



Article Trends in the Use of Driving-Impairing Medicines According to the DRUID Category: A Population-Based Registry Study with Reference to Driving in a Region of Spain between 2015 and 2019

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Abstract: The European DRUID (Drive Under the Influence of drugs, alcohol, and medicines) program classifies medications into three categories according to their effect on one's fitness to drive. The trend in the use of driving-impairing medicines (DIMs) in a region of Spain between 2015 and 2019 was analyzed through a population-based registry study. Pharmacy dispensing records for DIMs are provided. The use of DIMs on drivers was weighted according to the national driver's license census. The analysis was performed considering the population distribution by age and sex, treatment length, and the three DRUID categories. DIMs were used by 36.46% of the population and 27.91% of drivers, mainly chronically, with considerable daily use (8.04% and 5.34%, respectively). Use was more common in females than in males (42.28% vs. 30.44%) and increased with age. Among drivers, consumption decreases after 60 years of age for females and after 75 years of age for males. There was a 34% increase in the use of DIMs between 2015 and 2019, with a focus on daily use (>60%). The general population took 2.27 \pm 1.76 DIMs, fundamentally category II (moderate influence on fitness to drive) (20.3%) and category III (severe influence on fitness to drive) (19.08%). The use of DIMs by the general population and drivers is significant and has increased in recent years. The integration of the DRUID classification into electronic prescription tools would assist physicians and pharmacists in providing adequate information to the patient about the effects of prescribed medications on their fitness to drive.

Keywords: driving-impairing medicines; automobile driving; drug utilization; traffic accidents; driving under influence; DRUID classification

1. Introduction

According to the World Health Organization (WHO), injuries from traffic crashes will be the fifth leading cause of death by 2030, representing a serious public health problem worldwide [1].

Two of the most important factors to consider when assessing the risk of a traffic collision are age and sex. Older drivers have been shown to be up to 20 times more likely to be involved in a traffic crash than younger people [2], constituting the second leading cause of unintentional injury death for people over the age of 55 [3]. This is primarily because the skills needed to drive, such as attention, executive function, and visuospatial



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Copyright: © 2023 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). ability, decline with aging [4]. On the other hand, in terms of sex, males have higher risk values than females, especially at younger ages [5]. This may be due to several factors: females drive half as much as males [6] and show greater self-regulation behind the wheel, avoiding risky behaviors [7].

A significant proportion of these injuries can be attributed to driving after consuming psychoactive substances such as alcohol, illicit drugs, or certain medicines. Awareness of the role of these substances in driving is increasing, as is the implementation of appropriate interventions [8–10]. It is well known that alcohol and illicit drugs such as opioids, cocaine, and cannabis can impair a driver's psychomotor performance [11–14].

Driving a motor vehicle is a complex task that requires different cognitive and psychomotor capacities. The ability to drive a vehicle depends on the vision, brain, and musculoskeletal system performing the tasks involved in coordinated driving [15]. Some of the most commonly prescribed medicines can impair visual, cognitive, and/or motor skills necessary for safe driving [16]. Several groups of medicines can impair psychomotor performance, while the relationship between medication use and the risk of being involved in a traffic crash has been analyzed in several studies [9,10,17]. This association is evident for benzodiazepines [18–20], as well as for other commonly prescribed medicines such as antipsychotics, antidepressants, anxiolytics, opioid analgesics, etc. [12,15,21–24]. According to the European DRUID (Driving Under Influence of Alcohol and Drugs) project, the third and second most detected substance in traffic crashes and fatalities, respectively, was a benzodiazepine [25].

Due to the association between medicines and potential driving impairment, the European Union (EU) requires pharmaceutical companies to provide data on the effects of each drug on driving ability before marketing [26]. This information warns of possible effects that drivers may experience (drowsiness, dizziness, blurred vision, etc.) and that may significantly impair their fitness to drive [27].

Thus, some countries have chosen to print a driving pictogram on the outer packaging of certain medicines to warn of the potential risks while driving [28]. Currently, many European countries such as Spain, France, Austria, Denmark, Finland, among others, includes pictograms on certain medicines, which it is legally binding [29–31].

However, the population does not receive adequate information about the effects of medicines on driving ability. In Spain, for example, a study revealed that only 15.9% of the population was aware of the existence of the "medicines and driving" pictogram. The study also showed that most patients did not receive adequate information from healthcare professionals (physicians and pharmacists) about the effects of medicines on driving [28].

To address the problem of misinformation and to assist healthcare professionals in the medication selection process, several approaches to categorize medicines according to their effect on driving have been conducted [25,32]. One of the most comprehensive classifications is provided by the European DRUID project. According to this classification, the medicines are distinguished into three categories based on their effects on one's fitness to drive: I (minor influence), II (moderate influence) and III (severe influence) [27]. The definition of the DRUID categories was based on pharmacodynamic and kinetic data, pharmacovigilance data, experimental and epidemiological data, and additional data [27]. Considering these topics, the definitions of each DRUID category are shown in Supplementary Table S1 [27].

However, the DRUID classification does not include natural products that may affect psychomotor performance [33–36], which may be a handicap. In this sense, beneficial nutrients and antioxidants, including coenzyme Q10 (CoQ10) and alpha-tocopherol (α -Toc), and genistein proved to exhibit a neuroprotective effect attributed to its antioxidant, and/or anti-apoptotic properties [37,38] as well as α -Toc and/or CoQ10, where the underlying molecular mechanism of the treating compounds is related to the vascular endothelial growth factor (VEGF) and enhancing the brain-derived neurotrophic factor (BDNF)/tyrosine kinase B (TrkB)/cAMP-response element-binding protein (CREB) signaling pathway [39]. Additionally, the effects of the administration of the anti-amnestic

dose of St. John's wort extract and hyperforin on the ability to manage one's cognitive disturbance in psychotic and Huntington's disease patients were studied, among which the manifestation of prepulse inhibition (PPI) deficit was detected [40], wherein hyperforin was found to be one of the active ingredients responsible for St. John's wort-induced PPI disruption with no relation to apoptotic processes [41].

Based on previous research conducted by our group, approximately one in four drivers in Spain uses driving-impairing medicines (DIMs), with chronic use being the most common [42]. Up to now, trends in the use of DIMs have been analyzed: benzodiazepines [43], antipsychotics [44], opioids [45], antidepressants [46], antidiabetics [47], and antihistamines [48]. Now, we aim to analyze the use trend of DIMs according to the DRUID classification.

This study presents data on the consumption of DIMs by each DRUID category (I, II, and III) between 2015 and 2019 in Castile and Leon, a region of Spain with 2,323,770 inhabitants. The length of treatment and concomitant use among the different DRUID categories were specially addressed. Similarly to previous manuscripts, an estimate of the use of these medications in drivers was made. Lastly, all data are reported by sex and age group.

2. Results

Approximately 57.5 million packages of DIMs were dispensed in Castile and Leon between 2015 and 2019. For every 10 packages dispensed, 8 were medicines in Group N (central nervous system) of the Anatomic Therapeutic Chemical (ATC) classification, and 1 was a medicine in Group A (alimentary tract and metabolism). For each DRUID category, the most commonly used medicines were: lorazepam (DRUID III, 4.464.023 packages), codeine, and paracetamol (DRUID II, 1.211.750 packages) and metamizole (DRUID I, 4.753.078 packages) (Supplementary Table S2).

Overall, 36.4% of the general population took at least one DIM, mainly anxiolytics (14.27%), opioids (11.75%), antidepressants (8.75%), and other analgesics/antipyretics (8.25%). Chronic use (23.3%) predominated over subacute (6.72%) and acute (6.44%) use. Females were more likely to use DIM than males (42.28% vs. 30.44%, p = 0.001), and the use increased with age (Table 1, Figures 1 and 2).

