

## STOPP START Tool to Support Medication Review

Older people are known to have increased risk of adverse effects with medication due to age related alteration in pharmacokinetics and pharmacodynamics. This can sometimes lead to harm rather than benefit from a particular treatment.

Polypharmacy and inappropriate prescribing are well known risk factors for adverse drug reactions (ADRs), which commonly cause adverse clinical outcomes in older people.<sup>1</sup>

Systematic reviews and published evidence suggest some common drug groups associated with preventable drug related admissions especially in the elderly.<sup>2,3</sup> Hence prioritising to review these drug or drug groups helps reduce polypharmacy and the burden of ineffective or unnecessary treatment in the frail and elderly.

The recently published NICE guidance on Medicines Optimisation<sup>4</sup> recommends using a screening tool – for example the STOPP/START tool in older people – to identify potential medicines-related patient safety incidents for those on multiple medicines or with long term conditions.

This document is an adaptation of the

### **STOPP START medication review screening tool (STOPP-Screening Tool of Older Persons Prescriptions START -Screening Tool to Alert doctors to Right i.e. appropriate, indicated Treatments)**

Which aids physicians appraise older patient's medication in the context of their current clinical condition<sup>5</sup>.

Eighteen experts in geriatric pharmacotherapy initially contributed to suggesting and then rating the criteria.

The tool was validated in patients aged 65 and over but physicians must use their clinical judgement when deciding if a person is "elderly" in terms of using the toolkit and also consider other drug interactions or contra-indications not listed here.

The final decision to stop the drug should be weighed against the daily symptomatic benefit or prevention of rapid worsening of symptoms.

Where there is any doubt with the above information please check that it is in line with manufacturers recommendations, published literature or changes in national and local guidance. All Wirral guidance can be found at <http://mm.wirral.nhs.uk/forumulary/> or <http://mm.wirral.nhs.uk/guidelines/>

Adapted by Abigail Cowan, Prescribing Adviser North West Commissioning Support Unit with permission from STOPP/START Tool V9 – Dr D O'Mahony ([denis.omahony@ucc.ie](mailto:denis.omahony@ucc.ie)) and Dr Simon Conroy ([spc3@le.ac.uk](mailto:spc3@le.ac.uk)). Acknowledgments also to NHS Cumbria STOPP/START Toolkit Feb 2013 & Leicestershire Medicines Strategy Group Nov 2014 (adapted with permission).

Date of preparation	Written by	Checked by	Date of next review	Version
March 2015	Abigail Cowan	Steve Riley	March 2016	1

STOP medications (age ≥ 65 years)	Circumstances to review	Reason to review
<b>α-blockers</b>	Long-term urinary catheter in situ >2 Months	Not indicated
<b>Antibiotics Review</b>	IV antibiotics - ensure review date has been discussed with microbiology (under OPAT team) with the aim to switch to oral if possible  Long-term antibiotics for UTI prophylaxis	Review individual patients as recommended by microbiology – guidance at <a href="http://www.rcgp.org.uk/TArGeTantibiotics/">http://www.rcgp.org.uk/TArGeTantibiotics/</a>  Risk of infected/phlebotic IV lines. Increased risk of adverse effects and errors in preparation are significantly higher with parenteral drugs, compared to oral formulations. Increased patient discomfort and reduced mobility  Refer to urology. Patients should be reviewed at regular intervals to assess the risk: benefits in relation to C. difficile infection. Prophylactic antibiotics should be reviewed after 6 months and stopping should be considered.
<b>Anticholinergics</b>	To treat extra-pyramidal side-effects of antipsychotic medications	Risk of anticholinergic toxicity, including confusion and urinary retention
<b>Anticholinergic antispasmodics (e.g. hyoscine butylbromide)</b>	For patients with chronic constipation	Risk of exacerbation of constipation
<b>Antidiarrhoeal drugs (co-phenotrope, loperamide or codeine phosphate)</b>	For treatment of diarrhoea of unknown cause <i>N.B. Please be aware of C. difficile in undiagnosed diarrhoea</i>	Risk of delayed diagnosis, may exacerbate constipation with overflow diarrhoea, may precipitate toxic mega colon in inflammatory bowel disease, may delay recovery in unrecognised gastroenteritis
<b>Antidiarrhoeal drugs (co-phenotrope, loperamide or codeine phosphate)</b>	For the treatment of severe infective gastroenteritis	Risk of exacerbation or protraction of infection Risk of colitis and toxic mega colon if Clostridium difficile
<b>Antimuscarinics (bladder)</b>	Dementia or glaucoma or constipation or prostatism	Risk of worsening respective condition NICE CG171 Urinary Incontinence in Women <a href="https://www.nice.org.uk/guidance/cg171">https://www.nice.org.uk/guidance/cg171</a> Joint Wirral Medicines Formulary – Urinary Tract Disorders <a href="http://mm.wirral.nhs.uk/guidelines/">http://mm.wirral.nhs.uk/guidelines/</a>
<b>Antipsychotics</b>  <i>NB. Reduce slowly monitoring effect</i>	>1 month use as long-term hypnotic (check notes for duration) >1 month use in parkinsonism  If fallen in last 3 months For treatment of behavioural and psychological symptoms of dementia patients (review ongoing need)	Confusion, ↓BP, extrapyramidal side effects, falls  Risk of worsening extrapyramidal symptoms  May cause gait dyspraxia, parkinsonism Risk of gait disturbances, dehydration, prolonged sedation, cognitive decline, falls, stroke and death
<b>Aspirin</b>	Dose >150mg / day, restart at 75mg if still indicated  With a concurrent bleeding disorder  Peptic ulcer disease without histamine H2 receptor antagonist or PPI  If being used as monotherapy for stroke prevention in AF	Risk of bleeding; no evidence of increased efficacy  High risk of bleeding  Risk of bleeding  Guidance at: <a href="https://www.nice.org.uk/guidance/cg180">https://www.nice.org.uk/guidance/cg180</a>
<b>Benzodiazepines – reduce slowly &amp; monitor effect</b>	>1 month use of long-acting benzodiazepines, eg. chlordiazepoxide, oxazepam, diazepam, flurazepam, nitrazepam If fallen in last 3 months	Risk of prolonged sedation, confusion, impaired balance, falls

