

Clinical implementation of medication review - screening to supervise effectiveness and appropriateness

Adapted from Dr Bertrand Guignard presentation

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Medicines Review – Needing and Sharing the
Hospital Pharmacist's
EAHP Academy seminar
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CONFLICT OF INTERESTS

**Member of the editorial committee of
PIM-Check**



Learning session questions

Mr X., with medical history of COPD (classification GOLD 3), is admitted in internal medicine for community-acquired pneumonia, with moderate hypokalaemia (2.8 mmol/L). His admission prescription includes :

- Beclometasone 200 mcg BID
- Ciprofloxacin 400 mg BID PO
- Ceftazidime 6 g IV continuous infusion (solvent: NaCl 0.9% 100mL)
- KCl 40mmol IV 6h-infusion (solvent: NaCl 0.9% 500 ml) QD

What verbal advice can you give during the medical round ?

1. Ceftazidime and KCl are physically incompatible. Propose stopping and rinsing the infusion line before KCl administration
True or false
2. Ciprofloxacin associated with hypokalaemia is at risk of torsade de pointes. Propose an electrocardiogram monitoring.
True or False
3. Inhaled corticosteroids should not be prescribed as a first-line treatment and/or as a monotherapy to treat COPD. Consider prescribing beta₂-agonist or anticholinergics as first line treatment
True or False

CLINICAL CASE

Clinical Case

Mrs X, 73 years old, 162 cm, 50 kg

✧ Admission pattern

- ✧ Back pain,
- ✧ Fever for 5 days,
- ✧ Increased frequency of urination

✧ Physical examination

- ✧ Fever (39.5° C),
- ✧ Tachycardia,
- ✧ Dyspnea on moderate levels of exertion
- ✧ BP : 140/90
- ✧ Heart rate : 110 beats per minute
- ✧ Respiratory rate : 28 breaths per minute

✧ Laboratory test results

- | | |
|-------------------------------------|-------------------|
| ✧ eGFR: 45mL/min/1.73m ² | (90 - 130 ml/min) |
| ✧ White blood cells: 20 G/L | (4 - 11 G/L) |
| ✧ CRP: 344 | (<10) |
| ✧ Potassium: 2.7 mmol/L | (3.5 - 5 mmol/L) |

✧ Medical diagnostic

- ✧ Acute pyelonephritis with sever sepsis
- ✧ Cardiac decompensation (LVEF 30%)

✧ Past medical history

- ✧ Stable ischemic heart disease
 - ✧ STEMI (2011, with stent),
 - ✧ NSTEMI (2014, with stent)
- ✧ HBP
- ✧ Persistent atrial fibrillation (AF)
- ✧ Chronic kidney disease (eGFR : 45 mL/min)
- ✧ Leading cause: High blood pressure
- ✧ Depression, anxiety, sleep disorders
- ✧ Ø OH, Ø Tobacco



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Clinical Case: treatment 48h after admission

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole 40 mg QD
- Acenocoumarol QD dose adjusted according to INR
- Diltiazem extended-release 90 mg BID
- Fluoxetine 20 mg QD
- Furosemide 20 mg QD
- Oxazepam 15 mg QD
- OTC : paracetamol, ibuprofen

► Added at admission

- KCl 40 mmol TID IV infusion over 1h (solvent: NaCl 0.9% 250 ml)
- Magnesium sulfate 2 g continuous IV infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min QD (solvent: NaCl 0.9% 100 ml)
- Tramadol 25 mg QID PO, then 50 mg QID, poorly effective

Drug-related problems (DRPs)

► Definition :

*“Event or circumstance involving drug therapy that **actually** or **potentially** interferes with desired health outcomes”*



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Hepler CD, Strand LM. Opportunities and responsibilities in pharmaceutical care. Am J Hosp Pharm. 1990 Mar;47(3):533-43.

