Psychotropic Drug Utilisation in Older People in New Zealand from 2005 to 2013

Henry Ndukwe¹, June Tordoff¹, Ting Wang², Prasad Nishtala¹

¹School of Pharmacy
²Department of Mathematics and Statistics,

University of Otago, Dunedin, New Zealand



Outline

- Background
- Aims
- Method
- Results
- Conclusions
- Acknowledgements

Background

 International studies on psychotropic drug utilisation have shown high consumption levels (≥65 years)¹

 Long-term use has been associated with an increased risk of adverse events²

 Limited epidemiological information on psychotropic drug use in older people in New Zealand³

- 1. Nishtala PS, McLachlan AJ, Bell JS, Chen TF. Am J Geriatr Psych. 2008 Aug;16(8):621-32.
- 2. Gnjidic D, Bell JS, Hilmer SN. J Am Geriatr Soc. 2013 Sep;61(9):1640-1.
- 3. Ndukwe HC, Tordoff JM, Wang T, Nishtala PS, *Drugs Aging*, 2014; 31(10):755-68.

Aims

To describe and characterise national utilisation trend of psychotropic drugs used in older people New Zealand from 2005 to 2013

Method

 Repeated cross-sectional analysis of populationlevel dispensing data

 De-identified dispensing data extraction from Pharmaceutical collections by a unique identifier

Categorised using WHO-DDD classification system

 Defined daily dose (DDD) per 1000 older people per day (TOPD)

Defined Daily Dose

For example, Citalopram 20mg; WHO assigned (20 mg)

DDD = Strength (20mg/tablet) * Quantity Dispensed (1 tablet/day)
WHO-DDD (20mg)

DDD per year for a hypothetical weighted DDD sum of 25,000DDDs normalised by population of say 500,000 will give a standard weighted utilisation:

DDD/1000 older people /day = <u>DDD per year (0.05) * 1000</u> 365

Results

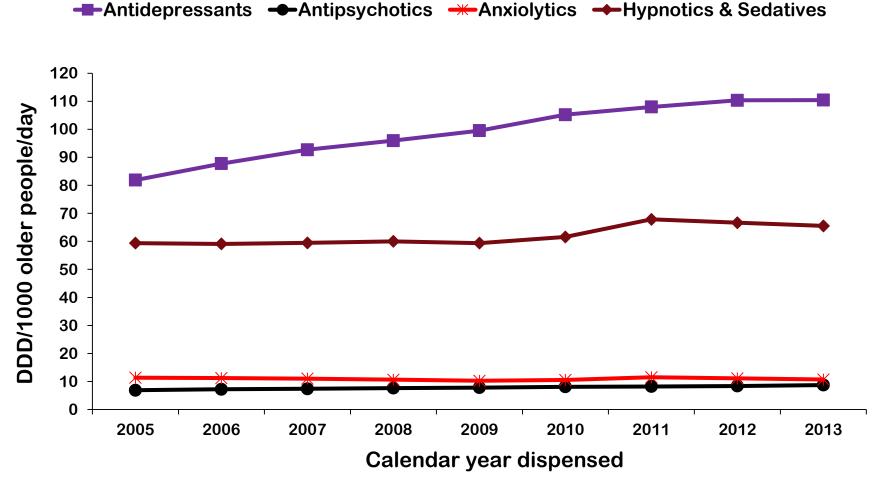


Fig 1. Psychotropic drug utilisation, yearly, in older people between 2005 to 2013. DDD defined daily dose

Published article for a part of this study can be found in Drugs Aging. 2014 Oct;31(10):755-68. doi: 10.1007/s40266-014-0205-1.

