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Title: Determinants of prescribing potentially inappropriate medications in a nationwide cohort of community dwellers with dementia receiving a comprehensive geriatric assessment

Running title: Inappropriate prescribing in dementia utilizing the interRAI dataset

Sharmin S Bala^{1*}, Hamish A Jamieson², Prasad S Nishtala³

¹Department of Preventive and Social Medicine, University of Otago, Dunedin, New Zealand

²Department of Medicine, University of Otago, Christchurch, New Zealand

³School of Pharmacy, University of Otago, Dunedin, New Zealand

***Corresponding author:**

Dr Sharmin S Bala, Department of Preventive and Social Medicine,
University of Otago, New Zealand.

Phone: +64-220893600

Email: sharmin.bala@postgrad.otago.ac.nz

ORCID: 0000-0003-1126-8291

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1 **Abstract:**

2 **Objective:** To identify the prevalence and predictors of prescribing potentially inappropriate
3 medications (PIMs) in a nationwide cohort of community dwellers with dementia requiring
4 complex care needs.

5 **Methods:** A cross-matched data of the International Resident Assessment Instrument-Home
6 Care (9.1) (interRAI-HC) ~~and with~~ prescribing data obtained from the Pharmaceutical Claims
7 Data Mart (Pharms) extract files for older adults (≥ 65 years) requiring complex care needs was
8 utilized for this study. The 2015 Beers criteria were applied to identify the prevalence of PIMs
9 in older adults with dementia. Sociodemographic and clinical predictors of PIMs were analysed
10 using a logistic regression model.

11 **Results:** The study population consisted of 16,568 individuals who had their first interRAI
12 assessment from 1st January 2015 to 31st December 2015. The estimated prevalence of
13 dementia was 13.2% (2,190/16,568). 66.9% (1,465/2,190) of the older adults diagnosed with
14 dementia were prescribed PIMs, of which anticholinergic medications constituted 59.6%
15 (873/1,465). Males and individuals who were prescribed a greater number of medications were
16 more likely to be prescribed PIMs. Individuals over 85 years of age, Māori ethnic group of
17 individuals, older adults who were being supervised with respect to their activities of daily
18 living, and individuals who reported good or excellent self reported health, had a lesser
19 likelihood of being prescribed PIMs.

20 **Conclusion:** We found that PIMs are prescribed frequently in older adults with dementia.
21 Comprehensive geriatric assessments can serve as a potential tool to decrease the occurrence
22 of PIMs in vulnerable groups with poor functional and cognitive status.

23

24 **Key words:** Dementia, potentially inappropriate medications, interRAI, prescribing in older
25 adults

26

27 **Key Points:**

28 The 2015 Beers criteria and interRAI assessments were used to determine the prevalence and
29 predictors of prescribing potentially inappropriate medications (PIMs) in older adults
30 diagnosed with dementia, and having complex care needs.

1 There was a high prevalence of PIMs in older adults with dementia. Anticholinergic
2 medications comprised the most commonly prescribed PIMs. Sociodemographic factors
3 including younger age, male gender, and New Zealand European ethnicity; and clinical factors
4 like the increased number of medications prescribed, functional status, and self-reported health
5 influenced the prescription of PIMs in this vulnerable population.

6 A due consideration of these attributes while prescribing may aid in reducing the prescribing
7 of PIMs in individuals with dementia.

8
9

10 **Introduction:**

11 Dementia is one of the principal syndromes linked with disability and dependence among older
12 adults and is a major challenge to individuals, communities, and societies globally.¹ The global
13 incidence of dementia is expected to rise to 81 million by 2040, primarily due to the progressive
14 nature of the disorder, which involves worsening of neurocognitive impairment and loss of
15 basic functions in daily life.²⁻⁴ In 2016, the estimated prevalence of dementia in New Zealand
16 (NZ) was more than 62,000, which is predicted to increase to 170,000 in 2050.⁵ One in every
17 four 'international Resident Assessment Instrument- Home Care' (interRAI-HC) evaluated
18 individuals in 2016-17 in NZ were diagnosed with dementia, of which 35% needed extensive
19 assistance or were completely dependent, and 30% ~~had showed~~ daily episodes of disturbing
20 behaviours, like wandering or being abusive.⁶ The interRAI was developed by an international
21 collaboration of experts and shows good inter-rater reliability. The interRAI assessors who
22 conduct the assessments are trained in a quality assurance programme conducted by the
23 Ministry of Health, NZ. A national competency framework supports and promotes the quality
24 assurance programs for the interRAI assessments.⁸ The interRAI-HC tool gathers information
25 on physical, mental, social, and cognitive domains of the health of residents/clients living in
26 home care settings.⁸