Drug-related problems (DRPs)

Hepler CD, et al. definition (1990)

- 1) Untreated indication (under-prescription)
- 2) Drug use without indication (over-prescription)
- 3) Improper Drug Selection (mis-prescription)
- 4) Subtherapeutic Dosage
- 5) Overdosage
- 6) Adverse drug reaction
- 7) Drug interaction
- 8) Failure to receive a drug

Drug-related problems (DRPs)

DRPs added in some other countries

- 1) Untreated indication (under-prescription)
- 2) Drug use without indication, **duplication** (over-prescription)
- 3) Improper Drug Selection (mis-prescription)
- 4) Subtherapeutic Dosage
- 5) Overdosage
- 6) Adverse drug reaction
- 7) Drug interaction
- 8) Failure to receive drug
- 9) **Contra-indication**
- 10) **Insufficient drug monitoring**
- 11) **Improper route of administration**
- 12) **Improper drug form**
- 13) **Improper treatment duration**



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SFPC. Standardisation et valorisation des activités de pharmacie clinique :
<http://www.actip.sfpc.eu/actip/index/ficheip/>

GSASA: Documentation des activités cliniques
<http://www.gsasa.ch/pages/activites/activites-cliniques/?oid=1587&lang=FR>

Drug therapy appropriateness

Medication Appropriateness Index (MAI)

Table 1. Medication Appropriateness Index*

To assess the appropriateness of the drug, please answer the following questions and circle the applicable score:				
1. Is there an indication for the drug? Comments:	1 Indicated	2	3 Not Indicated	9 DK†
2. Is the medication effective for the condition? Comments:	1 Effective	2	3 Ineffective	9 DK
3. Is the dosage correct? Comments:	1 Correct	2	3 Incorrect	9 DK
4. Are the directions correct? Comments:	1 Correct	2	3 Incorrect	9 DK
5. Are the directions practical? Comments:	1 Practical	2	3 Impractical	9 DK
6. Are there clinically significant drug-drug interactions? Comments:	1 Insignificant	2	3 Significant	9 DK
7. Are there clinically significant drug-disease/condition interactions? Comments:	1 Insignificant	2	3 Significant	9 DK
8. Is there unnecessary duplication with other drug(s)? Comments:	1 Necessary	2	3 Unnecessary	9 DK
9. Is the duration of therapy acceptable? Comments:	1 Acceptable	2	3 Unacceptable	9 DK
10. Is this drug the least expensive alternative compared to others of equal utility? Comments:	1 Least expensive	2	3 Most expensive	9 DK

*Complete instructions in the use of the scale are available upon request.

†Don't know.



What are we going to talk about today ?

- ▶ Drug – Drug Interactions
- ▶ Doses adjustment
- ▶ Drug administration
- ▶ Adverse drug reactions
- ▶ Inappropriate prescribing:
 - ▶ Under-prescriptions
 - ▶ Over-prescriptions
 - ▶ Mis-prescriptions



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DRUG – DRUG INTERACTIONS

Drug – Drug Interactions (DDI)?

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole 40 mg QD
- Acenocoumarol QD dose adjusted according to INR
- Diltiazem extended-release 90 mg BID
- Fluoxetine 20 mg QD
- Furosemide 20 mg QD
- Oxazepam 15 mg QD
- OTC : paracetamol, ibuprofen

► Added at admission

- KCl 40 mmol TID IV 1h-infusion (solvent: NaCl 0.9% 250 ml)
- Magnesium sulfate 2 g IV continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- Tramadol 25 mg QID PO, then 50 mg QID, poorly effective

Drug – Drug Interactions: Sources

► Software and Apps



- Lexi-Interact (Lexi Comp's):
www.lexi.com



- Micromedex Drugs interactions:
www.micromedexsolutions.com



- Epocrates:
www.epocrates.com

► Tables

► Cytochrome Tables:

[www.hug-ge.ch/sites/interhug/files/structures/pharmacologie et toxicologie cliniques/documents/interactions medicamenteuses et cyp450.pdf](http://www.hug-ge.ch/sites/interhug/files/structures/pharmacologie_et_toxicologie_cliniques/documents/interactions_medicamenteuses_et_cyp450.pdf)