| | Population Using DIM % (95CI) | | | | | | Drivers Using DIM % (95CI) | | | | | |
|-------------------|---------------------------------------|---------------------------------|----------------------------------|-------------------------------|---------------------------------------|----------------------------------|---------------------------------|---------------------------------|----------------------------------|--|--|--|
| | Yearly Use Daily Use | | Type of use | | | Yearly Use | Daily Use | Type of use | | | | |
| | icuity ese | | | Acute Subacute | | Chronic | | Acute | Subacute | Chronic | | |
| TOTAL DIM | | | | | | | | | | | | |
| Total | 36.46 (36.4–36.53) | 8.04 (8-8.07) | 6.44 (6.41–6.47) | 6.72 (6.69–6.75) | 23.3 (23.25–23.35) | 27.91 (27.83–27.98) | 5.34 (5.3–5.37) | 5.49 (5.46-5.53) | 6.26 (6.22–6.3) | $ \begin{array}{r} 16.15 \\ (16.09-16.21) \\ 15.42 \end{array} $ | | |
| Male | 30.44 (30.35–30.52) | 6.52 (6.47–6.56) | 6.27 (6.22–6.31) | 5.73 (5.69–5.78) | 18.44 (18.37–18.51) | 29.02 (28.93–29.12) | 6.18 (6.13–6.24) | 5.65 (5.6–5.69) | 5.96 (5.91–6.01) | 17.42 (17.34–17.5) | | |
| Female | 42.28 (42.19–42.37) | 9.5 (9.45–9.56) | 6.62 (6.57–6.66) | 7.67 (7.63–7.72) | 27.99 (27.91–28.07) | 26.25 (26.13–26.36) | 4.08 (4.03-4.13) | 5.26 (5.21–5.32) | 6.71 (6.65–6.77) | 14.27 (14.18–14.36) | | |
| DRUID I | $X^2 = 37,645.22;$ p = 0.001 | $X^2 = 19,471.66;$ p = 0.001 | $X^2 = 2716.22;$ p = 0.001 | $X^2 = 3222.11;$ p = 0.001 | $X^2 = 35,025.11;$ p = 0.001 | $X^2 = 282,817.29;$ p = 0.001 | $X^2 = 57,813.63;$ p = 0.001 | $X^2 = 42,935.41;$ p = 0.001 | $X^2 = 54,323.97;$ p = 0.001 | $X^2 = 185,912.23;$ p = 0.001 | | |
| Total | 17.11 (17.06–17.15) | 4.54 (4.51–4.56) | 3.68 (3.66–3.71) | 2.91 (2.89–2.94) | 10.5 (10.46–10.54) | 12.43 (12.38–12.48) | 3 (2.98–3.03) | 3.14 (3.11–3.17) | 2.25 (2.22–2.27) | 7.04 (7–7.08) | | |
| Male | 14.3 (14.24–14.36) | 3.83 (3.8–3.87) | 3.39 (3.35–3.42) | 2.3 (2.27–2.32) | 8.61 (8.56-8.66) | 13.57 (13.5–13.64) | 3.6 (3.56–3.64) | 3.33 (3.29–3.36) | 2.3 (2.27–2.33) | 7.94 (7.89–8) | | |
| Female | 19.82 (19.75–19.89) | 5.22 (5.18–5.26) | 3.97 (3.93–4) | 3.51 (3.48–3.55) | 12.33 (12.27–12.39) | 10.73 (10.65–10.81) | 2.12 (2.08–2.16) | 2.86 (2.82–2.9) | 2.17 (2.14–2.21) | 5.7 (5.64–5.76) | | |
| | $\dot{X}^2 = 30,920.70;$ p = 0.001 | $X^2 = 13,031.53;$ p = 0.001 | $X^2 = 3233.17;$ p = 0.001 | $X^2 = 2505.51;$ p = 0.001 | $\dot{X}^2 = 29,618.92;$ p = 0.001 | $X^2 = 142,137.07;$ p = 0.001 | $X^2 = 38,239.22;$ p = 0.001 | $X^2 = 21,685.44;$ p = 0.001 | $X^2 = 25,466.54;$ p = 0.001 | $X^2 = 96,994.06;$ p = 0.001 | | |
| DRUID II Total | 20.3 (20.25–20.35) | 2.72 (2.7–2.74) | 7.16 (7.13–7.19) | 4.12 (4.09–4.14) | 9.02 (8.98–9.06) | 15.2 (15.15–15.26) | 1.79 (1.77–1.81) | 5.7 (5.67–5.74) | 3.37 (3.34–3.4) | 6.13 (6.09–6.17) | | |
| Male | 16.55 (16.48–16.62) | 2.17 (2.14–2.2) | 6.21 (6.16–6.25) | 3.3 (3.27–3.34) | 7.03 (6.99–7.08) | 15.43 (15.35–15.5) | 2.04 (2.01–2.07) | 5.63 (5.58–5.67) | 3.24 (3.21–3.28) | 6.56 (6.51–6.61) | | |
| Female | 23.92 (23.84–24) | 3.24 (3.21–3.27) | 8.08 (8.03-8.13) | 4.9 (4.86–4.94) | 10.94 (10.88–10.99) | 14.87 (14.78–14.96) | 1.42 (1.39–1.45) | 5.82 (5.76–5.88) | 3.57 (3.52–3.62) | 5.49 (5.43–5.55) | | |
| DRUID III | $X^2 = 18,324.73;$ p = 0.001 | $X^2 = 6775.92;$ p = 0.001 | $X^2 = 4413.65.54;$ p = 0.001 | $X^2 = 2732.69;$ p = 0.001 | $X^2 = 13,583.15;$ P = 0.001 | $X^2 = 153,603.72;$ p = 0.001 | $X^2 = 15,132.36;$ p = 0.001 | $X^2 = 51,506.85;$ p = 0.001 | $X^2 = 36,306.531;$ p = 0.001 | $X^2 = 64,190.65;$ p = 0.001 | | |
| Total | 19.08 (19.03–19.13) | 2.62 (2.59–2.64) | 0.99 (0.98–1) | 4.58 (4.55-4.6) | 13.51 (13.47–13.56) | 13.62 (13.56–13.67) | 1.69 (1.67–1.71) | 0.81 (0.79–0.82) | 4.14 (4.11–4.18) | 8.67 (8.62-8.71) | | |
| Male | 13.27 (13.21–13.33) | 1.83 (1.81–1.86) | 0.76 (0.74–0.77) | 3.46 (3.43–3.5) | 9.05 (9–9.1) | 13.27 (13.2–13.34) | 1.84 (1.81–1.87) | 0.75 (0.73–0.77) | 3.73 (3.69–3.77) | 8.79 (8.73-8.85) | | |
| Female | 24.69 (24.61–24.76) | 3.37 (3.34–3.4) | 1.22 (1.2–1.23) | 5.65 (5.61-5.69) | 17.82 (17.75–17.89) | 14.13 (14.04–14.22) | 1.48 (1.45–1.51) | 0.9 (0.87–0.92) | 4.76 (4.7–4.81) | 8.48 (8.41-8.55) | | |
| | $X^2 = 13,929.87;$ p = 0.001 | $X^2 = 7836.51;$ p = 0.001 | $X^2 = 708.04;$ p = 0.001 | $X^2 = 648.79;$ p = 0.001 | $X^2 = 12,536.37;$ p = 0.001 | $X^2 = 139,899.19;$ p = 0.001 | $X^2 = 12,872.79;$ p = 0.001 | $X^2 = 8304.99;$ p = 0.001 | $X^2 = 34,128.12;$ p = 0.001 | $X^2 = 94,925.86;$ p = 0.001 | | |

Table 1. Use of DIMs according to the CONCYLIA database and the Castile and León drivers' license census data.

Abbreviations: DIM, driving-impairing medicines, 95 CI, confidence interval.

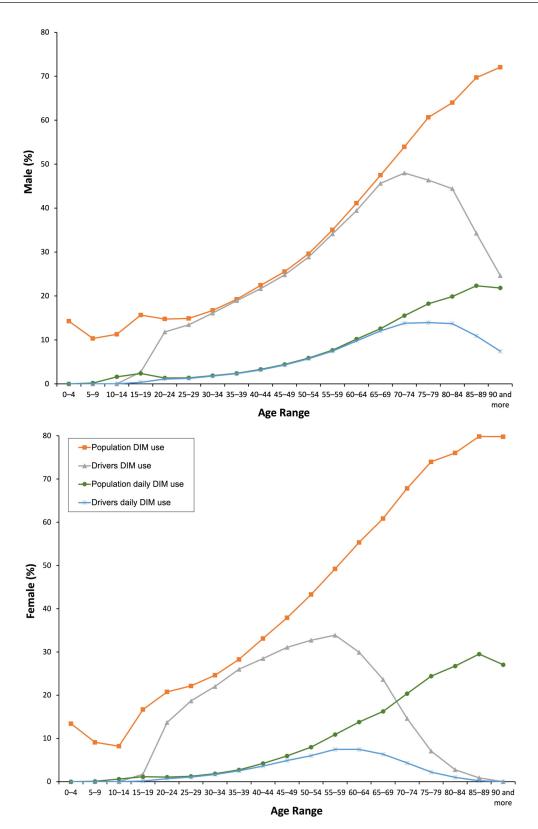


Figure 1. Frequency of the use of DIMs by the general population and the driver population.