27

28 Beers et al have defined potentially inappropriate medications (PIMs) as the prescription of
29 medications, where the risks outweigh clinical benefits, particularly when there is safer or more
30 effective alternative therapy for the same condition.⁹ A study conducted in rural community-
31 dwelling older adults in the United States of America (USA), utilizing the Beers criteria ~~has~~

1 found that half of the population of older adults utilized over-the-counter and prescribed
2 inappropriate medications.¹⁰ Prescribing medications for older adults with dementia is
3 challenging because of the risks associated with cognitive decline, behavioural and
4 psychological disturbances, prescription of multiple medications, and their associated
5 costs.^{12,13} Older adults with dementia also experience a greater sensitivity to the adverse effects
6 of medications acting on the central nervous system.^{14,15} Long-term utilization of medications
7 in older individuals with impaired physical and cognitive function has been associated with
8 increased risks of hospitalization and mortality.¹⁶

9 The 2015 Beers Criteria is often used as a tool for assessing the appropriateness of prescribing
10 medications in the geriatric population.¹⁷ A systematic review among older adults with
11 cognitive impairment and dementia reported a varied prevalence of prescription of PIMs of
12 10.2%–56.4% across Europe, Australia, and the United States; and the Beers criteria were
13 applied for assessing PIMs in ~~the~~ majority of the studies.¹⁸ Another recent literature review
14 reported that among the five studies conducted in ambulatory home-dwelling patients
15 diagnosed with dementia that used the Beers criteria, the prevalence of PIMs use ranged from
16 16.2% to 33%, and PIMs were found to be associated with the gender, ethnicity, number of
17 medications prescribed, and varied medical conditions.¹⁹ An understanding about the
18 prevalence and determinants of PIMs can help prevent adverse effects, and improve the quality
19 of prescribing in this vulnerable age-group. To our knowledge, this is the first study conducted
20 to evaluate the predictors of prescribing PIMs in older adults with dementia receiving
21 comprehensive geriatric risk assessments. These assessments capture a suite of
22 sociodemographic and clinical variables that have not been previously investigated. The study
23 aims to identify the prevalence and associations of prescribing PIMs in a nationwide cohort of
24 community dwellers with dementia. The overarching objective of the study is to identify the
25 factors associated with inappropriate prescribing in older adults diagnosed with dementia,
26 identified by the 2015 version of Beers criteria, by multinomial regression, utilizing the 2015
27 interRAI dataset.

28

29

30 **Materials and Methods:**

31 The Human Ethics Committee, University of Otago, NZ has approved the proceedings of the
32 study (ethical approval number 15/CEN/45/AM02).

1 **Data source:** The international Resident Assessment Instrument-Home Care (9.1) (interRAI-
2 HC) dataset is a comprehensive geriatric risk assessment for community-dwelling older adults
3 with complex care needs needing publicly funded long-term community services or aged
4 residential care in NZ.²⁰ In NZ, a standardized interRAI-HC has been implemented for
5 conducting all community care assessments in older adults requiring publically funded long-
6 term community services or aged residential care.²¹ Assessors in District Health Boards
7 throughout NZ utilize this standardised interRAI-HC assessment to assist in defining the level
8 of support required for the geriatric population.²⁰ Older adults are referred by a health
9 practitioner to have their health requirements assessed by the trained interRAI-HC assessors.
10 Assessors visit the individuals at their residence to develop individualized treatment plans
11 according to a standardized protocol.²² Once the participating individuals consent to the de-
12 identified interRAI-HC data being used for planning and research purposes, the information is
13 then collated into the electronic interRAI-HC database, maintained by New Zealand's
14 Technical Advisory Services (TAS) to provide facts and figures at the provider, regional and
15 national level.²² The interRAI-HC assessments provide information on 20 domains, including
16 social demographics, medical ailments, frailty, cognitive and physical function.²¹ The interRAI
17 database is linked to various NZ Ministry of Health nationwide collections, including
18 prescription use (the Pharms database), hospital discharges (National Minimum Dataset),
19 laboratory collections and mortality data, among others.²¹ The information on all the
20 prescription claims funded by PHARMAC (Pharmaceutical management agency) was sourced
21 from the Pharmaceutical Claims Data Mart (Pharms) extract files 2015, prepared by
22 community pharmacists, and includes the sociodemographics and medication details of the
23 individuals.²⁴ The cross-matched 2015 interRAI-Pharms dataset was utilized for the present
24 study.