► Books

- Stockley's Drug Interactions
- Drugdex

► Specific tools



- HIV: www.hiv-druginteractions.org



- HCV: www.hep-druginteractions.org

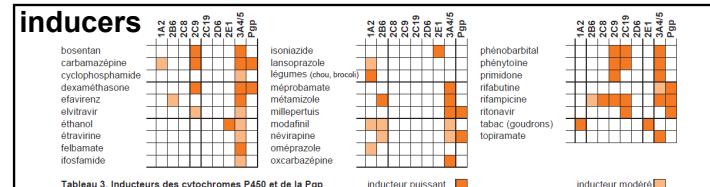
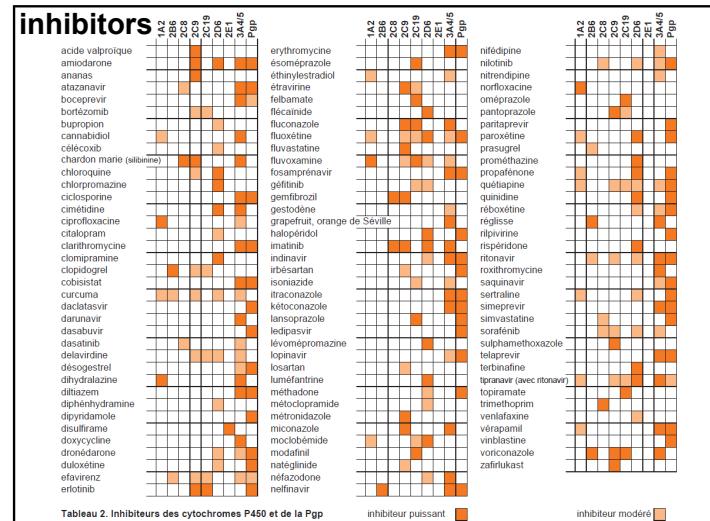
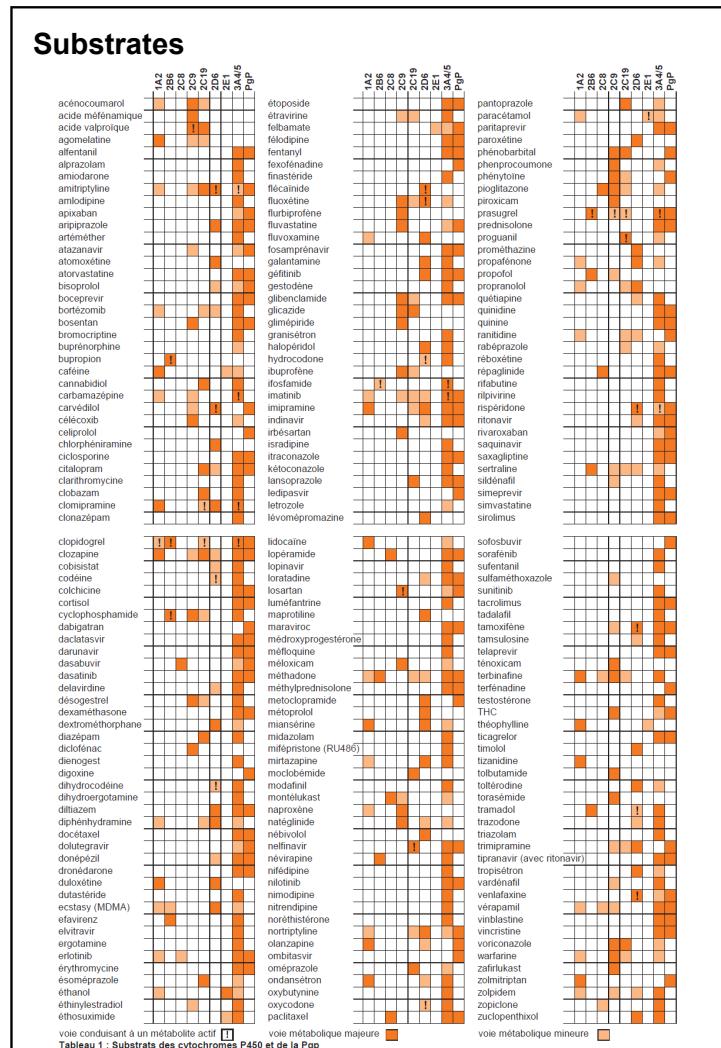


- Opioids DDIs:
www.opioiddruginteractions.com



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Drug – Drug Interactions: Cytochrome Tables



Inhibition

Cytochromes : L'impact dépend de : a) importance relative de la voie d'élimination inhibée par rapport à la clairance totale; b) présence ou non de métabolites actifs et c) concentrations d'inhibiteur. A l'arrêt du traitement inhibiteur, l'activité du CYP revient progressivement à la normale (>2 semaines après disparition de l'inhibiteur dans le sang). Exemple : Le millepertuis induit progressivement et puissamment l'activité du CYP3A4. Il accélère fortement l'élimination de l'éthynodiolésteradiol, substrat majeur du CYP3A4, et l'efficacité contraceptive ne sera plus assurée, il faudra donc prévoir un autre mode de contraception.