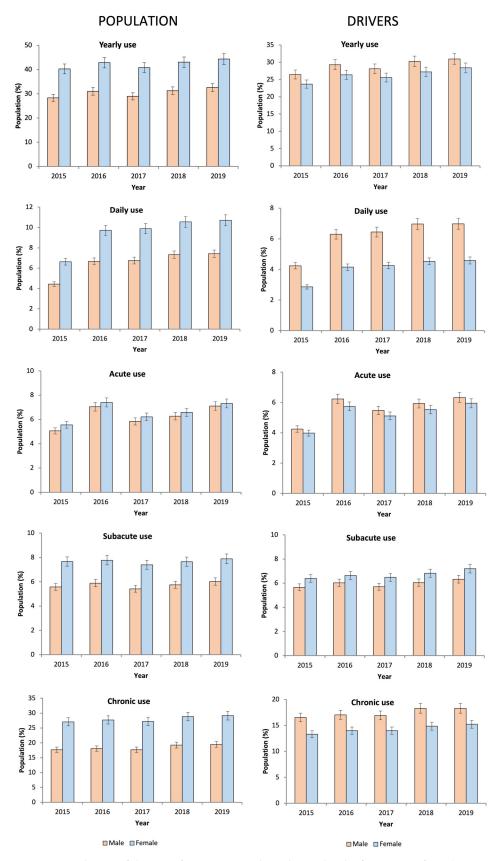
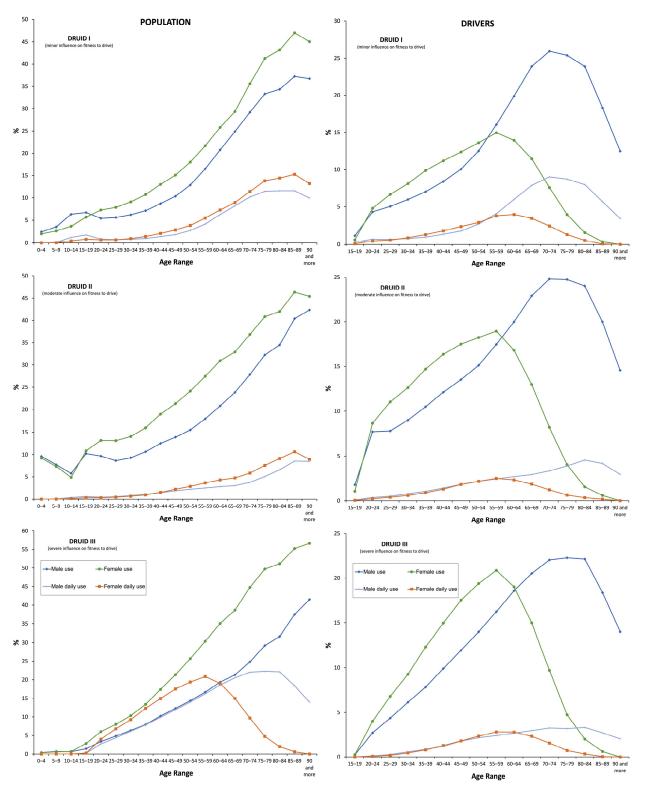


Figure 2. Evolution of the use of DIMs in Castile and Leon by the frequency of use (2015–2019).

The yearly use of DIMs increased by approximately 12% (38.60% in 2019 vs. 34.41% in 2015, p = 0.001), with a particular impact on acute use, which increased by 35.89%



(7.22% in 2019 vs. 5.31% in 2015, p = 0.001), followed by chronic use (8.68%, 24.41% in 2019 vs. 22.46% in 2015, p = 0.001) and subacute use (4.98%, 6.97% in 2019 vs. 6.64% in 2015, p = 0.001) (Figure 3).

Figure 3. Frequency of the use of DIMs use by DRUID classification.

The general population preferentially used DRUID II medicines (20.3%), followed by DRUID III medicines (19.08%) and DRUID I medicines (17.11%) (Table 1, Figure 2).

Nevertheless, the highest increase in use was observed for DRUID III medicines (19.59%, 20.33% in 2019 vs. 17% in 2015, p = 0.001), followed by DRUID I medicines (17.42%, 18.26% in 2019 vs. 15.55% in 2015, p = 0.001) and DRUID II medicines (12.93%, 21.91% in 2019 vs. 19.4% in 2015, p = 0.001) (Table 2).

Regardless of medicine use, yearly users took 2.27 ± 1.76 DIMs, increasing to 2.87 ± 1.94 for chronic users. Consumption was higher in females than in males, except for the medicines in the DRUID I category (Table 3).

As for daily use, 8.04% of the general population (Table 1) took a mean of 1.35 ± 0.68 DIMs (1.36 ± 0.7 in males vs. 1.35 ± 0.67 in females, p = 0.001) (Table 3). The greatest increase in use was observed among daily users (63.95%, 9.10% in 2019 vs. 5.55% in 2015, p = 0.001) (Figure 2), which was higher for DRUID II medicines (33.49%, 2.97% in 2019 vs. 2.23% in 2015, p = 0.001) than for DRUID III medicines (31.9%, 2.84% in 2019 vs. 2.15% in 2015, p = 0.001) and DRUID I medicines (29.3%, 5% in 2015 vs. 3.87 in 2015, p = 0.001).

In terms of concomitant use, 46.56% of the general population using a DRUID III medicine also used a DRUID II and 44.74% used a DRUID I medicine. However, with daily use, the concomitant use of DRUID III medicines increased to 61.67% for DRUID II and 52.81% for DRUID I medicines.

Concerning drivers, 27.91% took at least one DIM, which was higher in males than in females (29.02% vs. 26.25%). As in the general population, the use of DIMs was predominantly chronic (16.15%) with a higher use of DRUID II category medicines (15.20%) (Table 1). The use of DIMs increased with age, peaking at 55–59 years for females and 70–74 years for males (Figures 1 and 2, respectively). The increase in the use of DIMs was higher than in the general population, especially in yearly use (17.96%, 29.91% in 2019 vs. 25.35% in 2015, p = 0.001) (Figure 3) and in DRUID III medicines use (26.96%, 14.71% in 2019 vs. 11.61% in 2015, p = 0.001) (Table 2). Finally, the concomitant use of medicines from different DRUID categories was similar to the general population.

| | | Popula | tion Using DIM ^o | % (95CI) | | Drivers Using DIM % (95CI) | | | | | |
|----------------|------------------|------------------|-----------------------------|---------------------------|------------------|----------------------------|--------------------|--------------------|--------------------|-------------------|--|
| _ | 2015 | 2016 | 2017 | 2018 | 2019 | 2015 | 2016 | 2017 | 2018 | 2019 | |
| DRUID category | | | | | | | | | | | |
| I | | | | | | | | | | | |
| Total | 15.55 | 17.14 | 16.73 | 17.85 | 18.26 | 11.02 | 12.41 | 12.24 | 13.03 | 13.45 | |
| Total | (15.5–15.59) | (17.09–17.19) | (16.69–16.78) | (17.81 - 17.9) | (18.21–18.31) | (10.97 - 11.07) | (12.36 - 12.47) | (12.18-12.29) | (12.97 - 13.08) | (13.39–13.5) | |
| Mala | 12.69 | 14.23 | 13.97 | 15.09 | 15.52 | 12.01 | 13.5 | 13.4 | 14.34 | 14.6 | |
| Male | (12.63–12.76) | (14.17 - 14.29) | (13.9–14.03) | (15.02 - 15.15) | (15.45–15.58) | (11.95–12.08) | (13.43–13.58) | (13.33–13.47) | (14.27 - 14.42) | (14.53–14.67) | |
| Female | 18.31 | 19.95 | 19.41 | 20.53 | 20.89 | 9.51 | 10.77 | 10.51 | 11.1 | 11.76 | |
| remale | (18.24–18.38) | (19.88–20.02) | (19.34–19.48) | (20.46–20.6) | (20.82-20.97) | (9.44–9.59) | (10.69 - 10.85) | (10.43 - 10.59) | (11.02–11.18) | (11.67–11.84) | |
| | $X^2 = 6242.63;$ | $X^2 = 6220.82;$ | $X^2 = 6371.31;$ | $X^2 = 6240.85;$ | $X^2 = 6243.2;$ | $X^2 = 26,691.39;$ | $X^2 = 28,349.01;$ | $X^2 = 27,655.09;$ | $X^2 = 29,166.77;$ | $X^2 = 30,466.79$ | |
| | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | |
| II | | | | | | | | | | | |
| Total | 19.4 | 21.07 | 18.68 | 20.43 | 21.91 | 14.02 | 15.52 | 14.23 | 15.72 | 16.54 | |
| Iotal | (19.35–19.45) | (21.02–21.12) | (18.63–18.73) | (20.38–20.49) | (21.86–21.97) | (13.96–14.07) | (15.46–15.58) | (14.17 - 14.28) | (15.66–15.78) | (16.48–16.6) | |
| Male | 15.84 | 17.34 | 14.98 | 16.66 | 17.94 | 14.53 | 15.82 | 14.26 | 15.89 | 16.64 | |
| Iviale | (15.77–15.9) | (17.27 - 17.41) | (14.91 - 15.04) | (16.59–16.73) | (17.87 - 18.01) | (14.46 - 14.6) | (15.75–15.9) | (14.18–14.33) | (15.82–15.97) | (16.56–16.72) | |
| Female | 22.84 | 24.68 | 22.27 | 24.08 | 25.74 | 13.24 | 15.06 | 14.18 | 15.47 | 16.38 | |
| remate | (22.77–22.92) | (24.6–24.75) | (22.19–22.34) | (24.01–24.16) | (25.66–25.81) | (13.15–13.32) | (14.97–15.15) | (14.09 - 14.27) | (15.37–15.56) | (16.29–16.48) | |
| | $X^2 = 4212.31;$ | $X^2 = 3734.73;$ | $X^2 = 3259.54;$ | X ² = 3234.94; | $X^2 = 4093.43;$ | $X^2 = 29,478.34;$ | $X^2 = 32,470.1;$ | $X^2 = 28,803.92;$ | $X^2 = 30,527.18;$ | $X^2 = 32,463.86$ | |
| | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | |
| III | | | | | | | | | | | |
| Total | 17 | 19.17 | 18.85 | 20.05 (20-20.1) | 20.33 | 11.61 | 13.65 | 13.57 | 14.51 | 14.74 | |
| Iotai | (16.95–17.04) | (19.12–19.22) | (18.8 - 18.9) | | (20.28–20.38) | (11.56–11.66) | (13.6–13.71) | (13.52–13.63) | (14.46 - 14.57) | (14.68 - 14.79) | |
| Male | 11.52 | 13.27 | 13.02 | 14.14 | 14.39 | 11.49 | 13.24 | 13.15 | 14.18 | 14.3 | |
| wale | (11.46–11.58) | (13.21–13.34) | (12.96–13.08) | (14.08 - 14.21) | (14.33 - 14.46) | (11.42–11.56) | (13.17–13.31) | (13.07–13.22) | (14.11–14.25) | (14.23–14.38) | |
| Female | 22.29 | 24.87 | 24.48 | 25.76 | 26.04 | 11.8 | 14.28 | 14.21 | 15 (14.91–15.1) | 15.38 | |
| i cinule | (22.22–22.37) | (24.79–24.94) | (24.4–24.56) | (25.68–25.84) | (25.96–26.12) | (11.72–11.88) | (14.19–14.36) | (14.12 - 14.3) | . , , , | (15.28–15.47) | |
| | $X^2 = 3115.65;$ | $X^2 = 2544.38;$ | $X^2 = 2879.54;$ | $X^2 = 2704.39;$ | $X^2 = 2772.62;$ | $X^2 = 24,930.35;$ | $X^2 = 28,892.67;$ | $X^2 = 27,808.57;$ | $X^2 = 29,052.18;$ | $X^2 = 29,466.38$ | |
| | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | p = 0.001 | |