25

26 **Study population:** The eligible study population comprised community dwellers (≥ 65 years
27 of age) in NZ who have undergone the first interRAI comprehensive geriatric risk assessment
28 in 2015. The present study had a retrospective design, tracking the prevalence of PIMs and
29 dementia, and delineating the factors associated with prescription of PIMs in dementia in older
30 individuals. The personal information of the individuals is de-identified in the dataset.

31

32 **PIMs exposure:** PIMs were defined according to the list of medications to be avoided in
33 individuals diagnosed with dementia as per the 2015 Beers criteria²⁵ developed by the
34 American Geriatrics Society; which comprised the prescription of psychotropics, including

1 antipsychotics, medications with anticholinergic properties, benzodiazepines,
2 nonbenzodiazepines, benzodiazepine receptor agonist hypnotics, and H₂ receptor antagonists
3 **(Table 1)**. We have excluded medications listed under the 2015 Beers criteria which were not
4 available or not subsidized in NZ **(Appendix 1)**.

5
6 **Diagnosis of dementia:** Dementia was diagnosed by the Minimum Data Set (MDS) Cognitive
7 Performance Scale (CPS).²⁶ The CPS is a valid and reliable seven-point hierarchical scale
8 derived from the MDS that rates impairment from intact memory to very severe memory loss.²⁷
9 The CPS collates data on comatose status, short-term memory, cognitive skills for daily
10 decision-making, being understood by others, and self-performance in eating, with scores
11 ranging from 0 (intact memory) to 6 (very severe memory impairment). The CPS has been
12 shown to be highly correlated with the Mini-Mental State Examination in a number of
13 validation studies.

14
15 **Covariates:** The individuals were grouped according to age: 65-74 years, 75-84 years, 85-94
16 years, and 95 years and above. Based on the literature review of known predictors of
17 inappropriate prescribing in older adults with and without dementia, explanatory variables that
18 were tested for their influence on prescribing patterns were:

19 **1. Sociodemographics:** Age³⁰, gender³⁰ ethnicities³¹, marital status³⁰, living
20 arrangements^{32,33}, alcohol intake³⁴, and smoking history³⁵.

21 **2. Clinical:** Activities of daily living^{30,36}, self-reported health³⁰, hospitalization^{32,36}, and
22 number of medications.^{37,38}

23
24 **Statistical analysis:** The potential impact of different explanatory variables on the outcome
25 variable (PIMs) was analysed using logistic regression models.

26 Individuals with a diagnosis of dementia were flagged as a binary variable; those with the
27 diseased condition were coded as 1, the coding for no dementia was 0. Descriptive analysis
28 was conducted utilizing the IBM SPSS version 24. Logistic regression analysis was performed
29 using StataCorp® Release 14.2. We utilized the ‘STrengthening the Reporting of Observational
30 studies in Epidemiology’ (STROBE) guidelines (www.strobe-statement.org) as the research
31 reporting guiding principle³⁹ **(Appendix 2)**

32
33

1 **Results:**

2 The current analysis is based on data collected from 16,568 Home-Care assessments from 1st
3 January 2015 to 31st December 2015, who have received at least one prescription medication
4 funded by PHARMAC. Our observations suggest that the female population comprised 60.1%
5 (9,964). The mean (SD) age of the population was 82.35 (7.6) years. Individuals of all
6 ethnicities were included in the study. NZ Europeans and Māori were studied particularly, as
7 they represent the largest ethnicities in NZ.³¹ 13.2% (2,190) of the study population was
8 diagnosed with dementia. Dementia was marginally more prevalent in males (14.6%)
9 compared to females ($p=0.001$), and among the Māori ethnicity (16.6%). The
10 sociodemographic characteristics of the study population are depicted in **Table 2**, which also
11 displays the associations of PIMs with the corresponding 95% confidence interval (CI)
12 ($p<0.05$), after adjustment for confounders, in individuals diagnosed with dementia. Overall,
13 we observed that 66.9% (1,465/2,190) of the older adults diagnosed with dementia were
14 prescribed PIMs, of which 59.6% (873/1,465) constituted anticholinergic inappropriate
15 medications. Overall, 39.9% (873/2,190) of the individuals diagnosed with dementia were
16 prescribed anticholinergic PIMs (**Figure 1**).