P-glycoprotéine : L'impact dépend de l'affinité du substrat pour ce transporteur, de la concentration et de la puissance de l'inhibiteur. Comme pour les CYP, l'activité de la Pgp revient progressivement à la normale à l'arrêt du traitement inhibiteur (>2 semaines après disparition de l'inhibiteur dans le sang). Exemple : La clोzapine inhibe fortement l'activité de la Pgp. Associée à l'ritonavir, substrat de la Pgp, elle entraînera une augmentation de sa biodisponibilité.

Des tableaux dynamiques régulièrement mis à jour et comprenant davantage de molécules sont accessibles sur le site www.pharmacodin.ch, rubrique Centre d'informations thérapeutiques et de pharmacovigilance > outils > carte dynamique des interactions médicamenteuses et CYP.

Centre d'informations thérapeutique et de pharmacovigilance
Service de pharmacologie et toxicologie cliniques, Hôpitaux Universitaires, 1211 Genève 14
Tél. 022 372 89 32 - Fax 022 372 89 45 - www.pharmacodin.ch - Copyright SFTC - Genève, Janvier 2015

Major metabolic pathway

Minor metabolic pathway

Active metabolite



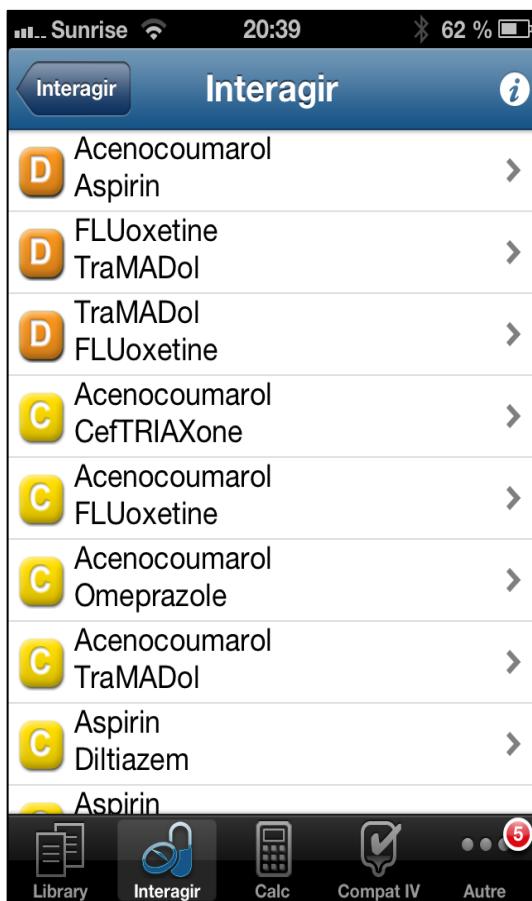
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Extract from:

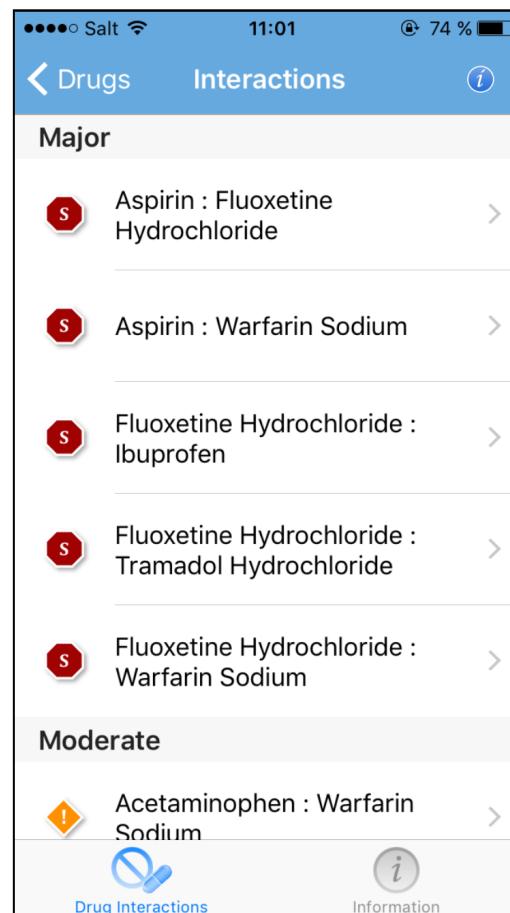
http://www.hug-ge.ch/sites/interhug/files/structures/pharmacologie_et_toxicologie_cliniques/documents/interactions_medicamenteuses_et_cyp450.pdf

