances in oral fluid from randomly selected drivers in Denmark. Forensic Sci Int 2012; 221: 33-8. 10. Bezemer KD, Smink BE, van Maanen R, Verschraagen M, de Gier JJ. Prevalence of medicinal drugs in suspected impaired drivers and a comparison with the use in the general Dutch population. Forensic Sci Int 2014; 241: 203-11. 11. Rudisill TM, Zhao S, Abate MA, Coben JH, Zhu M. Trends in drug use among drivers killed in U.S. traffic crashes, 1999-2010. Accid Anal Prev 2014; 70: 178-87. 12. Council Directive 83/570/EEC of 26 October 1983 amending Directives 65/65/EEC, 75/318/EEC and 75/319/EEC on the approximation of provisions laid down by law, regulation or administrative action relating to proprietary medicinal products. OJ L 332, 28.11.1983, p. 1–10. http://data.europa.eu/eli/dir/1983/570/oj (Accessed 9 January 2017) 13. European Commission. Enterprise and Industry Directorate-GeneralA guideline on summary of product characteristics (SmPC) 2009. http://ec.europa.eu/health/files/eudralex/vol-2/c/smpc guideline rev2 en.pdf and http://ec.europa.eu/health/documents/eudralex/vol-2/index en.htm (Accessed 9 January 2017) 14. de Gier JJ, Alvarez FJ, Mercier-Guyon C, Verstraete AG. Prescribing and dispensing
 - guidelines for medicinal drugs affecting driving performance. In: Drugs, Driving and Traffic

BMJ Open

2
3
4
5
6
7
/ 0
0
9
10
11
12
13
14
15
16
17
17
10
19
20
21
22
23
24
25
26
27
20
20
29
30
31
32
33
34
35
36
37
38
20
39
40
41
42
43
44
45
46
47
48
49
50
51
50
52 50
53
54
55
56
57
58
59
60

Safety, eds Verster JC, Pandi-Perumal SR, Ramaekers JG, de Gier JJ. Basel, Switzerland: Birkaeuser Verlag AG. 2009: 121-34.

- 15. Gómez-Talegón T, Fierro I, Del Río MC, Álvarez FJ, Ravera S, Monteiro S, de Gier JJ, Van der Linden G, Legrand SA, Pil K, Verstraete A, Mallaret M, Mercier-Guyon C, Mercier-Guyon I, Touliou K, Heiβing M. Establishment of framework for classification/categorisation and labelling of medicinal drugs and driving. DRUID project Deliverable 4.3.1. Cologne, Germany: BAST. 2011. http://www.druid-project.eu/Druid/EN/deliveraleslist/downloads/Deliverable_4_3.pdf?__blob=publicationFile&v=1 (Accessed 17 July 2017)
- 16. Ravera S, Monteiro S, de Gier J J, van der Linden T, Gómez-Talegón T, Álvarez FJ. & the DRUID Project WP4 Partner. A European approach to categorizing medicines for fitness to drive: Outcomes of the DRUID project. Br J Clin Pharmacol 2012; 74: 920-31.
- 17. Ministère de la Santé et des Solidarités. Direction Generale de la Santé. Arrêté du 18 Juillet 2005 pris pour l'application de l'article R.5121-139 du code de la santé publique et relative à l'opposition d'un pictogramme sur le conditionnement extérieur de certain médicaments et produits. Journal Officiel de la République Française. Août 2005 (SAN/P0522726A).
- 18. Real Decreto 1345/2007, de 11 de octubre, por el que se regula el procedimiento de autorización, registro y condiciones de dispensación de los medicamentos de uso humano fabricados industrialmente. pp. 45652–45698. BOE de 7 de Noviembre de 2007. http://www.boe.es/diario_boe/txt.php?id=BOE-A-2007-19249 (Accessed 9 January 2017).
- 19. Patrício Monteiro S. Driving-impairing Medicines and Traffic Safety: Patient's Perspectives.
 PhD thesis. Groningen, The Netherlands; University of Groningen. 2014.
 <u>http://www.rug.nl/research/portal/files/6563594/volledigedissertatie_1_.pdf</u> (Accessed 9 January 2017).