Table 2. Evolution of the use of DIMs according to CONCYLIA database and the Castile and León drivers' license census data (2015–2019).

Abbreviations: DIM, driving-impairing medicines, 95CI, confidence interval.

| | DRUID Category | | Population U | Jsing DIM (Mea | $n \pm SD$) | Drivers Using DIM(Mean \pm SD) | | | | |
|------------------|-------------------|-----------------|------------------|-----------------|------------------------|----------------------------------|-----------------|-----------------|-----------------------|--|
| Frequency of Use | | Males | Females | TOTAL | t, <i>p</i> | Males | Females | TOTAL | t, <i>p</i> | |
| | DRUID I | 1.02 ± 0.13 | 1.02 ± 0.15 | 1.02 ± 0.14 | t = -3.21; p = 0.078 | 1.02 ± 0.13 | 1.02 ± 0.14 | 1.02 ± 0.13 | t = -5.27; p = 0.093 | |
| A / | DRUID II | 1.03 ± 0.17 | 1.04 ± 0.19 | 1.03 ± 0.18 | t = -23.70; p = 0.001 | 1.03 ± 0.17 | 1.03 ± 0.19 | 1.03 ± 0.18 | t = -79.87; p = 0.043 | |
| Acute | DRUID III | 1 ± 0.06 | 1 ± 0.05 | 1 ± 0.05 | t = 1.28; p = 0.199 | 1 ± 0.05 | 1 ± 0.04 | 1 ± 0.05 | t = 3.51; p = 0.134 | |
| | TOTAL | 1.03 ± 0.19 | 1.04 ± 0.21 | 1.03 ± 0.2 | t = -21.9; p = 0.001 | 1.03 ± 0.19 | 1.04 ± 0.2 | 1.03 ± 0.19 | t = -11.9; p = 0.001 | |
| | DRUID I | 1.12 ± 0.34 | 1.12 ± 0.35 | 1.12 ± 0.35 | t = -2.34; p = 0.123 | 1.12 ± 0.35 | 1.14 ± 0.36 | 1.13 ± 0.35 | t = -7.89; p = 0.001 | |
| Sub-acute | DRUID II | 1.25 ± 0.49 | 1.32 ± 0.55 | 1.29 ± 0.53 | t = -42.65; p = 0.001 | 1.25 ± 0.49 | 1.32 ± 0.55 | 1.28 ± 0.51 | t = -31.06; p = 0.001 | |
| | DRUID III | 1.08 ± 0.29 | 1.11 ± 0.32 | 1.1 ± 0.31 | t = -25.87; p = 0.001 | 1.08 ± 0.28 | 1.1 ± 0.31 | 1.09 ± 0.3 | t = -17.77; p = 0.001 | |
| | TOTAL | 1.37 ± 0.61 | 1.44 ± 0.66 | 1.41 ± 0.64 | t = -44.32; p = 0.001 | 1.38 ± 0.61 | 1.43 ± 0.65 | 1.4 ± 0.63 | t = -28.8; p = 0.001 | |
| | DRUID I | 1.64 ± 0.92 | 1.58 ± 0.9 | 1.6 ± 0.91 | t = 31.810; p = 0.001 | 1.65 ± 0.93 | 1.46 ± 0.79 | 1.59 ± 0.89 | t = 75.72; p = 0.001 | |
| Chronic | DRUID II | 1.9 ± 1.26 | 1.98 ± 1.25 | 1.95 ± 1.25 | t = -29.71; p = 0.001 | 1.95 ± 1.31 | 2.06 ± 1.37 | 1.99 ± 1.33 | t = -27.55; p = 0.001 | |
| | DRUID III | 1.56 ± 0.87 | 1.68 ± 0.94 | 1.64 ± 0.92 | t = -74.74; p = 0.001 | 1.58 ± 0.88 | 1.7 ± 0.97 | 1.63 ± 0.92 | t = -53.21; p = 0.001 | |
| | TOTAL | 2.66 ± 1.84 | 3 ± 1.99 | 2.87 ± 1.94 | t = -44.32; p = 0.001 | 2.69 ± 1.87 | 2.92 ± 2.01 | 2.77 ± 1.92 | t = -65.66; p = 0.001 | |
| | DRUID I | 1.41 ± 0.78 | 1.39 ± 0.77 | 1.4 ± 0.77 | t = 15.09; p = 0.001 | 1.4 ± 0.78 | 1.28 ± 0.64 | 1.36 ± 0.74 | t = 84.04; p = 0.001 | |
| Voorly | DRUID II | 1.44 ± 0.95 | 1.52 ± 0.99 | 1.49 ± 0.97 | t = -62.85; p = 0.001 | 1.46 ± 0.98 | 1.48 ± 0.99 | 1.47 ± 0.99 | t = -9.29; p = 0.001 | |
| Yearly | DRUID III | 1.4 ± 0.77 | 1.51 ± 0.85 | 1.47 ± 0.83 | t = -94.53; p = 0.001 | 1.4 ± 0.77 | 1.46 ± 0.83 | 1.43 ± 0.8 | t = -31.72; p = 0.001 | |
| | TOTAL | 2.06 ± 1.61 | 2.404 ± 1.83 | 2.27 ± 1.76 | t = -204.39; p = 0.001 | 2.07 ± 1.64 | 2.15 ± 1.73 | 2.10 ± 1.67 | t = -32.08; p = 0.001 | |

Table 3. Average DIMs consumed in Castile and Leon by frequency of use.

Abbreviations: DIM, driving-impairing medicines, SD, standard deviation.

3. Discussion

According to our findings, nearly 36% of the general population and 26% of drivers took a DIM from 2015 to 2019. The use of DIMs increased with age and was prevalent among females in the general population and among male drivers. The chronic use and consumption of DRUID II medicines were the most prominent. The general population chronically used about three different DIMs, mainly central nervous system (ATC group N) medicines, especially anxiolytics and opioids, and alimentary tract and metabolism (ATC group A) medicines, especially oral antidiabetics and insulins. In each year of the study period, we detected a relevant concomitant use of different DIMs, such that half of the DRUID III medicines users were also taking a DRUID II and a DRUID I medicine. In addition, the trend of DIMs use was increasing, especially for DRUID III medicines.

To our knowledge, this is the first study to evaluate the use of DIMs according to the DRUID classification. DIMs in the DRUID II category (severe influence on fitness to drive) were the most frequently used, followed closely by DIMs in the DRUID III category (severe influence on fitness to drive), whose use trend showed the greatest increase between 2015 and 2019. These data constitute a serious road safety issue, since many patients, especially drivers, lack sufficient information on how to act when driving vehicles [28].

The use of DIMs, regardless of the DRUID category, is generally higher in females among the general population and in male drivers. This is most likely because in Spain males are the majority behind the wheel. Moreover, the number of driving licenses for females decreases with age [42–46,48].

Like other medicines, DIM use increases with age [49]. These values were predictable, since our region is characterized by an elderly population with an increasing rate of polymedication [50], especially in patients with mental disorders [51], for whom DIMs are mostly indicated.

However, the ability of the elderly to drive safely is a controversial issue. Mental and physical changes accompany aging [52], such as the decline of cognitive, visual, motor, and sensory functions [53]. These factors increase the risk of traffic crashes [52], especially after the age of 80 [54].

In this sense, the Spanish General Regulations for Drivers [55] state that drivers over the age of 65 must renew their license every 5 years, instead of the 10 years required for drivers under the age of 65. In addition, a medical examination to assess the driver's psychophysical condition is mandatory for license renewal. Related to this, Appendix 4 of these regulations [55] contains a list of diseases that restrict the renewal of a driver's license. In short, in Spain, the renewal of a driver's license depends on the state of the driver's abilities and aptitudes.