Drug – Drug Interactions: Sources

Lexi-Interact



Micromedex Drugs



Epocrates



Drug – Drug Interactions: Software comparisons

	Compendium	Lexi-Interact	Epocrates	Thériaque
INN search	Non	Oui	Oui	Oui
Brand Name search	Oui	Oui (noms commerciaux américains)	Oui (noms commerciaux américains)	Oui (noms commerciaux français)
Levels of risk rating	4 (6)	5	4	4
Colour code	Oui (rouge-orange-jaune)	Non	Oui (rouge-bleu)	Non
Risk Rating	✓	✓	✓	✓
Interaction frequency	X	X	X	X
Interaction mechanism	✓	✓	X	✓
Interaction effect	✓	✓	✓	✓
Clinical management	✓	✓	✓	(✓) → dépend du médicament
References	✓	✓	X	(✓) → seulement Thesaurus
Update	Tous les 14 jours	Continuelle?	Continuelle?	Quotidienne
Smartphone app	X	✓	✓	X

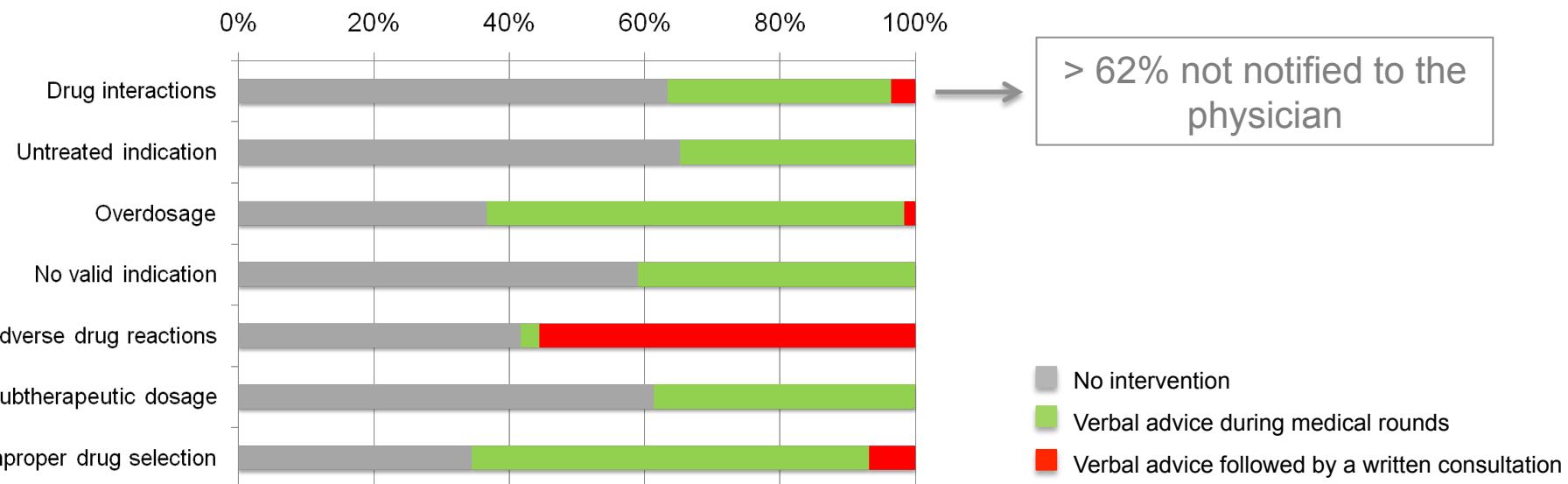
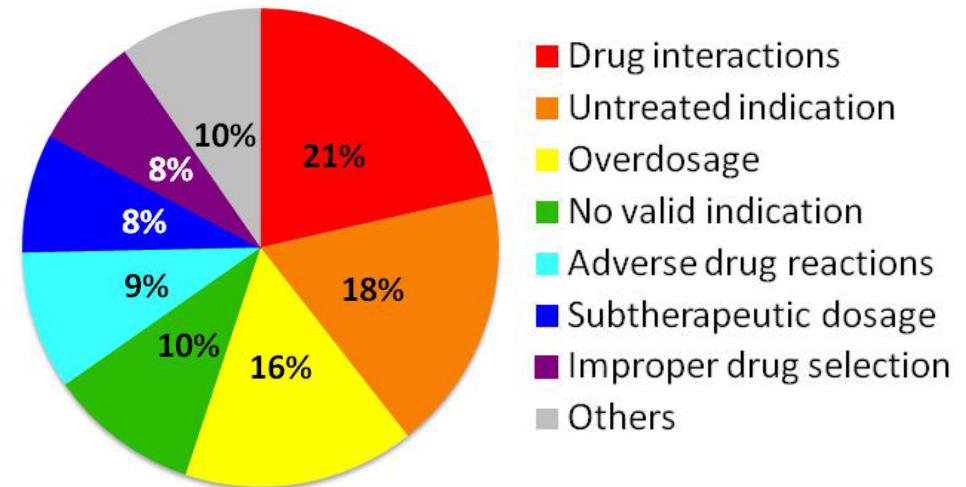
	Compendium suisse des médicaments	Lexi-Interact	Epocrates	Thériaque
True positives	50	59	58	45
False positives	0	0	2	1
True negatives	12	12	10	11
False negatives	12	3	4	17
Sensitivity	80%	95%	94%	73%
Specificity	100%	100%	83%	92%