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<b>Beta-blocker</b>	In combination with verapamil  In those with diabetes mellitus and frequent hypoglycaemic episodes	Risk of symptomatic heart block  Risk of masking hypoglycaemic symptoms
<b>Beta-blocker (non-cardioselective)</b>	In patients with asthma	Risk of bronchospasm
<b>Bisphosphonates (oral)</b>	Unable to sit upright / patient experiencing swallowing difficulties / compliance issues	Instruction for administration of medication if not followed causes increased risk of serious upper GI disorder Wirral Guidelines for the Management of Osteoporosis <a href="http://mm.wirral.nhs.uk/guidelines/">http://mm.wirral.nhs.uk/guidelines/</a>
<b>Calcium Channel Blocker</b>	If ankle oedema present  Verapamil and diltiazem should usually be avoided in heart failure	This may be an adverse effect of the Calcium Channel Blocker see UKMI Q&A 330.2; <a href="http://www.ukmi.nhs.uk/activities/medicinesQAs/default.asp">http://www.ukmi.nhs.uk/activities/medicinesQAs/default.asp</a>  They may further depress cardiac function and cause clinically significant deterioration.
<b>Carbocisteine</b>	If no benefit after 4 weeks	Unnecessary if no benefit shown
<b>Clopidogrel</b>	With concurrent bleeding disorder	High risk of bleeding
<b>Corticosteroids</b>	Systemic instead of inhaled corticosteroids for maintenance therapy in moderate-severe COPD >3 months as monotherapy for rheumatoid or osteoarthritis	Unnecessary exposure to long-term side effects of systemic steroids.  Risk of major systemic corticosteroids side effects
<b>Digoxin</b>	At doses >125 microgram per day with impaired renal function (eGFR <50ml/minute)	Can be increased levels of toxicity (e.g. nausea, diarrhoea, arrhythmias)
<b>Dipyridamole monotherapy</b>	With concurrent bleeding disorder	High risk of bleeding
<b>Diuretics (loop)</b>	Dependent ankle oedema and no signs of heart failure  As first line monotherapy for hypertension	No benefit; compression hosiery more appropriate  Safer, more effective alternatives available
<b>Diuretics (thiazides)</b>	With history of gout	Risk of exacerbating gout
<b>Domperidone</b>	Indications except nausea/vomiting Underlying Cardiac conditions, impaired cardiac conduction, co-prescribed other medications known to prolong QT interval or potent CYP3A4 inhibitors or with severe hepatic impairment	See MHRA warning issued <a href="https://www.gov.uk/drug-safety-update/domperidone-risks-of-cardiac-side-effects">https://www.gov.uk/drug-safety-update/domperidone-risks-of-cardiac-side-effects</a>
<b>Ipratropium (nebulised)</b>	Prescribing as required (prn) in addition to regular prescribing With glaucoma	Can lead to exceeding licensed dosage and therefore exacerbate side effects May exacerbate glaucoma
<b>Laxatives – stimulant (e.g. bisacodyl, senna)</b>	For patients with intestinal obstruction	Risk of bowel perforation  <a href="http://mm.wirral.nhs.uk/guidelines/">http://mm.wirral.nhs.uk/guidelines/</a>