The chronic use of DIMs was the most common, with the significant prevalence of daily use. In this sense, the impairment of fitness to drive decreases with chronic and stable medication use [56] due to the tolerance factor [24]. On the other hand, the initiation of treatment, dosage changes, acute use, and multiple daily doses are the most dangerous circumstances for drivers, especially after the age of 50 [8,18,56,57]. The number of medicines consumed should also be considered, as polypharmacy is associated with a higher likelihood of being involved in a traffic crash [57,58]. In the present study, the concomitant use of different DIMs was considerable. This scenario must be considered, especially when using DIMs classified as DRUID II and III (moderate and severe influence on fitness to drive, respectively). This use dramatically increases the risk of a traffic crash, especially if the driver has also consumed alcohol [58].

The present findings seem to be consistent with other European research which has found that central nervous system medicines (ATC group N) are the most commonly consumed [42,58]. Basically, the use of benzodiazepines (anxiolytics), antidepressants and opioid analgesics is predominant and has been increasing in recent years [43,45,46]. The relationship between traffic crashes and the use of benzodiazepines [18–20] and opioid analgesics [12,24] is well established, particularly in acute use and with increasing doses. Furthermore, among people taking antidepressants, the effect on fitness to drive is not only

due to the influence of the medication but also to the depression symptoms [59–61]. However, contradictory data on antidepressants are available in the literature. Medicines such as mirtazapine or tricyclic antidepressants can influence driving, mainly at the beginning of treatment [22,23], although selective serotonin reuptake inhibitors (SSRIs) affect driving only at high doses or when used in combination with other DIMs [62–64]. Most of these medicines, except for some types of antidepressants, are classified as DRUID II and DRUID III (severe influence on fitness to drive) [27].

Far behind the above, the next most used DIMs were medicines for the alimentary tract and metabolism (ATC group A). This group includes oral antidiabetics and insulins, which are classified as DRUID I (minor influence on fitness to drive) [27]. These figures, although higher than in other European countries [65], are consistent with a previous study conducted by our group [47] and other national data [66]. Diabetes mellitus is known to affect the fitness to drive due to chronic complications such as retinopathy and neuropathy [67,68] and especially acute complications such as hypoglycemia and hyperglycemia [68,69]. Indeed, hypoglycemia can be a direct consequence of the use of medicines such as insulin [70–72] and oral antidiabetic agents such as sulfonylureas and methyglinides [71,73]. Therefore, restrictions have been imposed in Europe and the United States for diabetics to obtain a driver's license [69,74–76].

The DRUID classification was developed by the EU in order to harmonize the criteria for classifying the medicines marketed in Europe according to their influence on fitness to drive (https://www.emcdda.europa.eu/publications/thematic-papers/druid_en, accessed on 25 February 2020). However, this categorization is not without limitations. The DRUID category is assigned based on the assumption that the medicine is used for its primary indication, is administered to an adult, at a normal dose, and at the beginning of the treatment [27]. Nevertheless, the DRUID classification has proven to be suitable for being incorporated into computerized pharmaceutical dispensing systems [77].

Information effectively provided by healthcare professionals, especially physicians and pharmacists, has been shown to have the potential to reduce the annual rate of traffic crashes by up to 45% [28]. In Spain, other warning systems, such as the "medicines and driving" pictogram, have not been fully effective [28]. Therefore, the incorporation of DRUID categorization into prescribing and dispensing tools must be a primary objective for healthcare authorities.

The most important recommendation for physicians and pharmacists is the need to effectively counsel their patients about the effects of DIMs on fitness to drive [78]. According to the conclusions of the DRUID project, health counseling should be brief, individualized, and provided at the time that a DIM is prescribed or dispensed [77]. In addition, following the recommendations of the DRUID project, prescribing/dispensing guidelines should be established based on the use of safer driving medications when available [77].

Specifically, ensuring the safe and effective use of medicines is one of the most important roles of a pharmacist [79]. In this sense, the pharmacist plays an important role in achieving patients' health goals, monitoring pharmacologic treatment, collaborating with the primary care team, and working in health promotion and disease prevention. Definitely, the pharmaceutical profession is well positioned in the healthcare system and should take the lead in educational programs for the driving population using DIMs.

Finally, the limitations of a retrospective observational design and those inherent to data extraction from health administrative databases should be mentioned. CONCYLIA does not include information on hospital dispensing, consumption derived from private practice, or "over the counter" dispensing. However, this bias is not considered relevant, since nearly 95% of the population in our region is covered by public health insurance and a medical prescription is mandatory for the dispensation of DIMs. Another bias to consider is that CONCYLIA does not include information about whether the patient is a driver. Therefore, a weighting method was established using the driver's license census, adjusting for age and sex. This method has been extensively tested in previous research by our team [42–48]. On the other hand, it has been assumed that the individual with a

driver's license is an active driver, but this is not always the case, especially among the elderly. Furthermore, comorbidities in the elderly, such as depression, Parkinson's disease, insomnia, hypertension, congestive heart failure, osteoarthritis, diabetes mellitus, and others, which certainly affect one's fitness to drive, were not considered [4]. Lastly, an important limitation is the lack of data on cannabis and marijuana use, which can seriously affect cognition, psychomotor function, and driving performance [80].

4. Materials and Methods

This work followed the same methodology as previously described in previous studies conducted by our group [42–48] and is presented below.

4.1. Real-World Study Details

The findings in this manuscript are presented from a population-based registry study following the STROBE [81] (Strengthening the Reporting of Observational Studies in Epidemiology) and RECORD [82] (Reporting of studies Conducted using Observational Routinely-collected Data) recommendations for observational studies.

All medicines marketed in Spain with the pictogram "medicines and driving" are considered DIMs. In this sense, all DIM dispensations in pharmacies in Castile and Leon from 2015 to 2019 were considered. As in previous manuscripts [42–48], dispensing was considered to be an approximation of the actual consumption. The total population of Castile and Leon was considered (Table S3). Figure 4 shows the flow chart of the inclusion of the study population.

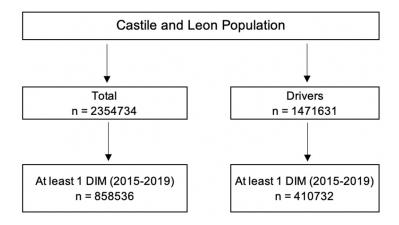


Figure 4. Flowchart of the inclusion of the study population.

Data on the use of DIMs were extracted from the Pharmacy Information System of the Castile and Leon Health System, CONCYLIA (http://www.saludcastillayleon.es/portalmedicamento/es/indicadores informes/concylia, accessed on 3 February 2020). This database includes, but is not limited to: dispensing code and date; pharmacy identification; patient sex and age; generic medicine name; number of packages dispensed; length of treatment; etc. In the CONCYLIA database, medication classification is based on the Anatomical Therapeutic Chemical Code (ATC) [83]. However, CONCYLIA does not collect data on dispensations from hospitals, private practices, and "over the counter medicines".

In addition, the National Department of Traffic (Ministry of Interior) (http://www. dgt.es/es/seguridad-vial/estadisticas-e-indicadores/permisos-conduccion/, accessed on 25 January 2020) provided access to drivers' license census data. Subsequently, as in previous manuscripts [42–48], the use of DIMs in drivers was adjusted (Table S3).

The Valladolid East Health Area Ethics Committee approved this study on 17 March 2016 (reference number PI 16-387).

4.2. Variables

The variables considered were: (1) yearly frequency of the use of DIMs; (2) acute, sub-acute, and chronic use of DIMs; (3) daily use of DIMs; (4) yearly frequency of DIMs use by DRUID categories; and (5) concomitant use of DIMs of diverse DRUID categories.

The length of treatment was determined to be between 1 and 7 days for acute use, between 8 and 29 days for subacute use, and more than 29 days for chronic use. Yearly use is equivalent to the use of at least one DIM during the year.

4.3. Statistical Analysis

Sex and age were considered for all analyses. Frequencies (percentages) with their 95% confidence interval (95% CI) or means with their standard deviation (SD) were used to report the results. Differences between the continuous variables were analyzed by the Student's *t*-test (t), while differences between the categorical variables were analyzed by the Chi-squared test (χ^2). The trend in DIM use during the study period was analyzed using the Cochran–Armitage trend test. The statistical significance level was set at $p \leq 0.05$. Finally, all statistical analyses were performed using the Statistical Package for the Social Sciences (SPSS version 24.0., SPSS Inc, Chicago, IL, USA).

5. Conclusions

During the study period, the use of DIMs was frequent and increasing. The use and chronic use of DRUID II and III medicines, which have the greatest impact on fitness to drive, are highlighted. The aging of our region, the high use of DIMs, and polypharmacy increase the risk of having a traffic crash [8,18,56–58]. Thus, the knowledge of how medications affect one's fitness to drive is clearly a relevant road safety issue for patients, physicians, pharmaceutical companies, policymakers, and the general population [84].

The DRUID classification has the necessary characteristics to be useful for physicians and pharmacists to inform drivers about the side effects (cognitive and psychomotor impairment) associated with the use of DIMs [85]. This action is more meaningful for the pharmacist, as they are usually the last opportunity for the patient to be informed before starting treatment [86].