62 pairs of interactions, 12 pairs without interaction

Adapted from : Ing Lorenzini K et al. Quel programme informatique de détection des interactions médicamenteuses néfastes ? Rev Med Suisse. 2012 Oct 17;8(358):1978-82.

Drug – Drug Interactions: Software limits

6 months
2 internal medicine wards
1 visit per week
145 patients included
69 years old (21-99)
1523 lines of prescription
383 DRPs identified



Lexi-interact

Lexicomp® Lexi-Interact™

Lookup

Enter item name to lookup.

Analyze New List

- [Acenocoumarol](#)
- [Aspirin](#)
- [CefTRIAxone](#)
- [DiltiaZEM CD](#)
- [FLUoxetine](#)
- [Furosemide](#)
- [Ginkgo Biloba](#)
- [HydroCHLORothiazide](#)
- [Ibuprofen](#)
- [Magnesium Sulfate](#)
- [Omeprazole](#)
- [Oxazepam](#)
- [Paracetamol \(INT\)](#)
- [Potassium Chloride](#)
- [TraMADol](#)

- Display complete list of interactions for an individual item by clicking item name.
- Add another item(s) [Lookup] to Analyze for potential interactions between items in the list.
- Remove item from the list by clicking the check mark next to the item name.

Lexi-Comp Online™ Interaction Analysis

Customize Analysis

Only interactions at or above the selected [risk rating](#) will be displayed. A: ▾
View interaction detail by clicking on link.

Acenocoumarol

- [D] [Aspirin](#) (Salicylates)
- [C] [CefTRIAxone](#) (Cephalosporins)
- [C] [FLUoxetine](#) (Agents with Antiplatelet Properties)
- [C] [FLUoxetine](#) (Selective Serotonin Reuptake Inhibitors)
- [D] [Ginkgo Biloba](#) (Ginkgo Biloba)
- [D] [Ibuprofen](#) (NSAID (Nonselective))
- [C] [Omeprazole](#) (Omeprazole)
- [C] [Paracetamol \(INT\)](#) (Acetaminophen)
- [C] [TraMADol](#) (TraMADol)

Aspirin

- [D] [Acenocoumarol](#) (Vitamin K Antagonists)
- [C] [DiltiaZEM CD](#) (Calcium Channel Blockers (Nondihydropyridine))
- [C] [FLUoxetine](#) (Selective Serotonin Reuptake Inhibitors)
- [C] [Furosemide](#) (Loop Diuretics)
- [D] [Ginkgo Biloba](#) (Ginkgo Biloba)
- [D] [Ibuprofen](#) (NSAID (Nonselective))

