

Recognising and Managing Polypharmacy in Practice

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Overview

- Background to polypharmacy
- How to manage polypharmacy
- Tools for use in practice

Why do we prescribe?

Primary prevention
Treatment
Symptom control

How do we make prescribing decisions?

National guidelines
Evidence based
Experience

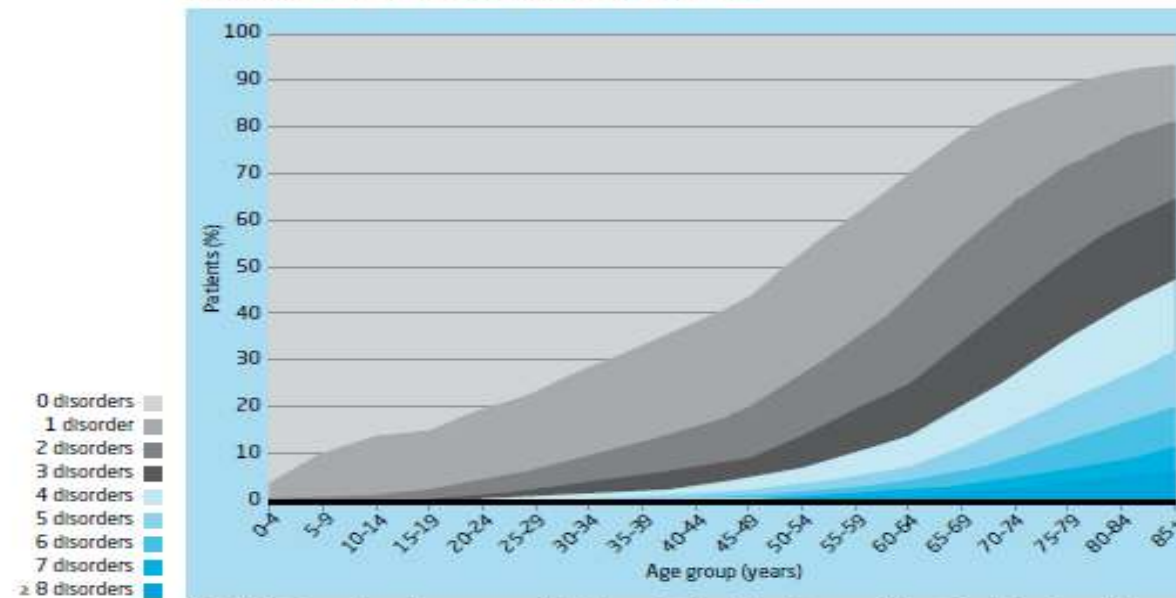
Multi-morbidity

Multi-morbidity defined as the presence in an individual of 2 or more long term conditions (LTC)

Can include:

- Defined physical and mental health conditions
- Ongoing conditions such as learning disability
- Symptom complexes such as frailty or chronic pain
- Sensory impairment such as sight or hearing loss
- Alcohol and substance misuse

Figure 5 Number of chronic disorders by age group



Note: This figure shows how common it is to have significant long-term conditions in relation to age. Few people (fewer than 30 per cent) do not have at least one condition by the age of 60, and many people will have two or three.

Source: Barnett et al (2012)⁵

Multi-morbidity

- Growing number of people with a LTC – around 15 million people in England now have a LTC and the number of LTC increases with age.
- Average number of prescription items per year for any one person in England is increasing
- Estimated between 30% – 50% of medicines for LTC are not taken as intended

<https://www.nice.org.uk/guidance/ng56>

Polypharmacy

The Kings Fund definition polypharmacy:

Appropriate polypharmacy

‘Prescribing for an individual for complex conditions or for multiple conditions in circumstances where medicines use has been optimised and where the medicines are prescribed according to best evidence’

Problematic polypharmacy

‘The prescribing of multiple medicines inappropriately, or where the intended benefit of the medicines are not realised’

Polypharmacy epidemiology

In 2015/16 48% of adults had taken at least 1 prescribed medicine (not including contraception or nicotine replacement)

This increases with age to more than 90% of those aged 75 and over

24% of adults were taking 3 or more medications

<http://healthsurvey.hscic.gov.uk/media/63790/HSE2016-pres-med.pdf>

Adherence

- Adherence is an important but often overlooked factor in polypharmacy. WHO in 2003, highlighted that 30-50% of medicines are not taken as intended (www.who.int/chp/knowledge/publications/adherence_full_report.pdf).
- Often further medicines are prescribed in response to 'treatment failure' rather than a check of the level of treatment adherence. This was illustrated by a recent study ([BJ Clinical Pharmacology Vol 84 Issue 1 Jan 2018 18-24](#)) which highlighted that hypertension is only controlled in 35% of people. Those in this study were assessed for adherence to their blood pressure medicines (physicians and people under their care were unaware of adherence measurements), and 68% of the people prescribed medicines were non-adherent.