Supplementary Materials: The following supporting information can be downloaded at: https: //www.mdpi.com/article/10.3390/ph16040508/s1, Table S1: Definition of DRUID categories; Table S2: List of the 20 most consumed DIMs according to DRUID classification in Castile and Leon during the study period (packages/year); Table S3: Evolution of the Castile and Leon population and driver's licenses (2015–2019).

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Institutional Review Board Statement: This study was conducted according to the guidelines of the Declaration of Helsinki and approved by the East Valladolid Health Area Ethics Committee (PI 16-387, 17 March 2016).

Informed Consent Statement: Patient consent was waived due to the fact that this was a retrospective observational study, and anonymized databases provided by the health authorities were used.

Data Availability Statement: Restrictions apply to the availability of these data. Data were obtained from regional health authorities (Gerencia Regional de Salud (GRS)) and may be requested from conciertofco@saludcastillayleon.es (GRS).

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References

- 1. World Health Organization (WHO). Report on the Global Situation of Road Safety. Available online: https://www.who.int/publications/i/item/9789241565684 (accessed on 21 June 2020).
- Pitta, L.S.R.; Quintas, J.L.; Trindade, I.O.A.; Belchior, P.; Gameiro, K.d.S.D.; Gomes, C.M.; Nóbrega, O.T.; Camargos, E.F. Older Drivers Are at Increased Risk of Fatal Crash Involvement: Results of a Systematic Review and Meta-Analysis. *Arch. Gerontol. Geriatr.* 2021, 95, 104414. [CrossRef]
- Centers for Disease Control Prevention. National Center for Injury Prevention and Control: Web-Based Injury Statistics Query And Reporting System (WISQARS). Washington, DC, US Department of Health and Human Services. Available online: https://www.cdc.gov/injury/wisqars/index.html (accessed on 20 March 2023).
- Hagiya, H.; Takase, R.; Honda, H.; Nakano, Y.; Otsuka, Y.; Kataoka, H.; Uno, M.; Ueda, K.; Takahashi, M.; Ogawa, H.; et al. Prevalence of Medical Factors Related to Aging among Older Car Drivers: A Multicenter, Cross-Sectional, Descriptive Study. BMC Geriatr. 2022, 22, 792. [CrossRef] [PubMed]
- 5. Santamariña-Rubio, E.; Pérez, K.; Olabarria, M.; Novoa, A.M. Gender Differences in Road Traffic Injury Rate Using Time Travelled as a Measure of Exposure. *Accid. Anal. Prev.* 2014, *65*, 1–7. [CrossRef] [PubMed]
- Mateos-Granados, J.; Martín-delosReyes, L.M.; Rivera-Izquierdo, M.; Jiménez-Mejías, E.; Martínez-Ruiz, V.; Lardelli-Claret, P. Sex Differences in the Amount and Patterns of Car-Driving Exposure in Spain, 2014 to 2017: An Application of a Quasi-Induced Exposure Approach. *Int. J. Environ. Res. Public Health* 2021, 18, 13255. [CrossRef] [PubMed]
- Gwyther, H.; Holland, C. The Effect of Age, Gender and Attitudes on Self-Regulation in Driving. *Accid. Anal. Prev.* 2012, 45, 19–28. [CrossRef]
- 8. Ravera, S.; Hummel, S.A.; Stolk, P.; Heerdink, R.E.; de Jong-van den Berg, L.T.W.; de Gier, J.J. The Use of Driving Impairing Medicines: A European Survey. *Eur. J. Clin. Pharmacol.* **2009**, *65*, 1139–1147. [CrossRef]
- 9. Engeland, A.; Skurtveit, S.; Mørland, J. Risk of Road Traffic Accidents Associated with the Prescription of Drugs: A Registry-Based Cohort Study. *Ann. Epidemiol.* 2007, 17, 597–602. [CrossRef]
- 10. Orriols, L.; Salmi, L.-R.; Philip, P.; Moore, N.; Delorme, B.; Castot, A.; Lagarde, E. The Impact of Medicinal Drugs on Traffic Safety: A Systematic Review of Epidemiological Studies. *Pharmacoepidemiol. Drug Saf.* **2009**, *18*, 647–658. [CrossRef]
- Herrera-Gómez, F.; García-Mingo, M.; Álvarez, F.J. Prevalence of Alcohol and Other Psychoactive Substances in Motor Vehicle Drivers in Spain, 2018: Cross-Sectional Dataset Analysis with Studies from 2008 and 2013. *Forensic Sci. Int.* 2020, 313, 110266. [CrossRef]
- Herrera-Gómez, F.; García-Mingo, M.; Colás, M.; González-Luque, J.C.; Álvarez, F.J. Opioids in Oral Fluid of Spanish Drivers. Drug Alcohol Depend. 2018, 187, 35–39. [CrossRef]
- 13. Herrera-Gómez, F.; Gutiérrez-Abejón, E.; García-Mingo, M.; Álvarez, F.J. Positivity to Cocaine and/or Benzoylecgonine in Confirmation Analyses for On-Road Tests in Spain. *Int. J. Environ. Res. Public Health* **2021**, *18*, 5371. [CrossRef]
- 14. Herrera-Gómez, F.; García-Mingo, M.; Colás, M.; González-Luque, J.C.; Alvarez, F.J. Drivers Who Tested Positive for Cannabis in Oral Fluid: A Longitudinal Analysis of Administrative Data for Spain between 2011 and 2016. *BMJ Open* **2019**, *9*, e026648. [CrossRef]
- 15. Hetland, A.; Carr, D.B. Medications and Impaired Driving: A Review of the Literature. *Ann. Pharmacother.* **2014**, *48*, 494–506. [CrossRef]
- 16. Ravera, S.; van Rein, N.; de Gier, J.J.; de Jong-van den Berg, L.T.W. A Comparison of Pharmacoepidemiological Study Designs in Medication Use and Traffic Safety Research. *Eur. J. Epidemiol.* **2012**, *27*, 473–481. [CrossRef]
- Orriols, L.; Delorme, B.; Gadegbeku, B.; Tricotel, A.; Contrand, B.; Laumon, B.; Salmi, L.-R.; Lagarde, E. CESIR research group Prescription Medicines and the Risk of Road Traffic Crashes: A French Registry-Based Study. *PLoS Med.* 2010, 7, e1000366. [CrossRef]
- 18. Barbone, F.; McMahon, A.D.; Davey, P.G.; Morris, A.D.; Reid, I.C.; McDevitt, D.G.; MacDonald, T.M. Association of Road-Traffic Accidents with Benzodiazepine Use. *Lancet* **1998**, *352*, 1331–1336. [CrossRef]
- 19. Walsh, J.M.; de Gier, J.J.; Christopherson, A.S.; Verstraete, A.G. Drugs and Driving. Traffic Inj. Prev. 2004, 5, 241–253. [CrossRef]
- Orriols, L.; Gbaguidi, G.N.; Contrand, B.; Gadegbeku, B.; Lagarde, E. Trends in Benzodiazepine Anxiolytics and Z-Hypnotics Use among French Drivers Involved in Road Traffic Crashes from 2005 to 2015: A Responsibility Case-Control Study. *Inj. Epidemiol.* 2019, *6*, 32. [CrossRef]
- 21. Rudisill, T.M.; Zhu, M.; Kelley, G.A.; Pilkerton, C.; Rudisill, B.R. Medication Use and the Risk of Motor Vehicle Collisions among Licensed Drivers: A Systematic Review. *Accid. Anal. Prev.* **2016**, *96*, 255–270. [CrossRef]