17 **Sociodemographic predictors:** Individuals over 85 years of age were less likely to be
18 prescribed PIMs, compared to individuals aged 65-74 years (aOR=0.64, CI=0.53, 0.77 for
19 individuals aged 85-94 years, aOR=0.53, CI=0.38, 0.73 for individuals over 95 years). Males
20 were more likely to be prescribed PIMs (aOR=1.24, CI=1.09, 1.41) compared to females. The
21 Māori ethnic group (aOR=0.59, CI=0.47, 0.76) and the other ethnicity groups (aOR=0.68,
22 CI=0.54, 0.87) were less likely to be prescribed PIMs, compared to the NZ Europeans.

23 **Clinical factors associated with PIMs**

24 Older adults who were prescribed a greater number of medications were more likely to be
25 prescribed PIMs (aOR=1.15, CI=1.14, 1.16), compared to those prescribed a single medication.
26 With respect to the activities of daily living, the older adults who were being supervised
27 (aOR=0.83, CI=0.69, 0.99) were less likely to be prescribed PIMs, compared to individuals
28 who were independent in their self-performance and capacity. Older adults who reported
29 excellent (aOR=0.62, CI=0.43, 0.89) and good self-health (aOR=0.65, CI=0.49, 0.85) had a
30 lesser likelihood to be prescribed PIMs, compared to those who reported poor self-health.

31

32

1 **Discussion:**

2 We reported the prevalence of prescription of PIMS and identified the sociodemographic and
3 clinical variables associated with the prescription of PIMS among community-dwelling older
4 adults with dementia.

5

6 Several studies investigating PIMS in older individuals have been carried out in NZ that made
7 use of the Beers criteria, and have focused on community-dwelling or hospitalized older
8 adults.^{31,35,40} This study appears to be the first to apply the 2015 Beers criteria to examine the
9 associations of inappropriate prescribing exclusively in community-dwelling older individuals
10 with dementia in NZ, who have received a comprehensive geriatric risk assessment.

11

12 The NZ interRAI annual report 2016-17⁶ noted a 1.2 times higher prevalence of dementia in
13 males, compared to females, findings consistent with those in our study. Likewise, in the
14 present study, male individuals diagnosed with dementia were more prone to be prescribed
15 PIMS, compared to females, similar to the findings of a recent study in Finland⁴¹ which reported
16 the male gender as a risk factor for initiation of PIMS in community-dwelling older adults with
17 Alzheimer's disease; and the research conducted in Korea which analysed the trends in
18 prescribing of atypical antipsychotics in geriatric patients with dementia.⁴² Numerous studies
19 have reported a higher rate of inappropriate medication use in older women than in men of the
20 same age-group⁴³⁻⁴⁵, although the clinical relevance of this association remains uncertain.⁴⁶

21

22 Our study observed a high prevalence of dementia in the Māori group, comparable to that
23 observed in a study conducted in NZ to assess an indigenous approach for the diagnosis and
24 management of dementia⁴⁷, and to the findings of the NZ interRAI annual report 2016-17.⁶ It
25 has been observed that the Māori population over the age of 50 have worse health outcomes
26 and a greater burden of chronic ailments than non-Māori of the same age-group.⁴⁸ The
27 prevalence of PIMS (68.4%) in NZ Europeans with dementia exceeded the occurrence in all
28 other ethnic groups of individuals, identical to other PIMS prevalence studies in the geriatric
29 age group conducted in NZ.^{31,40} This could be attributed to NZ Europeans being the
30 predominant ethnic group in New Zealand, comprising 71.2% of the country's inhabitants.⁴⁹