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STOP medications (age ≥ 65 years)	Circumstances to review	Reason to review
<b>Metformin</b>	Renal impairment: Review dose if eGFR <45mL/minute Avoid if eGFR <30ml/minute	Increased risk of lactic acidosis
<b>NSAID (oral)</b>	History of PUD or GI bleed unless with H2 antagonist, PPI or misoprostol  Moderate severe hypertension (moderate 160/100mm Hg - 179/109mm Hg; severe: >180/110mm Hg  Heart failure  With Warfarin  65+ and e GFR <60ml/min or heart failure  On long-term NSAID and colchicine for chronic treatment of gout when there is no C/I to allopurinol  Long-term corticosteroids as monotherapy (>3 month for arthritis)  Cox-2 inhibitors and diclofenac in cardiovascular disease Ibuprofen (at total daily dose above 1200mg per day) in cardiovascular disease	Risk of peptic ulcer relapse  Risk of exacerbation of hypertension  Risk of exacerbation of heart failure  Risk of GI bleeding  Risk of deterioration in renal function  Allopurinol first choice prophylactic in gout  Risk of GI bleed and systemic corticosteroids side effects  Increased risk of thrombotic events  Increased risk of thrombotic events
<b>Oestrogen (systemic)</b>	With history of breast cancer or venous thromboembolism  Without progesterone in patients with intact uterus	Increased risk of reoccurrence  Risk of endometrial cancer
<b>Omega-3 fatty acids</b>	Prescribed for secondary prevention of MI	Review as per <a href="http://www.nice.org.uk/guidance/cg172/resources/guidance-mi-secondary-prevention-pdf">http://www.nice.org.uk/guidance/cg172/resources/guidance-mi-secondary-prevention-pdf</a>
<b>Opioids (all type)</b>	Long-term use of powerful opiates (e.g. morphine, fentanyl) as first line therapy for mild-moderate pain  Regular prescription >2 weeks in chronic constipation without concurrent use of laxatives	WHO analgesic ladder not observed  Risk of severe constipation
<b>Pioglitazone (glitazones)</b>	Heart failure and elderly patients	Increased risk of fracture, bladder cancer and heart failure
<b>Prochlorperazine or metoclopramide</b>	With Parkinsonism	Risk of exacerbating Parkinsonism. Metoclopramide is for 5 days use only <a href="#">Drug Safety Update August 2013</a>
<b>PPI treatment</b>	Dose for PUD for more than 8 weeks	Earlier discontinuation or dose reduction for maintenance/prophylactic treatment of PUD, oesophagitis or GORD indicated Increased risk of <i>C. difficile</i> infection, pneumonia, bone fractures and hypomagnesaemia <a href="http://mm.wirral.nhs.uk/guidelines/">http://mm.wirral.nhs.uk/guidelines/</a> <a href="http://www.nice.org.uk/guidance/CG184/">http://www.nice.org.uk/guidance/CG184/</a>