- 22. Verster, J.C.; van de Loo, A.J.A.E.; Roth, T. Mirtazapine as Positive Control Drug in Studies Examining the Effects of Antidepressants on Driving Ability. *Eur. J. Pharmacol.* **2015**, 753, 252–256. [CrossRef]
- 23. Ramaekers, J.G. Antidepressants and Driver Impairment: Empirical Evidence from a Standard on-the-Road Test. J. Clin. Psychiatry 2003, 64, 20–29. [CrossRef] [PubMed]
- Schumacher, M.B.; Jongen, S.; Knoche, A.; Petzke, F.; Vuurman, E.F.; Vollrath, M.; Ramaekers, J.G. Effect of Chronic Opioid Therapy on Actual Driving Performance in Non-Cancer Pain Patients. *Psychopharmacology* 2017, 234, 989–999. [CrossRef] [PubMed]
- 25. Schulze, H.; Shumacher, M.; Urmeew, R.; Auerbach, K.; Alvarez, F.J.; Bernhoft, I.M.; De Gier, H.; Hagenzieker, M.; Houwing, S.; Knoche, A.; et al. Driving Under the Influence of Drugs, Alcohol and Medicines in Europe Findings from the DRUID Project. Lisbon: European Monitoring Centre for Drugs and Drug Addiction (EMCDDA). Available online: http://www.emcdda.europa. eu/system/files/publications/743/TDXA12006ENN_402402.pdf (accessed on 12 May 2022).
- 26. European Union (EU). European Council Directive 83/570/EEC of 26 October 1983 on the Approximation of Provisions Laid Down by Law, Regulation or Administrative Action Relating to Proprietary Medicinal Products. Available online: https: //eur-lex.europa.eu/legal-content/EN/TXT/?uri=celex%3A31983L0570 (accessed on 20 June 2020).
- Ravera, S.; Monteiro, S.P.; de Gier, J.J.; van der Linden, T.; Gómez-Talegón, T.; Alvarez, F.J. DRUID Project WP4 Partners A European Approach to Categorizing Medicines for Fitness to Drive: Outcomes of the DRUID Project. *Br. J. Clin. Pharmacol.* 2012, 74, 920–931. [CrossRef] [PubMed]
- Fierro, I.; Gómez-Talegón, T.; Alvarez, F.J. 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, 1056–1061. [CrossRef]
- 29. Monteiro, S.P. Driving-Impairing Medicines and Traffic Safety: Patients' Perspectives; Rijksuniversiteit Groningen: Groningen, The Netherlands, 2014.
- Ministry of Health. ROYAL DECREE 1345/2007, of October 11, 2007, Which Regulates the Procedure for Authorization, Registration and Conditions of Dispensing of Industrially Manufactured Medicines for Human Use. Available online: https: //www.boe.es/buscar/doc.php?id=BOE-A-2007-19249 (accessed on 20 June 2020).
- 31. de la Santé et des Solidarités, M.; de la Santé, D.G. 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; Ministère de la Santé et des Solidarité: Paris, France, 2005.
- 32. Alvarez, F.J.; Mercier-Guyon, C.; Verstraete, A.G. Prescribing and Dispensing Guidelines for Medicinal Drugs Affecting Driving Performance. In *Drugs, Driving and Traffic Safety*; BirkhäuserVerlag: Basel, Switzerland, 2009; pp. 121–134.
- Lamport, D.J.; Lawton, C.L.; Merat, N.; Jamson, H.; Myrissa, K.; Hofman, D.; Chadwick, H.K.; Quadt, F.; Wightman, J.D.; Dye, L. Concord Grape Juice, Cognitive Function, and Driving Performance: A 12-Wk, Placebo-Controlled, Randomized Crossover Trial in Mothers of Preteen Children. Am. J. Clin. Nutr. 2016, 103, 775–783. [CrossRef]
- 34. LaSala, G.S.; McKeever, R.G.; Patel, U.; Okaneku, J.; Vearrier, D.; Greenberg, M.I. Effect of Single-Dose Ginkgo Biloba and Panax Ginseng on Driving Performance. *Clin. Toxicol.* **2015**, *53*, 108–112. [CrossRef]
- 35. Irwin, C.; McCartney, D.; Grant, G.; Delang, N.; Bartrim, K.; Cox, G.R.; Desbrow, B. Effects of Different Sources of Low-Dose Caffeine on Mood/Arousal and Cognitive Performance. *Percept. Mot. Ski.* 2022, *129*, 1672–1690. [CrossRef]
- 36. Thomas, K.; Canedo, J.; Perry, P.J.; Doroudgar, S.; Lopes, I.; Chuang, H.M.; Bohnert, K. Effects of Valerian on Subjective Sedation, Field Sobriety Testing and Driving Simulator Performance. *Accid. Anal. Prev.* **2016**, *92*, 240–244. [CrossRef]
- Nagib, M.M.; Tadros, M.G.; Al-Khalek, H.A.A.; Rahmo, R.M.; Sabri, N.A.; Khalifa, A.E.; Masoud, S.I. Molecular Mechanisms of Neuroprotective Effect of Adjuvant Therapy with Phenytoin in Pentylenetetrazole-Induced Seizures: Impact on Sirt1/NRF2 Signaling Pathways. *Neurotoxicology* 2018, 68, 47–65. [CrossRef]
- Elsayed, A.A.; Menze, E.T.; Tadros, M.G.; Ibrahim, B.M.M.; Sabri, N.A.; Khalifa, A.E. Effects of Genistein on Pentylenetetrazole-Induced Behavioral and Neurochemical Deficits in Ovariectomized Rats. *Naunyn Schmiedebergs Arch. Pharmacol.* 2018, 391, 27–36. [CrossRef]
- Nagib, M.M.; Tadros, M.G.; Rahmo, R.M.; Sabri, N.A.; Khalifa, A.E.; Masoud, S.I. Ameliorative Effects of α-Tocopherol and/or Coenzyme Q10 on Phenytoin-Induced Cognitive Impairment in Rats: Role of VEGF and BDNF-TrkB-CREB Pathway. *Neurotox. Res.* 2019, 35, 451–462. [CrossRef]
- 40. Tadros, M.G.; Mohamed, M.R.; Youssef, A.M.; Sabry, G.M.; Sabry, N.A.; Khalifa, A.E. Proapoptotic and Prepulse Inhibition (PPI) Disrupting Effects of Hypericum Perforatum in Rats. *J. Ethnopharmacol.* **2009**, *122*, 561–566. [CrossRef]
- Tadros, M.G.; Mohamed, M.R.; Youssef, A.M.; Sabry, G.M.; Sabry, N.A.; Khalifa, A.E. Involvement of Serotoninergic 5-HT1A/2A, Alpha-Adrenergic and Dopaminergic D1 Receptors in St. John's Wort-Induced Prepulse Inhibition Deficit: A Possible Role of Hyperforin. *Behav. Brain Res.* 2009, 199, 334–339. [CrossRef]
- Gutiérrez-Abejón, E.; Herrera-Gómez, F.; Criado-Espegel, P.; Alvarez, F.J. Use of Driving-Impairing Medicines by a Spanish Population: A Population-Based Registry Study. *BMJ Open* 2017, 7, e017618. [CrossRef]
- Herrera-Gómez, F.; Gutiérrez-Abejón, E.; Criado-Espegel, P.; Álvarez, F.J. The Problem of Benzodiazepine Use and Its Extent in the Driver Population: A Population-Based Registry Study. *Front. Pharmacol.* 2018, 9, 408. [CrossRef]
- 44. Herrera-Gómez, F.; Gutiérrez-Abejón, E.; Álvarez, F.J. Antipsychotics in the General Population and the Driver Population: Comparisons from a Population-Based Registry Study. *Int. Clin. Psychopharmacol.* **2019**, *34*, 184–188. [CrossRef]