31

32 A higher proportion (72.3%) of relatively younger group (65-75 years) of individuals with
33 dementia were prescribed PIMS, which reflects the findings of the study performed by Hyttinen
34 et al to evaluate PIMS prevalence in community-dwelling older adults with and without

1 Alzheimer's Disease⁴¹. These findings could potentially be attributed to prescriber awareness
2 concerning the prescription of PIMs in individuals of the older age group.⁵⁰

3
4 There was a high prevalence of PIMs (66.9%) in older adults diagnosed with dementia in our
5 study, comparable to the prevalence (62%) in a study conducted in community-dwelling older
6 adults in the USA.⁵¹ This is a significant finding of our study. Similarly, researchers in
7 Australia⁵² reported a prescription of at least one PIM as 56.4% among individuals with
8 dementia living in residential aged care facilities. A study of six residential care homes in
9 England⁴⁶ observed the prevalence of at least 1 PIM prescribed in 46.2% and 40.9% of the
10 older individuals with dementia, utilizing the STOPP criteria, reviewed at two time-points, 16
11 weeks apart. A direct international comparison of the prevalence of PIM prescriptions with our
12 findings is challenging because of the differences between the PIMs lists used, and the
13 population under study. Holmes et al have developed a tool for assessment of appropriate
14 medication prescribing in advanced dementia, in which the primary goal is palliation of
15 symptoms.⁵³ There is a similar need to arrive at a global consensus through research on
16 appropriate prescribing in older adults presenting with different stages of dementia.

17
18 One of the most significant findings of our research suggest that 59.6% of the PIMs prescribed
19 belonged to the anticholinergic class of medications; and 39.9% of the population under study
20 were prescribed anticholinergic medications which were termed inappropriate to prescribe in
21 older individuals diagnosed with dementia, according to the 2015 Beers criteria.²⁵ Bhattacharya
22 et al⁵⁴ reported a prevalence of 43% of anticholinergic medication prescription among elderly
23 outpatients with dementia. A study by Somers et al⁵⁵ reported a high anticholinergic burden of
24 PIMs in residential aged care facilities in Melbourne. A study by Cross et al⁵⁶ in Australia also
25 reported a clinically significant anticholinergic burden in older adults attending Memory
26 Clinics. Anticholinergic agents are specifically associated with negative outcomes in older
27 adults diagnosed with dementia, such as risk of falls, delirium, worsening of cognitive function,
28 and increased mortality.^{54,57} The anticholinergic agents are notorious for their peripheral side
29 effects, which include dry mouth, constipation, urinary retention, and bowel obstruction; and
30 the central side effects such as impaired concentration, confusion, attention deficit, and
31 impairment of memory.⁵⁴ PIMs with anticholinergic properties may also inhibit the potential
32 benefits of cholinesterase inhibitors, which is the main pharmacological class, currently
33 approved for the management of Alzheimer's disease.⁵⁸ Several researchers have attested that

1 anticholinergics may be associated with an increased risk for the development of sustained
2 cognitive deficits, which can range from mild cognitive impairment to dementia.⁵⁹⁻⁶¹

3
4 The results of the logistic regression analysis showed that the likelihood of PIMs increased
5 with the number of medications prescribed, which is akin to the findings of the research in
6 older people with dementia in care homes in the United Kingdom (UK),⁴⁶ the study by
7 Wucherer et al in community-dwelling primary care patients screened positive for dementia,⁶²
8 and the research conducted in Sweden using the EU(7)-PIM list to evaluate the prevalence of
9 PIMs in older people with cognitive impairment.⁵⁷ A study performed in UK utilizing the
10 primary care database of anonymised electronic health records from general practice
11 witnessed that patients over 65 years of age diagnosed with dementia, and taking multiple
12 medications, were more likely to be prescribed antipsychotics.⁶³ A higher number of
13 medications being prescribed may indicate multiple comorbidities. Drug interactions and non-
14 adherence are other risk factors that may have adverse consequences among older adults with
15 dementia, which are linked to a high number of prescribed medications.⁵⁷

16
17 In our study, individuals who reported poor self-health had an increased likelihood of
18 developing PIMs, identical to the results of a study conducted in the USA to assess potentially
19 inappropriate anticholinergic medication use in home-dwelling older adults with dementia.⁵⁰
20 The Bronx Aging Study⁶⁴ revealed that patients with poor or fair ratings of self-perceived
21 health utilized more prescription medications. Scores on self-perceived health status may be an
22 appropriate measure of the syndrome. This is supported by the conclusion that subjects
23 reporting fair and poor ratings on self-perceived health have increased numbers of physician
24 visits.⁶⁴

25 Contrary to the analysis of studies conducted in individuals diagnosed with dementia in
26 Sweden⁶⁵ and in eight European countries⁶⁶, a striking result to emerge from our data is that
27 the PIMs prevalence was higher in individuals who were functionally independent with respect
28 to ADL.