20. Jomaa I, Odisho M, Cheung JM, Wong K, Ellis JG, Smyth T, et al. Pharmacists' perceptions

and communication of risk for alertness impairing medications. Res Social Adm Pharm.

https://doi.org/10.1016/j.sapharm.2016.12.010
21. Agencia Española de Medicamentos y Productos Sanitarios. Medicamentos y Conduccion.
http://www.aemps.gob.es/industria/etiquetado/conduccion/home.htm (Accessed 9 July 2016)
22. Agencia Española de Medicamentos y Productos Sanitarios. Medicamentos y Conduccion:
Listados de principios activos por grupos ATC* e incorporación del pictograma de la
conducción
http://www.aemps.gob.es/industria/etiquetado/conduccion/listadosPrincipios/home.htm
(Accessed 9 July 2016)
23. Fierro I, Gómez-Talegón T, Alvarez FJ. The Spanish pictogram on medicines and driving:
The population's comprehension of and attitudes towards its use on medication packaging.
Accid Anal Prev 2013; 50: 1056-61.
24. Documento de consenso sobre medicamentos y conducción en España: información a la
población general y papel de los profesionales sanitarios. Madrid, Spain: Ministerio de
Sanidad, Servicios Sociales e Igualdad and Ministerio del Interior. 2016.
http://www.msssi.gob.es/profesionales/saludPublica/prevPromocion/docs/Medicamentos_con
duccion_DocConsenso.pdf (Accessed 10 May 2016)
25. Instituto Nacional de Toxicología y Ciencias Forenses. Víctimas mortales en accidentes de
tráfico: memoria 2016. Madrid, Spain: Ministerio de Justicia. 2016.
https://administraciondejusticia.gob.es/paj/PA_WebApp_SGNTJ_NPAJ/descarga/Memoria%
20Trafico%20INTCF202014.pdf?idFile=00359cf9-26d5-4d33-96ea-ec9703c78470
(Accessed 9 January 2017).

2017, in press.

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml

BMJ Open

26	. GOV:UK. Drugs and driving: the law. Collection Drug driving. <u>https://www.gov.uk/drug-</u>
	driving-law (Accessed 9 January 2017). https://www.gov.uk/government/collections/drug-
	driving#table-of-drugs-and-limits (Accessed 9 January 2017).
27	. Department for Transport. Guidance for healthcare professionals on drug driving. London,
	United Kingdom: Department for Transport. 2014.
	https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/325275/health
	care-profs-drug-driving.pdf (Accessed 9 January 2017).
28	. Álvarez FJ, González-Luque JC, Seguí-Gómez M. Drugs, Substance Use Disorder and
	Driving: Intervention of Health Professionals in the Treatment of Addictions. Adicciones
	2015; 27: 161-7.
29	. Norwegian Ministry of Transport and Communications. Driving under the influence of non-
	alcohol drugs-legal limits implemented in Norway. Oslo, Norway: Norwegian Government
	Security and Service Organization. 2014.
	https://www.regjeringen.no/globalassets/upload/sd/vedlegg/brosjyrer/sd_ruspavirket_kjoring_
	net.pdf (Accessed 9 January 2017)
30	. Council Directive 91/439/EEC of 29 July 1991 on driving licences. Official Journal, 1991; L
	237: 0001-0024. http://eur-lex.europa.eu/legal-content/ES/TXT/?uri=CELEX:31991L0439
	(Accessed 17 July2017).
31	. Austroads. Assessing Fitness to drive for commercial and private vehicle drivers. Fifth
	Edition. Sydney, Australia: Austroads Ltd., 2016.
	https://www.onlinepublications.austroads.com.au/downloads/AP-G56-16 (Accessed 17
	July2017).

