

TOP 5 Low-value practices and interventions

The Australasian Society of Clinical and Experimental Pharmacologists and Toxicologists (ASCEPT) is the professional and independent society in Australia and New Zealand with expertise in the use and toxicity of medicines and chemicals.

- 1** Recognise and stop the prescribing cascade

- 2** Reduce the use of medicines when there is a safer or more effective non-pharmacological management strategy

- 3** Avoid using a higher or lower dose than is necessary for the patient to optimise the 'benefit-to-risk' ratio and achieve the patient's therapeutic goals

- 4** Stop medicines when no further benefit will be achieved or the potential harms outweigh the potential benefits for the individual patient

- 5** Reduce use of multiple concurrent therapeutics (hyper-polypharmacy)

EVOLVE is a physician-led initiative to ensure the highest quality patient care through the identification and reduction of low-value practices and interventions.

EVOLVE is patient-centred and evidence-based, with rigorous and transparent processes. Its focus is to stimulate clinical conversations – between colleagues, across specialties, and with patients – to ensure the care that's delivered is the best for each patient.

EVOLVE is part of a worldwide movement to analyse medical practices and reduce unnecessary interventions. It is an initiative in partnership between the RACP and the Specialty Societies, Divisions, Faculties and Chapters.

1 Recognise and stop the prescribing cascade

A prescribing cascade occurs when a new medicine is prescribed to 'treat' an adverse reaction to another drug in the mistaken belief that a new medical condition requiring treatment has developed. Prescribing cascades should be avoided because they are associated with adverse outcomes due to the second or additional drugs, which places the patient at risk. One example of a prescribing cascade is when a patient is prescribed a non-steroidal drug for pain, and is then prescribed proton pump inhibitors (PPIs) to reduce the risk of stomach side effects caused by the first prescribed drug.

As prescribing cascades are precipitated by adverse drug reactions, they can be prevented by avoidance and early detection of the initial adverse drug reaction. For instance, many adverse drug reactions in the elderly are dose-related. It is advised that starting treatment at low doses and titrating to effect may reduce their risk. Most adverse drug reactions occur within a few months of starting a medicine. Clinicians should consider the potential for an adverse drug reaction to be the cause of any new symptoms, particularly if a drug has been recently started or changed. Patients should be asked about new symptoms, as many patients do not report adverse drug reactions. When such reactions occur, non-drug treatment strategies should be considered as the most appropriate first-line management, rather than starting a second medicine to counteract adverse effects.

2 Reduce the use of medicines when there is a safer or more effective non-pharmacological management strategy

Pharmacological treatments should be avoided or minimised if safer or more effective non-pharmacological alternatives are available. Pharmacological treatments may become a panacea for chronic lifestyle-related problems, and may detract from behaviour management tools that have proven effective in managing these same problems.

There is also a risk of adverse effects from particular pharmacological treatments which may be avoidable by using non-pharmacological management strategies. For instance, physiotherapy should be used instead of oxycodone for addressing non-cancer pain, because of the risk of adverse effects. Another example is the use of psychotropic medicines for behavioural and psychological symptoms of dementia when non-pharmacological management strategies are both more effective and safer.

3 Avoid using a higher or lower dose than is necessary for the patient to optimise the 'benefit-to-risk' ratio and achieve the patient's therapeutic goals

Therapeutic dosage should be adjusted to optimise the benefit-to-risk ratio of the treatment. Dosage should be no higher or lower than needed to achieve the patient's therapeutic goals. As patients become more frail, potential harms usually increase and potential benefits usually decrease for a given dosage of pharmacological treatment. For example, carefully assessing the risk and benefits when initiating non-steroidal inflammatory drugs in elderly patients is important, because of the increased risk of stroke associated with NSAID therapy; and use of proton pump inhibitors in the elderly should be stepped down after an initial course of therapy. Related to this, high drug doses are not necessarily more effective than low doses. An example of this is the relationship between doses of a selective serotonin re-uptake inhibitor for patients with major depressive disorder and useful clinical improvements.

4 Stop medicines when no further benefit will be achieved or the potential harms outweigh the potential benefits for the individual patient

Pharmacological treatments should cease when there are no further benefits to be achieved from the treatment, or when the potential harms from the treatment start to outweigh the potential benefits. This is particularly pertinent for elderly patients with a limited life expectancy where the treatments are unlikely to prevent disease events, and may in fact lead to adverse effects that reduce quality of life. These patients are at an increased risk of polypharmacy and increased drug events. For example, bisphosphonate treatment should not be administered to patients living in residential aged care facilities when these patients are already too frail to swallow drugs or have a life expectancy which is significantly less than 12 months.