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<b>PPI (omeprazole or esomeprazole)</b>	If co-prescribed clopidogrel	MHRA Drug Safety Update 2010 advises that concurrent use should be discouraged due to reduced antiplatelet effect, see <a href="https://www.gov.uk/drug-safety-update/clopidogrel-and-proton-pump-inhibitors-interaction-updated-advice">https://www.gov.uk/drug-safety-update/clopidogrel-and-proton-pump-inhibitors-interaction-updated-advice</a>
<b>SSRIs</b>	If sodium less than 130 in past 2 months Citalopram & escitalopram – risk of QT prolongation	SSRIs can cause/worsen hyponatraemia  Don't use in patients with congenital long QT syndrome or known pre-existing QT interval prolongation In combination with other drugs known to prolong the QT intervals
<b>Statins</b>	Prognosis of less than six months unless there is an acute vascular syndrome  In patients displaying symptoms of muscle weakness and pain	In the absence of a recent acute coronary syndrome or cerebrovascular event, the discontinuation of a statin toward the end of life is reasonable  Risk of myopathy and rhabdomyolysis. Check creatinine kinase if patient presents with muscular symptoms.
<b>Sulfonylureas (particularly Glibenclamide or Chlorpropamide)</b>	With Type 2 diabetes	Risk of prolonged hypoglycaemia
<b>Theophylline</b>	Monotherapy for COPD	Safer, more effective alternatives, risk of adverse effects due to narrow therapeutic index
<b>Tricyclic antidepressants</b>  <i>NB. Withdraw gradually over at least 4 weeks – monitor effect</i>	Dementia  Glaucoma  Cardiac conductive abnormalities  Constipation  Combination with opiate or calcium channel blocker  Prostatism or history of urinary retention  Patients taking dosulepin	Risk of worsening cognitive impairment  May exacerbate glaucoma if untreated  Pro-arrhythmic effects  May worsen constipation  Risk of severe constipation  Risk of urinary retention  Increased cardiac risk & toxicity in overdose
<b>Vasodilator drugs (e.g. hydralazine, minoxidil)</b>	With persistent postural hypotension i.e. recurrent > 20 mmHG drop in systolic blood pressure	Risk of syncope and falls
<b>Warfarin</b>	For 1 <sup>st</sup> uncomplicated DVT for longer than 6 months or PE for longer than 12 months  Hepatic impairment with impaired clotting ability and raised INR	No proven added benefit  Increased risk of bleeding as a result of impaired ability to produce clotting factors
<b>Any regular duplicate drug class prescription</b>	E.g. two concurrent opiates, multiple NSAIDs, multiple diuretics  Two or more anticholinergics (antimuscarinics)	Optimisation of monotherapy within a single drug class prior to considering a new drug class  Increased risk of side-effects including confusion falls and death

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START medications (age ≥ 65 years)	Circumstances
<b>ACE Inhibitor</b>	Chronic heart failure Following acute myocardial infarction Diabetes with nephropathy (overt urinalysis, proteinuria or microalbuminuria) >30mg / 24 hours ± serum biochemical renal impairment
<b>Antidepressants</b>	In presence of moderate to severe depressive symptoms lasting at least three months
<b>Antihypertensive</b>	Systolic blood pressure consistently >160mm Hg
<b>Aspirin</b>	Documented history of atherosclerotic coronary, cerebral or peripheral vascular disease in patients with sinus rhythm Following an acute MI
<b>Beta-blocker (oral)</b>	With chronic stable angina
<b>Beta-agonist (inhaled)</b>	Guidance at <a href="http://mm.wirral.nhs.uk/document_uploads/guidelines/COPDguidelinesv2.pdf">http://mm.wirral.nhs.uk/document_uploads/guidelines/COPDguidelinesv2.pdf</a> Review patients with mild, moderate or severe COPD at least once a year, and very severe COPD at least twice a year as per NICE guidance - <a href="http://www.nice.org.uk/guidance/cg101">http://www.nice.org.uk/guidance/cg101</a>
<b>Bisphosphonates</b>	In patients taking maintenance oral corticosteroid therapy with previous fragility fractures or incident fractures during glucocorticoid therapy. Ensure there are no absorption interactions e.g. Calcium. Counsel patient on the correct way to take a bisphosphonate.
<b>Calcium and vitamin D</b>	In patients with known osteoporosis (radiological evidence or previous fragility fracture) or acquired dorsal kyphosis
<b>Clopidogrel</b>	For ischaemic stroke or PVD as per <a href="http://www.nice.org.uk/guidance/ta210">http://www.nice.org.uk/guidance/ta210</a>
<b>DMARD</b>	With active moderate-severe rheumatoid disease lasting >12 weeks
<b>Fibre supplement</b>	For chronic symptomatic diverticular disease with constipation
<b>Laxatives</b>	In patients taking opioids - to prevent constipation
<b>PPI</b>	For severe reflux or peptic stricture requiring dilatation For patients over 80 years old on antiplatelets and SSRI
<b>Statins</b>	Documented history of coronary, cerebral or peripheral vascular disease, where the patient's functional status remains independent for activities of daily living and life expectancy >5 years Diabetes mellitus plus ≥ 1 co-existing major cardiovascular risk factor present
<b>Anticoagulation (warfarin or a NOAC)</b>	Chronic atrial fibrillation as per <a href="http://www.nice.org.uk/guidance/cg180">http://www.nice.org.uk/guidance/cg180</a> Following diagnosis of DVT and PE if benefit outweighs the risk of treatment

## References

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