- 32. CONCYLIA. Sistema de Información de Farmacia. Gerencia Regional de Salud de Castilla y León. Valladolid, Spain: Junta de Castilla y León.
 33. Instituto Nacional de Estadística. Cifras oficiales de población de los municipios españoles: Revisión del Padrón Municipal. Madrid, Spain. <u>http://www.ine.es/dyngs/INEbase/es/operacion.htm?c=Estadistica_C&cid=1254736177011&menu=resultados&secc=1254736195458&idp=1254734710990</u> (Accessed 15 Jun 2016)
 34. Tarjeta Sanitaria de SACYL. Recursos y Gestión Poblacional. Gerencia Regional de Salud. 2015. Valladolid, Spain: Junta de Castilla y León.
 35. Ministerio del Interior. Dirección General de Tráfico. Estadísticas e Indicadores. Permisos de conducción. <u>http://www.dgt.es/es/seguridad-vial/estadisticas-e-indicadores/permisos-</u> conducción/ (Accessed 15 June 2016).
 - 36. Bernhoft IM (coordinator). Results from epidemiological research prevalence, risk and characteristics of impaired drivers. DRUID project Deliverable 2.4.1. Cologne, Germany: BAST. 2011. <u>http://www.druid-project.eu/Druid/EN/deliverales-</u>list/downloads/Deliverable 2.4.1.pdf? blob=publicationFile&v=1 (Accessed 20 July 2017)
 - 37. Del Río MC, Alvarez FJ. Medication and fitness to drive. Pharmacoepidemiol Drug Saf 2003;12: 389-94.
 - 38. Barbone F, McMahon AD, Davey PG, Morris AD, Reid IC, McDevitt DG, MacDonald TM. Association of road-traffic accidents with benzodiazepine use. Lancet 1998; 352: 1331-6.
 - Wilhelmi BG, Cohen SP. A framework for "driving under the influence of drugs" policy for the opioid using driver. Pain Physician 2012; 115(3 Suppl): ES215-30.
 - Monárrez-Espino J, Laflamme L, Elling B, Möller J. Number of medications and road traffic crashes in senior Swedish drivers: a population-based matched case-control study. Inj Prev 2014; 20: 81-7.

BMJ Open

2	
3	
4	
5	
6	
7	
8	
0	
3	
10	
11	
12	
13	
14	
15	
16	
17	
17	
18	
19	
20	
21	
22	
23	
24	
27	
20	
26	
27	
28	
29	
30	
31	
32	
22	
33	
34	
35	
36	
37	
38	
39	
40	
11	
41	
42	
43	
44	
45	
46	
47	
48	
49	
50	
50	
51	
52	
53	
54	
55	
56	
57	
52	
50	
09	
$\mathbf{n}(\mathbf{l})$	

41. Schumacher MB, Jongen S, Knoche A, Petzke F, Vuurman EF, Vollrath M, Ramaekers JG. Effect of chronic opioid therapy on actual driving performance in non-cancer pain patients. Psychopharmacology (Berl) 2017; 234: 989-99.

- 42. Monteiro SP, Huiskes R, Van Dijk L, Van Weert JC, De Gier JJ. How effective are pictograms in communicating risk about driving-impairing medicines? Traffic Inj Prev 2013; 14: 299-308.
- 43. Emich B, van Dijk L, Monteiro SP, de Gier JJ. A study comparing the effectiveness of three warning labels on the package of driving-impairing medicines. Int J Clin Pharm 2014; 366: 1152-9.
- 44. Smyth T, Sheehan M, Siskind V, Mercier-Guyon C, Mallaret M. Consumer perceptions of medication warnings about driving: a comparison of French and Australian labels. Traffic Inj Prev 2013; 14: 557-64.
- 45. Schulze H, Schumacher M, Urmeew R, Auerbach K. DRUID (Driving Under the Influence of Drugs, Alcohol and Medicines) project. Final Report: Work performed, main results and recommendations. DRUID project Deliverable 0.1.8. Revision 2.0. Cologne, Germany: BAST. 2010. <u>http://www.druid-</u>

project.eu/Druid/EN/Dissemination/downloads_and_links/Final_Report.pdf?__blob=publicati onFile&v=1 (Accessed 9 January 2017).