- 45. Herrera-Gómez, F.; Gutiérrez-Abejón, E.; Ayestarán, I.; Criado-Espegel, P.; Álvarez, F.J. The Trends in Opioid Use in Castile and Leon, Spain: A Population-Based Registry Analysis of Dispensations in 2015 to 2018. *J. Clin. Med.* **2019**, *8*, 2148. [CrossRef]
- 46. Gutiérrez-Abejón, E.; Herrera-Gómez, F.; Criado-Espegel, P.; Álvarez, F.J. Trends in Antidepressants Use in Spain between 2015 and 2018: Analyses from a Population-Based Registry Study with Reference to Driving. *Pharmaceuticals* **2020**, *13*, *61*. [CrossRef]
- 47. Gutiérrez-Abejón, E.; Criado-Espegel, P.; Herrera-Gómez, F.; Álvarez, F.J. Population-Based Registry Analysis of Antidiabetics Dispensations: Trend Use in Spain between 2015 and 2018 with Reference to Driving. *Pharmaceuticals* **2020**, *13*, 165. [CrossRef]
- 48. Gutiérrez-Abejón, E.; Herrera-Gómez, F.; Álvarez, F.J. Trends in the Use of Antihistamines with Reference to Drivers between 2015 and 2019: A Population-Based Registry Analysis. *Fundam. Clin. Pharmacol.* **2021**, *35*, 1168–1178. [CrossRef]
- Hernández-Rodríguez, M.Á.; Sempere-Verdú, E.; Vicens-Caldentey, C.; González-Rubio, F.; Miguel-García, F.; Palop-Larrea, V.; Orueta-Sánchez, R.; Esteban-Jiménez, Ó.; Sempere-Manuel, M.; Arroyo-Aniés, M.P.; et al. Evolution of Polypharmacy in a Spanish Population (2005–2015): A Database Study. *Pharmacoepidemiol. Drug Saf.* 2020, 29, 433–443. [CrossRef] [PubMed]
- Green, I.; Stow, D.; Matthews, F.E.; Hanratty, B. Changes over Time in the Health and Functioning of Older People Moving into Care Homes: Analysis of Data from the English Longitudinal Study of Ageing. *Age Ageing* 2017, *46*, 693–696. [CrossRef] [PubMed]
- 51. Topinková, E.; Baeyens, J.P.; Michel, J.-P.; Lang, P.-O. Evidence-Based Strategies for the Optimization of Pharmacotherapy in Older People. *Drugs Aging* **2012**, *29*, 477–494. [CrossRef] [PubMed]
- 52. Falkenstein, M.; Karthaus, M.; Brüne-Cohrs, U. Age-Related Diseases and Driving Safety. Geriatrics 2020, 5, 80. [CrossRef]
- 53. Anstey, K.J.; Wood, J.; Lord, S.; Walker, J.G. Cognitive, Sensory and Physical Factors Enabling Driving Safety in Older Adults. *Clin. Psychol. Rev* 2005, 25, 45–65. [CrossRef]
- 54. Cicchino, J.B.; McCartt, A.T. Trends in Older Driver Crash Involvement Rates and Survivability in the United States: An Update. *Accid. Anal. Prev.* **2014**, 72, 44–54. [CrossRef]
- 55. Ministry of the Interior. Royal Decree 818/2009, of May 8, 2009, Approving the General Regulations for Drivers. Available online: https://www.boe.es/buscar/doc.php?id=BOE-A-2009-9481 (accessed on 22 March 2023).
- 56. Wilhelmi, B.G.; Cohen, S.P. A Framework for "Driving under the Influence of Drugs" Policy for the Opioid Using Driver. *Pain Physician* **2012**, *15*, ES215-230. [CrossRef]
- 57. 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*, 81–87. [CrossRef]
- 58. Bernhoft, I. Results from Epidemiological Research—Prevalence, Risk and Characteristics of Impaired Drivers; European Union: Brussels, Belgium, 2011.
- 59. Hill, L.L.; Lauzon, V.L.; Winbrock, E.L.; Li, G.; Chihuri, S.; Lee, K.C. Depression, Antidepressants and Driving Safety. *Inj. Epidemiol.* **2017**, *4*, 10. [CrossRef]
- 60. Wickens, C.M.; Mann, R.E.; Stoduto, G.; Ialomiteanu, A.; Smart, R.G.; Rehm, J. The Impact of Probable Anxiety and Mood Disorder on Self-Reported Collisions: A Population Study. J. Affect. Disord. 2013, 145, 253–255. [CrossRef]
- 61. Orriols, L.; Queinec, R.; Philip, P.; Gadegbeku, B.; Delorme, B.; Moore, N.; Suissa, S.; Lagarde, E. CESIR Research Group Risk of Injurious Road Traffic Crash after Prescription of Antidepressants. *J. Clin. Psychiatry* **2012**, *73*, 1088–1094. [CrossRef]
- 62. Brunnauer, A.; Buschert, V.; Fric, M.; Distler, G.; Sander, K.; Segmiller, F.; Zwanzger, P.; Laux, G. Driving Performance and Psychomotor Function in Depressed Patients Treated with Agomelatine or Venlafaxine. *Pharmacopsychiatry* **2015**, *48*, 65–71. [CrossRef]
- 63. Ravera, S.; Ramaekers, J.G.; de Jong-van den Berg, L.T.W.; de Gier, J.J. Are Selective Serotonin Reuptake Inhibitors Safe for Drivers? What Is the Evidence? *Clin. Ther.* 2012, *34*, 1070–1083. [CrossRef]
- 64. Brunnauer, A.; Laux, G. Driving Under the Influence of Antidepressants: A Systematic Review and Update of the Evidence of Experimental and Controlled Clinical Studies. *Pharmacopsychiatry* **2017**, *50*, 173–181. [CrossRef]
- 65. Simó Miñana, J. Utilización de Medicamentos en España y en Europa. Atención Primaria 2012, 44, 335–347. [CrossRef]
- 66. Agencia Española del Medicamento y Productos Sanitarios. *Informe De Utilización De Medicamentos U/AN/V1/03092015: Utilización de Medicamentos Antidiabéticos En España Durante El Periodo 2000–2014;* Agencia Española del Medicamento y Productos Sanitarios: Madrid, Spain, 2015.
- 67. Diabetes Canada Clinical Practice Guidelines Expert Committee; Houlden, R.L.; Berard, L.; Lakoff, J.M.; Woo, V.; Yale, J.-F. Diabetes and Driving. *Can. J. Diabetes* **2018**, 42 (Suppl. S1), S150–S153. [CrossRef]
- Cox, D.J.; Singh, H.; Lorber, D.; Hermayer, K. Diabetes and Driving Safety: Science, Ethics, Legality and Practice. *Am. J. Med. Sci.* 2013, 345, 263–265. [CrossRef]
- 69. Laberge-Nadeau, C.; Dionne, G.; Ekoé, J.M.; Hamet, P.; Desjardins, D.; Messier, S.; Maag, U. Impact of Diabetes on Crash Risks of Truck-Permit Holders and Commercial Drivers. *Diabetes Care* 2000, 23, 612–617. [CrossRef]
- 70. Inkster, B.; Frier, B.M. Diabetes and Driving. *Diabetes Obes. Metab.* 2013, 15, 775–783. [CrossRef]
- Ma, S.; Zhang, J.; Zeng, X.; Wu, C.; Zhao, G.; Lv, C.; Sun, X. Type 2 Diabetes Can Undermine Driving Performance of Middle-Aged Male Drivers through Its Deterioration of Perceptual and Cognitive Functions. *Accid. Anal. Prev.* 2020, 134, 105334. [CrossRef]
- 72. UK Hypoglycaemia Study Group. Risk of Hypoglycaemia in Types 1 and 2 Diabetes: Effects of Treatment Modalities and Their Duration. *Diabetologia* 2007, *50*, 1140–1147. [CrossRef] [PubMed]

- Signorovitch, J.E.; Macaulay, D.; Diener, M.; Yan, Y.; Wu, E.Q.; Gruenberger, J.-B.; Frier, B.M. Hypoglycaemia and Accident Risk in People with Type 2 Diabetes Mellitus Treated with Non-Insulin Antidiabetes Drugs. *Diabetes Obes. Metab.* 2013, 15, 335–341. [CrossRef] [PubMed]
- 74. Stork, A.D.M.; van Haeften, T.W.; Veneman, T.F. Diabetes and Driving: Desired Data, Research Methods and Their Pitfalls, Current Knowledge, and Future Research. *Diabetes Care* 2006, 29, 1942–1949. [CrossRef] [PubMed]
- 75. Kilpatrick, E.S.; Rigby, A.S.; Warren, R.E.; Atkin, S.L. Implications of New European Union Driving Regulations on Patients with Type 1 Diabetes Who Participated in the Diabetes Control and Complications Trial. *Diabet. Med.* **2013**, *30*, 616–619. [CrossRef]
- 76. American Diabetes Association; Lorber, D.; Anderson, J.; Arent, S.; Daniel, J.; Frier, B.M.; Greene, M.A.; Griffin, J.W.; Gross, G.; Hathaway, K.; et al. Diabetes and Driving. *Diabetes Care* **2012**, *35* (Suppl. S1), S81–S86. [CrossRef]
- 77. Legrand, S.-A.; Boets, S.; Meesmann, U.; Verstraete, A.G. Medicines and Driving: Evaluation of Training and Software Support for Patient Counselling by Pharmacists. *Int. J. Clin. Pharm.* **2012**, *34*, 633–643. [CrossRef]
- 78. Ravera, S.; van Rein, N.; de Gier, J.J.; de Jong-van den Berg, L.T.W. Road Traffic Accidents and Psychotropic Medication Use in The Netherlands: A Case-Control Study. *Br. J. Clin. Pharmacol.* **2011**, *72*, 505–513. [CrossRef]
- 79. Jomaa, I.; Odisho, M.; Cheung, J.M.Y.; Wong, K.; Ellis, J.G.; Smyth, T.; Saini, B. Pharmacists' Perceptions and Communication of Risk for Alertness Impairing Medications. *Res. Soc. Adm. Pharm.* **2018**, *14*, 31–45. [CrossRef]
- 80. Ramaekers, J.G.; Berghaus, G.; van Laar, M.; Drummer, O.H. Dose Related Risk of Motor Vehicle Crashes after Cannabis Use. *Drug Alcohol Depend.* **2004**, *73*, 109–119. [CrossRef]
- 81. von Elm, E.; Altman, D.G.; Egger, M.; Pocock, S.J.; Gøtzsche, P.C.; Vandenbroucke, J.P. STROBE Initiative The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) Statement: Guidelines for Reporting Observational Studies. *J. Clin. Epidemiol.* **2008**, *61*, 344–349. [CrossRef]
- 82. Benchimol, E.I.; Smeeth, L.; Guttmann, A.; Harron, K.; Moher, D.; Petersen, I.; Sørensen, H.T.; von Elm, E.; Langan, S.M.; RECORD Working Committee. The REporting of Studies Conducted Using Observational Routinely-Collected Health Data (RECORD) Statement. *PLoS Med.* **2015**, *12*, e1001885. [CrossRef]
- WHO Collaborating Centre for Drug Statistics Methodology ATC/DDD Index. Available online: https://www.whocc.no/atc_ddd_index/ (accessed on 12 August 2020).
- 84. Ramaekers, J.G. Drugs and Driving Research in Medicinal Drug Development. Trends Pharmacol. Sci. 2017, 38, 319–321. [CrossRef]
- 85. Alonso, F.; Esteban, C.; Montoro, L.; Tortosa, F. Psychotropic Drugs and Driving: Prevalence and Types. *Ann. Gen. Psychiatry* **2014**, *13*, 14. [CrossRef]
- 86. Emich, B.; van Dijk, L.; Monteiro, S.P.; de Gier, J.J. A Study Comparing the Effectiveness of Three Warning Labels on the Package of Driving-Impairing Medicines. *Int. J. Clin. Pharm.* **2014**, *36*, 1152–1159. [CrossRef]

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