How did we get here?

- Treating the disease and not the patient
- Prescribing pressures
- Medical and nursing specialism
- Working in silos
- Lack of research and evidence in relation to those who are very old
- Expectation that we can live forever

Tools to assess appropriateness of de-prescribing

Aim is to review whether the risk of harmful effects exceeds the potential benefit

- Beers criteria (American Geriatrics society Updated 2019)

https://nicheprogram.org/sites/niche/files/2019-02/Panel-2019-Journal_of_the_American_Geriatrics_Society.pdf

- STOPP/START

<http://ageing.oxfordjournals.org/content/early/2014/10/16/ageing.afu145.full>

- Anticholinergic burden

<http://www.medicheck.com/assessment>

- Scottish guidelines

Scottish Government Polypharmacy Model of Care group (2018)

Anticholinergic Burden Scale

Drugs with ACB Score of 1		Drugs with ACB Score of 2		Drugs with ACB Score of 3	
Generic Name	Brand Name	Generic Name	Brand Name	Generic Name	Brand Name
Alimemazine	Theralan™	Amantadine	Symmetrel™	Amitriptyline	Elavil™
Alverine	Spasmonal™	Belladonna	Multiple	Amoxapine	Asendin™
Alprazolam	Xanax™	Carbamazepine	Tegretol™	Atropine	Sal-Tropine™
Aripiprazole	Abilify™	Cyclobenzaprine	Flexeril™	Benztropine	Cogentin™
Asenapine	Saphris™	Cyproheptadine	Periactin™	Brompheniramine	Dimetapp™
Atenolol	Tenormin™	Loxapine	Loxitane™	Carbinoxamine	Histex™, Carbihist™
Bupropion	Wellbutrin™, Zyban™	Mependine	Domerol™	Chlorpheniramine	Chlor-Trimeton™
Captopril	Capoten™	Methotrimeprazine	Lavopromia™	Chlorpromazine	Thorazine™
Cotizine	Zyrtec™	Molindone	Moban™	Clemastine	Tavist™
Chlorthalidone	Diuril™, Hygroton™	Nefopam	Nefogesic™	Clomipramine	Anafranil™
Cimetidine	Tagamet™	Oxcarbazepine	Trileptal™	Clozapine	Clozaril™
Clidinium	Librax™	Rimozide	Orap™	Darifenacin	Enablix™
Clorazepate	Tranxene™			Desipramine	Norpramin™
Codine	Contin™			Dicyclomine	Bentyl™
Colchicine	Colcrys™			Dimenhydrinate	Dramamine™, others
Doxylorstatine	Clarinex™			Diphenhydramine	Benadryl™, others
Diazepam	Valium™			Doxepin	Sinequan™
Digoxin	Lanoxin™			Doxylamine	Unisom™, others
Dipyridamole	Persantine™			Fesoterodine	Toviaz™
Disopyramide	Niaropa™			Flavoxate	Urispas™
Fentanyl	Duragesic™, Actiq™			Hydroxyzine	Atarax™, Vistaril™
Furosemide	Lasix™			Hyoscyamine	Anaspaz™, Levsin™
Fluvoxamine	Luvex™			Imipramine	Tofranil™
Haloperidol	Haldol™			Meclizine	Antivert™
Hydralazine	Apresoline™			Methocarbamol	Robaxin™
Hydrocortisone	Cortef™, Cortaid™			Nortriptyline	Pamelor™
Iloperidone	Fanapt™			Olanzapine	Zyprexa™
Isoorbide	Isordil™, Ismo™			Orphenadrine	Norflex™
Levocetirizine	Xyzal™			Oxybutynin	Ditropan™
Loperamide	Immodium™, others			Paroxetine	Paxil™
Loratadine	Claritin™			Perphenazine	Trilafon™
Metoprolol	Lopressor™, Toprol™			Promethazine	Phenergan™
Morphine	MS Contin™, Avinza™			Propranolol	Pro-Banthine™
Nifedipine	Procardia™, Adalat™			Propiverina	Detrunorm™
Paliperidone	Invega™			Quetiapine	Seroquel™
Prednisone	Deltasone™, Sterapred™			Scopolamine	Transderm Scop™
Quinidine	Quinaglute™			Solifenacin	Vesicare™
Ranitidine	Zantac™			Thioridazine	Mellilan™
Risperidone	Risperdal™			Tolterodine	Detrol™
Theophylline	Theodur™, Uniphyll™			Trifluoperazine	Stelazine™
Trazodone	Desyrel™			Trihexyphenidyl	Artane™
Triamterene	Dyrenium™			Trimipramine	Surmontil™
Venlafaxine	Effexor™			Tropium	Sanctura™
Warfarin	Coumadin™				