46. International Council on Alcohol, Drugs and Traffic Safety (ICADTS): Working Group on Prescribing and Dispensing Guidelines for Medicinal Drugs Affecting Driving Performance. Utrecht, The Netherland: ICADTS. 2001. Available at <u>http://www.icadts.nl/medicinal.html</u> (Accessed 9 July 2016).

- 47. Legrand SA, Boets S, Meesmann U, Verstraete AG. Medicines and driving: evaluation of training and software support for patient counselling by pharmacists. Int J Clin Pharm 2012; 34: 633-43.
- 48. Agencia Española de Medicamentos y Productos Sanitarios. Utilización de medicamentos ansiolíticos e hipnóticos en España durante el periodo 2000-2012. Informe de Utilización de Medicamentos U/HAY/V1/17012014. 2014

https://www.aemps.gob.es/medicamentosUsoHumano/observatorio/docs/ansioliticos_hipnoti cos-2000-2012.pdf (Accessed 20 July 2017)

49. Agencia Española de Medicamentos y Productos Sanitarios. Utilización de medicamentos antidepresivos en España durante el periodo 2000-2013. Informe de Utilización de Medicamentos U/AD/V1/14012015. 2015.

https://www.aemps.gob.es/medicamentosUsoHumano/observatorio/docs/antidepresivos-2000-2013.pdf (Accessed 20 July 2017)

50. EUROSAT statistics explained. Medicine use statistics. <u>http://ec.europa.eu/eurostat/statistics-explained/index.php/Medicine_use_statistics#Prescribed_medicines</u> (Accessed 20 July 2017) <u>http://ec.europa.eu/eurostat/statistics-explained/index.php/File:Self-reported_use_of_prescribed_medicines_by_age_and_sex, 2014 (%25).png</u>

(Accessed 20 July 2017)

Page 27 of 37

BMJ Open

	Population in	Drivers'	All medicines	Medici	nes with the p	ctogram	Drivers using medicines
	Castile & León with	license census	%	"Medicines and Driving"			with the pictogram
	health insurance	(December	(95%CI)		%		"Medicines and Driving"
	card (December	2015)			(95%CI)	-	%
	2015)			Acute	Sub-acute	Chronic	(95%CI)
TOTAL	2 376 717	1 470 389	73.69	5.31	6.64	22.46	25.36
			(73.63–73.75)	(5.28–5.34)	(6.61–6.67)	(22.41–22.51)	(25.29–25.43)
Sex							
Male	1 168 591	887 357	68.78	5.06	5.58	17.69	26.46
			(68.70–68.86)	(5.02–5.10)	(5.54–5.62)	(17.62–17.76)	(26.37–26.55)
Female	1 208 126	583 032 🥒	78.43	5.56	7.66	27.06	23-69
			(78.36–78.50)	(5.52–5.60)	(7.61–7.71)	(26.98–27.14)	(23.58–23.8)
Age range							
(Male/Female)							
0-4	45 405 / 42 504	-	76.59 / 74.86	15.61 /	2.81 / 2.37	0.52 / 0.43	-
				15.31			
5-9	50 925 / 48 078	-	71.22 / 69.60	6.82 / 7.05	1.57 / 1.34	2.73 / 1.50	-
10-14	49 439 / 47 220	-	67.24 / 65.53	3.47 / 3.86	1.48 / 1.49	6.92 / 3.12	-
15-19	48 620 / 46 904	9 282 / 5 586	62.49 /68.16	3.17 / 4.18	2.88 / 4.71	7.06 / 5.33	2.50 / 1.69
20-24	54 724 / 53 382	43 294 / 35 387	53.00 / 67.78	3.48 / 4.77	3.81 / 6.75	4.51 / 5.87	9.34 / 11.53
25-29	62 787 / 61 247	55 831 / 50 618	47.83 / 65.06	3.51 / 5.06	4.15 / 7.36	4.81 / 7.23	11.09 / 16.24
30-34	75 089 / 71 664	69 810 / 61 387	46.72 / 66.41	3.51 / 5.27	4.62 / 7.87	6.02 / 9.44	13.16 / 19.34
35-39	90 372 / 87 031	86 841 / 75 838	50.20 / 68.35	3.95 / 5.20	5.25 / 8.40	7.84 / 12.28	16.38 / 22.56
40-44	92 686 / 89 879	89 294 / 76 277	54.63 / 69.28	4.10 / 5.27	5.72 / 8.89	10.12 / 16.23	19.20 / 25.79
45-49	93 082 / 91 643	90 151 / 74 310	59.23 / 72.04	4.32 / 5.47	5.99 / 9.55	12.90 / 20.68	22.48 / 28.95
50-54	93 252 / 90 618	90 450 / 67 282	66.50 / 77.55	4.87 / 5.83	6.47 / 9.86	16.37 / 25.63	26.88 / 30.67
55-59	87 280 / 84 212	85 820 / 56 346	74.43 / 82.71	5.09 / 5.83	7.04 / 9.60	20.38 / 31.45	31.96 / 31.37
60-64	72 448 / 69 337	71 450 / 36 255	82.67 / 87.96	5.57 / 5.98	7.34 / 9.87	25.65 / 37.38	38.03 / 27.84
65-69	65 430 / 66 777	62 572 / 23 964	89.09 / 91.10	5.88 / 5.90	7.97 / 9.69	31.21 / 43.95	43.09 / 21.37