Research in Pharmacoepidemiology (RiPE) Group @ National School of Pharmacy, University of Otago

Table 1: Psychotropic drug utilisation (in DDD/TOPD) compared by therapeutic class and subclass between 2005 and 2013 calendar years

Therapeutic class	Therapeutic subclass	ATC CODE	2005 DDD/TOPD	2013 DDD/TOPD
Antidepressants	SSRI	N06AB	53.9	73.4
	TCA	N06AA	23.2	22.2
	TeCA	N06AX/AA	0.2	3.1
	MAOI	N06AF	0.4	0.3
Antipsychotics	SNRI	N06AX	2.2	10.0
	RIMA	N06AG	2.1	1.4 👢
	FGA	N05A	2.2	1.6 👢
	SGA	N05A	4.6	7.1

Table 2: Psychotropic drug utilisation (in DDD/TOPD) compared by therapeutic class and subclass between 2005 and 2013 calendar years

Therapeutic class	Therapeutic subclass	ATC CODE	2005 DDD/TOPD	2013 DDD/TOPD
Anxiolytic	BDZ	N05BA	11.2	10.5
	Non-BDZ	N05BE	0.2	0.2
Hypnotic and Sedatives	BDZ Hypnotics	N05CD	25.5	17.5
	Zopiclone	N05CF	33.8	48.1
National Total			159.5	195.4

Red markings highlight increase in drug utilisation; **ATC** Anatomical Therapeutic Chemical, **BDZ** benzodiazepine, **DDD** defined daily dose, **FGA** first generation (typical) antipsychotic, **MAOI** monoamine oxidase inhibitor, **RIMA** Reversible inhibitor of monoamine oxidase-A, **SGA** second generation (atypical) antipsychotic, **SNRI** serotonin-norepinephrine reuptake inhibitor, **SSRI** selective serotonin reuptake inhibitor, **TCA** tricyclic antidepressant, **TeCA** tetracyclic antidepressant, **WHO** World Health Organization.

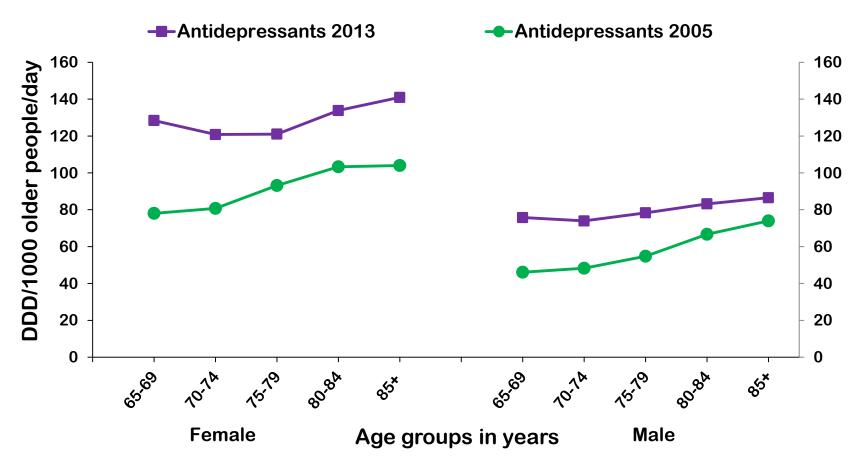


Fig 2: Utilization of antidepressant medicines normalized by sex and five-year age group

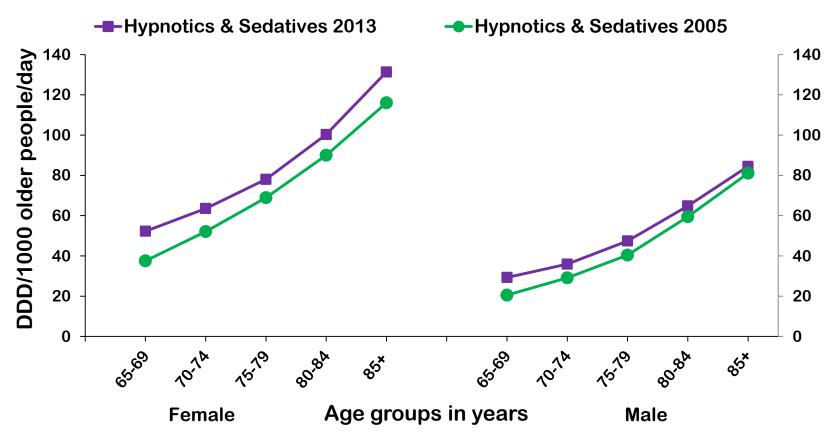


Fig 3: Utilization of hypnotic and sedative medicines normalized by sex and five-year age group