Categorical Scoring:

- Possible anticholinergics include those listed with a score of 1; Definite anticholinergics include those listed with a score of 2 or 3

Numerical Scoring:

- Add the score contributed to each selected medication in each scoring category
- Add the number of possible or definite Anticholinergic medications

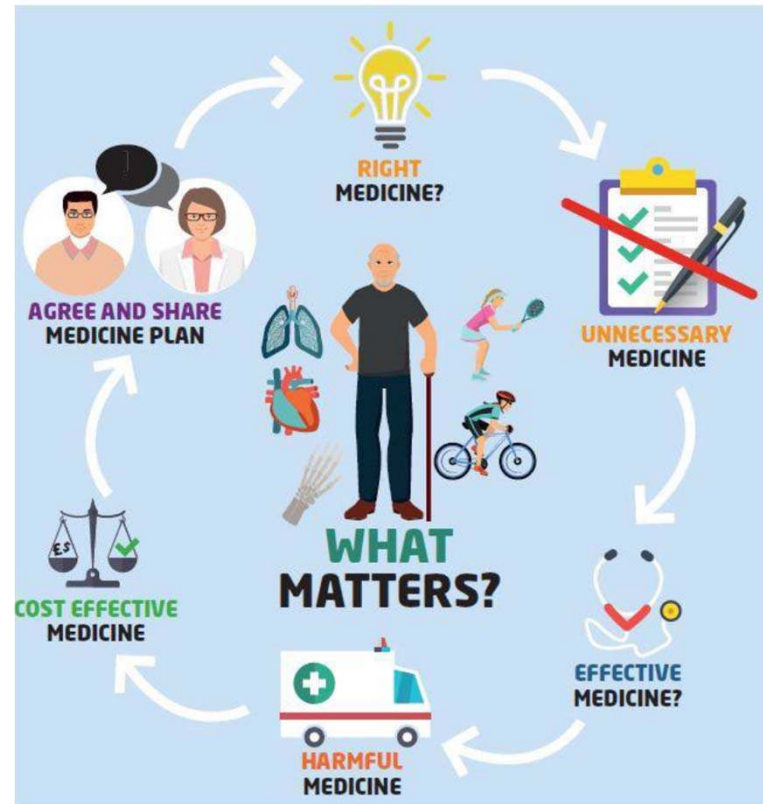
Notes:

- Each definite anticholinergic may increase the risk of cognitive impairment by 46% over 6 years.³
- For each one point increase in the ACB total score, a decline in MMSE score of 0.33 points over 2 years has been suggested.⁴
- Additionally, each one point increase in the ACB total score has been correlated with a 26% increase in the risk of death.⁴

Aging Brain Care

www.agingbraincare.org

Medicines Review



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What matters to the patient (Aim)

- Identify aims and objectives of drug therapy by asking the patient *what matters to you*
- Explain any key information such as laboratory markers
- Establish treatment objectives with patient through shared decision making

Does the patient take unnecessary drug therapy (Need)

- For the remaining drugs, it should be verified that each has a function in achieving the therapeutic goals or outcomes that matter most to the patient
- Review preventative treatment to ensure the patient is able to continue taking medicine for required time to gain benefit (Drug efficacy (NNT))
- Can lifestyle changes replace any unnecessary drug therapy?

Are therapeutic objectives being achieved (Effectiveness)

- Check treatment choice is the most effective to achieve intended outcomes
- If this is not the case, the possibility of patient non-adherence should be investigated as a potential explanation. Otherwise, the need for dose titration may also be considered. 50% of patients taking four or more medicines don't take them as prescribed

Is the patient at risk of ADR's or suffers actual ADR's (Safety)

- The presence of ADR's can sometimes be identified from laboratory data (eg hypokalaemia from diuretic use)
- The patient may report such symptoms (including drug-drug and drug-disease interactions)
- Ask the patient specific questions (eg about the presence of anticholinergic symptoms, dizziness or drowsiness). If patient is experiencing ADR's use Yellow card reporting

Is the drug therapy cost-effective? (Efficiency)

- Opportunities for cost minimisation should be explored, but changing drugs for cost reasons should only be considered if effectiveness, safety or adherence would not be compromised
- Ensure prescribing is in line with current formulary recommendations

Is the patient willing and able to take drug therapy as intended (Patient-centred)

- Does the patient understand the outcome of the review?
- Ensure drug therapy is tailored to patient preferences
- Agree and communicate plan with patients and/or welfare proxy
- Even if adult lacks capacity, their views should still be sought.

In summary:

Risk v benefit – when the risk of harmful effects exceeds the potential benefit for the patient – consider whether the medication should be stopped

Deprescribing is all of our responsibility

Use the tools to help you with your decision making