70-74	56 526 / 61 968	51 161 / 12 390	93.79 / 94.39	5.86 / 5.55	8.01/9.14	38.05 / 52.02	46.98 / 13.34
75-79	45 154 / 56 939	35 993 / 5 000	93.34 / 93.00	5.76 / 4.74	7.89 / 8.21	44.10 / 58.52	46.04 / 6.28
80-84	44 543 / 62 354	28 304 / 1 941	95.81 / 95.85	5.41/4.21	7.38 / 7.21	51.24 / 64.90	40.69 / 2.38
85-89	27 547 / 46 335	14 160 / 429	99.79 / 98.09	4.98 / 3.66	7.63 / 6.57	56.91 / 69.24	35.74 / 0.74
90 and more	13 282 / 30 034 📐	2 944 / 22	99.90 / 98.51	4.59 / 3.39	8.20 / 6.37	58.92 / 68.08	15.89 / 0.06

Abbreviations: 95%CI, confidence interval.

Page 29 of 37

BMJ Open

Frequency	Number of DIM	Patie	ents under treatr % (95%Cl)	nent	Drivers under treatment % (95%Cl)			
of use		Males	Females	Total	Males	Females	Total	
Acute	1	97.82 (97.71-97.94)	97.17 (97.05-97.3)	97.48 (97.39-97.56)	98.09 (97.95-98.22)	97.36 (97.16-97.57)	97.81 (97.69-97.93)	
	2	2.16 (2.04-2.28)	2.80 (2.67-2.92)	2.50 (2.41-2.58)	1.89 (1.75-2.03)	2.6 (2.39-2.8)	2.16 (2.04-2.28)	
	3 or more	0.02 (0.01-0.03)	0.03 (0.02-0.05)	0.03 (0.02-0.03)	0.02 (0.01-0.04)	0.04 (0.01-0.06)	0.03 (0.02-0.04)	
	Mean (±SD)	1.02 1.02-1.02)	1.03 (1.03-1.03)	1.03 (1.03-1.03)	1.02 (1.02-1.02)	1.03 (1.03-1.03)	1.02 (1.02-1.02)	
Sub-acute	1	71.59 (71.25-71.94)	67.39 (67.08-67.69)	69.13 (68.90-69.35)	71.75 (71.36-72.15)	68.59 (68.12-69.06)	70.41 (70.1-70.71)	
	2	24.71 (24.38-25.04)	27.35 (27.06-27.63)	26.26 (26.04-26.47)	24.53 (24.15-24.9)	26.32 (25.87-26.77)	25.29 (25-25.58)	
	3 or more	3.69 (3.55-3.84)	5.27 (5.12-5.41)	4.62 (4.51-4.72)	3.72 (3.56-3.89)	5.09 (4.86-5.31)	4.3 (4.17-4.44)	
	Mean (±SD)	1.32 (1.32-1.32)	1.38 (1.38-1.38)	1.36 (1.36-1.36)	1.32 (1.32-1.32)	1.37 (1.36-1.38)	1.34 (1.34-1.34)	
Chronic	1	31.36 (31.04-31.67)	23.84 (23.65-24.05)	26.76 (26.58-26.93)	30.91 (30.54-31.28)	25.58 (25.12-26.03)	29.07 (28.78-29.36)	
	2	28.34 (28.03-28.64)	26.96 (26.74-27.17)	27.49 (27.31-27.67)	28.33 (27.97-28.69)	28.22 (27.75-28.68)	28.29 (28-28.58)	