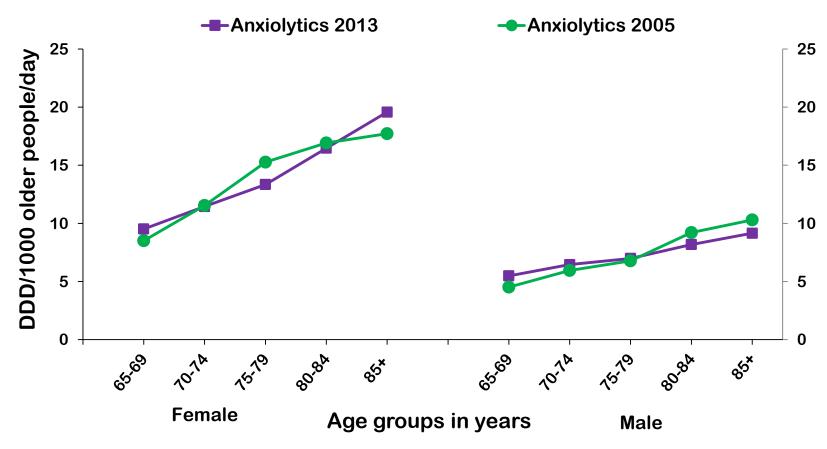


Fig 4: Utilization of psychotropic medicines normalized by sex and five-year age group

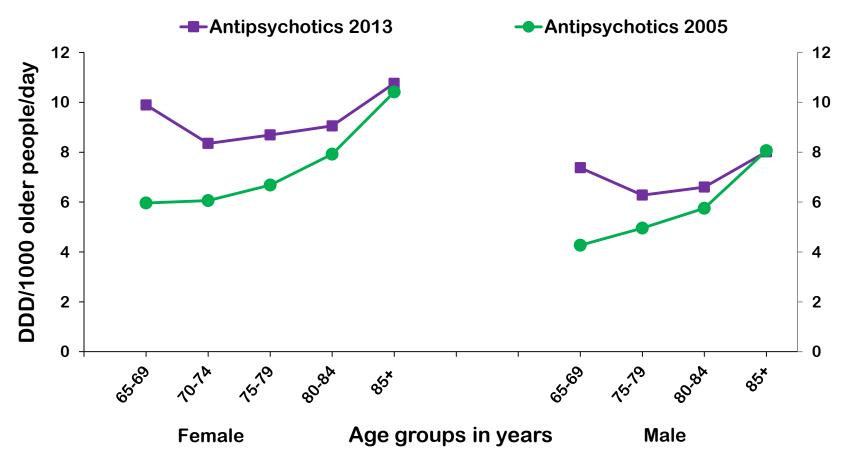


Fig 5: Utilization of antipsychotic medicines normalized by sex and five-year age group

Conclusions

 Overall, psychotropic drug utilisation in older people increased by one fifth (from 159.5 to 195.4 DDD/TOPD)
 from index date

 The utilisation of zopiclone was higher (>40%) despite it's association with adverse events in older people

 Compensatory substitution with newer psychotropic drugs like atypical antipsychotics and SSRI antidepressants

Acknowledgements

□ Research in Pharmacoepidemiology (R*i*PE) Group,
University of Otago, New Zealand

☐ Research Funding by University of Otago through a Doctoral Scholarship

□ Department of Analytical Services, Ministry of Health, New Zealand