29
30 The study found increased use of anticholinergic medicines in dementia. Several studies have
31 shown that prescription of anticholinergic medicines can adversely impact cognition, physical
32 function, and can also increase the risk of mortality. Future research will aim at utilizing the
33 interRAI assessments for reducing anticholinergic medicines, and whether this leads to

1 improved cognitive outcomes. Further research will be undertaken to develop safer alternatives
2 to anticholinergic medications in this vulnerable group.

3
4
5 **Strength of the study:** The strength of this study includes the use of a national comprehensive
6 geriatric assessment, interRAI tool to record social attributes, clinical diagnosis and medication
7 use information in a substantial number of subjects, and the inclusion of the geriatric population
8 with dementia. Selection bias is mitigated by the wide prescription coverage in this population.
9 Standardized interRAI HC assessments conducted by trained healthcare personnel facilitates
10 the provision of valid clinical, social and functional data for research purposes. Various
11 predictors incorporated in the multivariate regression model, such as living arrangements,
12 activities of daily living, self-reported health are seldom seen in studies using administrative
13 claims data; hence, this study provides a unique perspective to the determinants of prescription
14 of PIMs in older adults with dementia. An additional strength is the application of the updated
15 2015 Beers criteria.

16
17 **Limitations:** The prevalence and associations of PIMs with respect to the individual
18 psychotropic medications were not studied. It was not possible to delineate the subtypes of
19 dementia. The retrospective analysis may not have been as competent as a prospective research
20 in outlining the findings of the study. The geriatric risk assessments are conducted in older
21 individuals living in the community specifically requiring complex care needs, which is
22 different from surveyed populations of older adults living in the community. Hence, the
23 findings of this study might not be applicable to community-dwelling older adults in various
24 countries because of variances in the population, health systems, prescribing guidelines, and
25 the cost of medications, as all these factors influence prescribing patterns; however, country-
26 specific guidelines can be developed using this information. The study design is cross-
27 sectional, hence only the associations of prescribing PIMs have been highlighted, and the
28 causality cannot be established.

29
30 **Conclusion:** In the present study, we observed that the majority of the individuals diagnosed
31 with dementia were prescribed PIMs, indicating that the quality of prescribing needs to be
32 improved.³⁷ Furthermore, important sociodemographic predictors like male gender, European
33 ethnicity, relatively younger aged individuals, and clinical predictors like the prescription of
34 anticholinergic medications, a higher number of medications prescribed, poor self-health, and

1 functionally independent individuals were identified as risk factors for prescribing PIMs in
2 older adults diagnosed with dementia. Reviewing the modifiable predictors of prescribing
3 PIMs could significantly reduce the prevalence of inappropriate prescribing in this vulnerable
4 population.

5
6 **Statement of contributions of authors:** Dr Prasad Nishtala designed the study; Dr Sharmin
7 Bala performed the research, Dr Sharmin Bala analysed the data; Dr Prasad Nishtala and Dr
8 Hamish Jamieson contributed new methods and models; Dr Sharmin Bala wrote the paper. All
9 authors contributed to data interpretation, critically commented on manuscript for intellectual
10 content, and approved the final manuscript.

11
12 **Statement of human rights Ethical approval:** All procedures performed in studies involving
13 human participants were in accordance with the ethical standards of the institutional and/or
14 national research committee and with the 1964 Helsinki declaration and its later amendments
15 or comparable ethical standards.

16
17 **Disclosure of potential conflicts of interest:** The authors declare no conflict of interest.