	3 or more	40.31 (39.97-40.64)	49.20 (48.95-49.44)	45.75 (45.56-45.95)	40.76 (40.36-41.15)	46.21 (45.69-46.72)	42.64 (42.33-42.95)	
	Mean (±SD)	2.63 (2.62-2.64)	2.96 (2.95-2.97)	2.83 (2.82-2.84)	2.65 (2.64-2.66)	2.86 (2.85-2.87)	2.72 (2.71-2.73)	
Total	Mean (±SD)	2.08 (2.07-2.09)	2.39 (2.38-2.40)	2.27 (2.27-2.27)	2.1 (2.09-2.11)	2.15 (2.14-2.16)	2.12 (2.11-2.13)	

Table 2. Frequency of the consumption of driving-impairing medicines

Abbreviations: 95%CI, 95% confidence interval; DIM, Driving-impairing medicines.

ATC Groups	Pa	atients under treatmei	nt	Drivers under treatment				
		% (95%CI)			% (95%CI)			
	Males	Females	Total	Males	Females	Total		
А	5.3 (5.26-5.34)	5.5 (5.46-5.54)	5.4 (5.37-5.43)	5.13 (5.08-5.17)	2.24 (2.21-2.28)	3.98 (3.95-4.02)		
С	0.36 (0.35-0.37)	0.26 (0.25-0.26)	0.31 (0.3-0.31)	0.32 (0.31-0.33)	0.04 (0.04-0.05)	0.21 (0.2-0.22)		
D	0.12 (0.12-0.13)	0.11 (0.1-0.11)	0.11 (0.11-0.12)	0.07 (0.07-0.08)	0.08 (0.07-0.09)	0.08 (0.07-0.08)		
G	0.6 (0.59-0.62)	1 (0.99-1.02)	0.81 (0.79-0.82)	0.52 (0.5-0.53)	0.49 (0.47-0.51)	0.51 (0.49-0.52)		
J	0.02 (0.02-0.03)	0.03 (0.03-0.04)	0.03 (0.03-0.03)	0.03 (0.02-0.03)	0.03 (0.02-0.03)	0.03 (0.02-0.03)		
L	0.51 (0.49-0.52)	0.07 (0.06-0.07)	0.28 (0.28-0.29)	0.37 (0.36-0.38)	0.05 (0.04-0.05)	0.24 (0.23-0.25)		
М	0.7 (0.69-0.72)	1.11 (1.09-1.13)	0.91 (0.9-0.92)	0.78 (0.76-0.8)	1.02 (0.99-1.04)	0.87 (0.86-0.89)		
N	22.45 (22.37-22.52)	35.68 (35.6-35.77)	29.17 (29.12-29.23)	21.51 (21.42-21.6)	21.02 (20.91-21.12)	21.31 (21.25-21.38)		
Р	0.08 (0.07-0.08)	0.24 (0.23-0.25)	0.16 (0.15-0.16)	0.08 (0.07-0.09)	0.19 (0.18-0.2)	0.12 (0.12-0.13)		
R	3.62 (3.59-3.65)	4.38 (4.34-4.41)	4 (3.98-4.03)	2.51 (2.48-2.54)	1.98 (1.94-2.02)	2.3 (2.27-2.32)		
S	0.41 (0.4-0.42)	0.42 (0.41-0.43)	0.41 (0.41-0.42)	0.33 (0.32-0.35)	0.1 (0.09-0.1)	0.24 (0.23-0.25)		
V	0 (0-0)	0 (0-0)	0 (0-0)	0 (0-0)	0 (0-0)	0 (0-0)		