5 Reduce use of multiple concurrent therapeutics (hyper-polypharmacy)

Polypharmacy — variously defined as more than five or up to 10 or more medications taken regularly — is common among elderly patients. However, patients who are prescribed with multiple, concurrent therapeutics may be on as many as 15 to 20 drugs at time.

Research has confirmed a significant association between polypharmacy and adverse outcomes among older people living in the community because the toxicities and side effects associated with prescribed drugs are accrued over many years.

Polypharmacy in older people is associated with decreased physical and social functioning; increased risk of falls, delirium and other geriatric syndromes; hospital admissions; and, death.

How this list was developed...

A working party of members of the Australasian Society of Clinical and Experimental Pharmacologists and Toxicologists (ASCEPT) was established to propose an initial list of recommendations. ASCEPT's membership was then invited to participate in an online survey to comment on the appropriateness of the proposed recommendations and suggest additional items for consideration.

Based on the survey responses, six recommendations were shortlisted. Following an evidence review the top 5 list items were selected. The final list was signed off by the ASCEPT President in April 2016.

Australasian Society of Clinical and Experimental Pharmacologists and Toxicologists – Top 5 recommendations

EVIDENCE SUPPORTING RECOMMENDATION 1

Caughey GE, Roughead EE, Pratt N, et al. Increased risk of hip fracture in the elderly associated with prochlorperazine: is a prescribing cascade contributing? *Pharmacoepidemiol Drug Saf* 2010; 19: 977-82

Kalisch L, Caughey G, Roughead E, Gilbert A. The prescribing cascade. *Australian Prescriber* 34(6): December 2011, 162-166

Martin J, Coombes I. Mortality from common drug interactions systems, knowledge and clinical reasoning to optimise prescribing, *Internal Medicine Journal*, 44 621-624 (2014)

EVIDENCE SUPPORTING RECOMMENDATION 2

Declercq T, Petrovic M, Azermai M. Withdrawal versus continuation of chronic antipsychotic drugs for behavioural and psychological symptoms in older people with dementia. *Cochrane Database Syst Rev* 2013; 3: CD007726.

NSW Therapeutic Advisory Group 2015. Preventing and managing problems with opioid prescribing for chronic non-cancer pain

EVIDENCE SUPPORTING RECOMMENDATION 3

Caughey G, Roughead E, Pratt N, et al. Stroke risk and NSAIDs: an Australian population-based study. *Med J Aust* 2011; 195 (9): 525-529

Dimmitt S, Stampfer H. Low drug doses may improve outcomes in chronic disease. *Med J Aust* 2009; 191 (9): 511-513.

Weekes L. Proton pump inhibitors: too much of a good thing?. *Med J Aust* 2015; 202 (9): 464

EVIDENCE SUPPORTING RECOMMENDATION 4

Ballard C, Hanney ML, Theodoulou M, et al. The dementia antipsychotic withdrawal trial (DART-AD): long-term follow-up of a randomised placebo-controlled trial. *Lancet Neurol* 2009; 8: 151-15

Maddison A, Fisher J, Johnston G. Preventive medication use among persons with limited life expectancy. *Prog Palliat Care*. 2011 Jan; 19(1): 15-21

McKellar GE, Hampson R, Tierney A, et al. Nonsteroidal antiinflammatory drug withdrawal in patients with stable rheumatoid arthritis. *J Rheumatol*. 2011 Oct;38(10):2150-2. doi: 10.3899/jrheum.101162. Epub 2011 Jul 1

Reeve E, Shakib S, Hendrix I, et al. The benefits and harms of deprescribing. *Med J Aust* 2014; 201 (7): 386-389

Scott I, Anderson K, Freeman C, Stowasser D. First do no harm: a real need to deprescribe in older patients. *Med J Aust* 2014; 201 (7): 390-392

EVIDENCE SUPPORTING RECOMMENDATION 5

Davies E, Green C, Taylor S, et al. Adverse Drug Reactions in Hospital In-Patients: A Prospective Analysis of 3695 Patient-Episodes. *PLoS ONE* 4(2): e4439. doi:10.1371/journal.pone.0004439

Hubbard R, Peel N, Scott I. et al. Polypharmacy among inpatients aged 70 years or older in Australia. *Med J Aust* 2015; 202 (7): 373-377

Scott I, Anderson K, Freeman C, Stowasser D. First do no harm: a real need to deprescribe in older patients. *Med J Aust* 2014; 201 (7): 390-392