18
19
20 **Research involving Human Participants:** For this type of study formal consent is not
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22
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24
25
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1 **Table 1: List of medications to be avoided in individuals diagnosed with dementia**

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Anticholinergics		Antipsychotics	H ₂ -receptor antagonists	Benzodiazapines
Chlorpheniramine	Homatropine [†] (excludes ophthalmic)	Haloperidol	Cimetidine	Alprazolam
Orphenadrine		Trifluoperazine	Ranitidine	Lorazepam
Diphenhydramine (oral)	Propantheline	Fluphenazine	Famotidine	Oxazepam
Benzotropine	Chlorpromazine	Chlorpromazine		Temazepam
Amitriptyline	Clozapine	Thioridazine		Triazolam
Clomipramine	Olanzapine	Aripiprazole		Clonazepam
Doxepin (>6 mg)	Thioridazine	Quetiapine		Diazepam
Imipramine	Trifluoperazine	Ziprasidone		Flurazepam
Nortriptyline	Oxybutynin	Risperidone		Meprobamate
Paroxetine	Solifenacin	Clozapine		
Trimipramine	Tolterodine	Olanzapine		
Atropine [†] (excludes ophthalmic)	Disopyramide			
Promethazine	Prochlorperazine			

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3 ***†=(excludes ophthalmic)**

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	Total		PIMs		
	N	(%)	aOR	95% CI	p-value
<i>Age (years)</i>					
65-74	3,048	(18.4)	1†	1	
75-84	6,776	(40.9)	0.84	(0.69, 1.01)	0.058
85-94	6,192	(37.4)	0.64	(0.53, 0.77)	0.000
95+	552	(3.3)	0.53	(0.38, 0.73)	0.000
<i>Sex ‡</i>					
Female	9,964	(60.1)	1†	1†	
Male	6,603	(39.9)	1.24	(1.09, 1.41)	0.001
<i>Ethnicity</i>					
European	14,639	(88.4)	1†	1†	
Māori	957	(5.8)	0.59	(0.47, 0.76)	0.000
Other	972	(5.9)	0.68	(0.54, 0.87)	0.002
<i>Marital status</i>					
Married	6,607	(39.9)	1†	1†	
Other	9,961	(60.1)	1.01	(0.80, 1.28)	0.916
<i>Alcohol</i>					
No	13,225	(79.8)	1†	1†	
Yes	3,343	(20.2)	1.02	(0.88, 1.19)	0.760
<i>Smoking</i>					
No	15,653	(94.5)	1†	1†	
Yes	915	(5.5)	0.85	(0.66, 1.09)	0.209
<i>Living arrangements</i>					
Alone	8,019	(48.4)	1†	1†	
Spouse only	5,447	(32.9)	1.14	(0.88, 1.50)	0.327
Other	1,292	(7.8)	0.79	(0.63, 0.99)	0.050
With child §	1,810	(10.9)	0.86	(0.71, 1.03)	0.119
<i>Activities of daily living ¶</i>					
Independent	9,985	(60.3)	1†	1†	
Supervision	2,143	(12.9)	0.83	(0.69, 0.99)	0.035
Limited	1,782	(10.8)	0.94	(0.76, 1.18)	0.642
Extensive	1,046	(6.3)	0.77	(0.60, 1.00)	0.053
Maximal	730	(4.4)	0.74	(0.53, 1.02)	0.066
Dependent+	880	(5.3)	0.84	(0.60, 1.16)	0.288
<i>Self-reported health</i>					
Poor	1,925	(11.6)	1†	1†	
Excellent	522	(3.2)	0.62	(0.43, 0.89)	0.010
Good	6,806	(41.1)	0.65	(0.49, 0.85)	0.002
Fair	5,695	(34.4)	0.80	(0.60, 1.06)	0.124
Couldn't respond	1,620	(9.8)	0.82	(0.59, 1.13)	0.229
<i>Hospitalisation</i>					
No hospitalisation (in last 90 days)	8,602	(51.9)	1†	1†	
Other	7,966	(48.1)	1.11	(0.97, 1.27)	0.129
<i>Dementia</i>					

No	14,378	(86.8)	1†	1†	
Yes	2,190	(13.2)	1.17	(0.99, 1.37)	0.057
No of meds			1.15	(1.14, 1.16)	0.000

1 †= Reference value, ‡= 1 missing, §= not spouse / partner, ¶=2 missing.
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