Table 3. Frequency of the consumption of driving-impairing medicines by ATC group.

BMJ Open

Litic Chemical C. Lar system; D. Dermato. Sex hormones and insulins; Lusculo-skeletal system; N. Nervous sys. Torgans; V. Various. Abbreviations: ATC, Anatomical Therapeutic Chemical Classification System; A, Alimentary tract and metabolism; B, Blood and blood forming organs; C, Cardiovascular system; D, Dermatologicals; G, Genito-urinary system and sex hormones; H, Systemic hormonal preparations, excluding sex hormones and insulins; J, Antiinfectives for systemic use; L, Antineoplastic and immunomodulating agents; M, Musculo-skeletal system; N, Nervous system; P, Antiparasitic products, insecticides and repellents; R, Respiratory system; S, Sensory organs; V, Various.

ATC	Frequency	Ра	itients under treatm	ient	Drivers under treatment % (95%CD			
Group	of use		% (95%CI)					
		Males	Females	Total	Females	Males	Total	
Α	Acute	0.68	1.53	1.12	0.57	2.22	0.94	
		(0.61-0.74)	(1.44-1.62)	(1.06-1.18)	(0.5-0.64)	(1.96-2.47)	(0.86-1.02)	
	Sub-acute	17.7	29.63	23.87	17.69	53.18	25.62	
		(17.4 - 18)	(29.28-29.98)	(23.64-24.11)	(17.34-18.04)	(52.32-54.03)	(25.27-25.97)	
	Chronic	81.62	68.84	75.01	81.74	44.6	73.44	
		(81.32-81.93)	(68.49-69.19)	(74.77-75.24)	(81.38-82.09)	(43.75-45.46)	(73.09-73.8)	
Ν	Acute	18.06	12.79	14.79	18.02	16.55	17.45	
		(17.92-18.21)	(12.69-12.89)	(14.7-14.87)	(17.85-18.2)	(16.34-16.75)	(17.31-17.58)	
	Sub-acute	21.6	18.8	19.86	23.53	25.72	24.39	
		(21.44-21.76)	(18.68-18.91)	(19.76-19.95)	(23.34-23.72)	(25.48-25.97)	(24.24-24.54)	
	Chronic	60.33	68.41	65.36	58.45	57.73	58.17	
		(60.15-60.52)	(68.27-68.55)	(65.24-65.47)	(58.23-58.67)	(57.45-58.01)	(57.99-58.34)	
R	Acute	71.93	74.38	73.29	70.21	78.92	73.18	
		(71.5-72.36)	(74.01-74.76)	(73.01-73.58)	(69.61-70.81)	(78.17-79.66)	(72.71-73.65)	
	Sub-acute	22.63	21.93	22.24	23.86	18.49	22.03	
		(22.24-23.03)	(21.57-22.28)	(21.98-22.5)	(23.3-24.42)	(17.78-19.2)	(21.58-22.47)	
	Chronic	5.43	3.69	4.46	5.93	2.59	4.79	
		(5.22-5.65)	(3.53-3.85)	(4.33-4.6)	(5.62-6.24)	(2.3-2.88)	(4.56-5.02)	

Table 4. Frequency of the consumption of driving-impairing medicines A, N and R ATC group.

Abbreviations: ATC, Anatomical Therapeutic Chemical Classification System; A, Alimentary tract and metabolism; N, Nervous system; R, Respiratory system

BMJ Open

2 3	Figure 1 Frequency of modicing consumption in Castile and León in 2015
4	Figure 1. Frequency of medicine consumption in Castne and Leon in 2015.
5 6	
7	
8	
9 10	
11	
12 13	
14	
15	
17	
18	
19 20	
21	
22	
23 24	
25	
26 27	
28	
29 30	
31	
32	
33 34	
35	
36 37	
38	
39 40	
41	
42 43	
44	
45	
46 47	
48	
49 50	
51	
52 52	
53 54	
55	
56 57	
58	
59 60	
00	Sor poor roview only - http://hmionon?hmi.com/site/shout/guidelines.yhtm
	FOLDERLIEVIEW ONV = HUD.//OHHODEL.DHH.COH/SITE/ADDU//OHHOEHNES XNTH







152x187mm (300 x 300 DPI)

	No	Decommondation
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abet
The and abstract	1	Page 1
		(b) Provide in the abstract an informative and balanced summary of what was do
		and what was found
		Pages 2-3
In two dw ation		14503.2.5
Introduction Dealeground/rationala	2	Evaluin the scientific heatercound and rationals for the investigation being report
Dackground/rationale	2	Explain the scientific background and rationale for the investigation being report
Ohiaatiwaa	2	Fages 5-7
Objectives	3	State spectric objectives, including any prespectried hypotheses
		rage /
Methods		
Study design	4	Present key elements of study design early in the paper
		Pages 7-8
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitme
		exposure, follow-up, and data collection
		Pages 7-8
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of
		participants
		Page 7-8
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and eff
		modifiers. Give diagnostic criteria, if applicable
		Pages 9
Data sources/	8*	For each variable of interest, give sources of data and details of methods of
measurement		assessment (measurement). Describe comparability of assessment methods if the
		more than one group
		Page 9
Bias	9	Describe any efforts to address potential sources of bias
		Page 9
Study size	10	Explain how the study size was arrived at Pages 7-8
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable,
		describe which groupings were chosen and why
		Pages 9
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confoundi
		Page 10
		(b) Describe any methods used to examine subgroups and interactions
		Page 8 and 10
		(c) Explain how missing data were addressed
		Page 8
		(d) If applicable, describe analytical methods taking account of sampling strategy
		Page 10
		(<i>e</i>) Describe any sensitivity analyses
		Non applicable
Results		

For peer review only - http://bmjopen!bmj.com/site/about/guidelines.xhtml

		eligible, examined for eligibility, confirmed eligible, included in the study,
		Completing follow-up, and analysed
		Page 10
		(b) Give reasons for non-participation at each stage
		Non applicable
		(c) Consider use of a flow diagram
		Non applicable
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and
		information on exposures and potential confounders
		Page 10
		(b) Indicate number of participants with missing data for each variable of interest
		Pages 10-11
Outcome data	15*	Report numbers of outcome events or summary measures
		Pages 10-11
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and
		their precision (eg. 95% confidence interval). Make clear which confounders were
		adjusted for and why they were included
		Pages 11-12
		(b) Report category boundaries when continuous variables were categorized Pages
		11_12
		(a) If relevant, consider translating estimates of relative rick into absolute rick for a
		(c) If relevant, consider transfaring estimates of relative fisk into absolute fisk for a
		New empiricable
	1.7	
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and
		sensitivity analyses
		Non applicable
Discussion		
Key results	18	Summarise key results with reference to study objectives
		Page 12
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or
		imprecision. Discuss both direction and magnitude of any potential bias
		Page 16
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations,
		multiplicity of analyses, results from similar studies, and other relevant evidence
		Pages 12-16
Generalisability	21	Discuss the generalisability (external validity) of the study results
	21	Pages 12-15
Other information		
Funding	2.2.	Give the source of funding and the role of the funders for the present study and if
		applicable for the original study on which the present article is based
		Page 18
		1 460 10

*Give information separately for exposed and unexposed groups.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at

For peer review only - http://bmjopen?bmj.com/site/about/guidelines.xhtml

BMJ Open

http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.