CefTRIAxone

- [C] [Acenocoumarol](#) (Vitamin K Antagonists)
- DiltiaZEM CD (DiltiaZEM)
- [C] [Aspirin](#) (Salicylates)
- [B] [Ibuprofen](#) (Nonsteroidal Anti-Inflammatory Agents)
- [C] [Magnesium Sulfate](#) (Magnesium Salts)

FLUoxetine

- [C] [Acenocoumarol](#) (Anticoagulants)
- [C] [Acenocoumarol](#) (Vitamin K Antagonists)
- [C] [Aspirin](#) (Aspirin)
- [D] [Ginkgo Biloba](#) (Herbs (Anticoagulant/Antiplatelet Properties))
- [C] [HydroCHLORothiazide](#) (Thiazide and Thiazide-Like Diuretics)
- [D] [Ibuprofen](#) (NSAID (Nonselective))
- [C] [Omeprazole](#) (CYP2C9 Inhibitors (Moderate))
- [C] [Oxazepam](#) (CNS Depressants)
- [C] [TraMADol](#) (TraMADol)
- [C] [TraMADol](#) (TraMADol)

Furosemide

- [C] [Aspirin](#) (Salicylates)
- [D] [Ibuprofen](#) (Nonsteroidal Anti-Inflammatory Agents)
- [C] [TraMADol](#) (Analgesics (Opioid))

Ginkgo Biloba

- [D] [Acenocoumarol](#) (Vitamin K Antagonists)
- [D] [Aspirin](#) (Salicylates)
- [D] [FLUoxetine](#) (Agents with Antiplatelet Properties)
- [B] [HydroCHLORothiazide](#) (Thiazide and Thiazide-Like Diuretics)
- [D] [Ibuprofen](#) (Agents with Antiplatelet Properties)
- [D] [Ibuprofen](#) (Nonsteroidal Anti-Inflammatory Agents)

HydroCHLORothiazide

- [C] [FLUoxetine](#) (Selective Serotonin Reuptake Inhibitors)
- [B] [Ginkgo Biloba](#) (Ginkgo Biloba)
- [C] [Ibuprofen](#) (Nonsteroidal Anti-Inflammatory Agents)
- [C] [TraMADol](#) (Analgesics (Opioid))

Ibuprofen

- [D] [Acenocoumarol](#) (Vitamin K Antagonists)
- [D] [Aspirin](#) (Salicylates)
- [B] [DiltiaZEM CD](#) (Calcium Channel Blockers)
- [D] [FLUoxetine](#) (Selective Serotonin Reuptake Inhibitors)
- [D] [Furosemide](#) (Loop Diuretics)
- [D] [Ginkgo Biloba](#) (Herbs (Anticoagulant/Antiplatelet Properties))
- [D] [Ginkgo Biloba](#) (Herbs (Anticoagulant/Antiplatelet Properties))
- [C] [HydroCHLORothiazide](#) (Thiazide and Thiazide-Like Diuretics)

Magnesium Sulfate

- [C] [DiltiaZEM CD](#) (Calcium Channel Blockers)
- [C] [Oxazepam](#) (CNS Depressants)
- [C] [TraMADol](#) (CNS Depressants)

Omeprazole

- [C] [Acenocoumarol](#) (Vitamin K Antagonists)
- [C] [FLUoxetine](#) (CYP2C9 Substrates)

Oxazepam

- [C] [FLUoxetine](#) (Selective Serotonin Reuptake Inhibitors)
- [C] [Magnesium Sulfate](#) (Magnesium Sulfate)
- [C] [TraMADol](#) (CNS Depressants)

Paracetamol (INT)

- (Acetaminophen)
- [C] [Acenocoumarol](#) (Vitamin K Antagonists)
- [B] [TraMADol](#) (Analgesics (Opioid))

Potassium Chloride

No interactions identified with others in the selection list.

TraMADol

- [C] [Acenocoumarol](#) (Vitamin K Antagonists)
- [C] [FLUoxetine](#) (CYP2D6 Inhibitors (Strong))
- [C] [FLUoxetine](#) (Serotonin Modulators)
- [C] [Furosemide](#) (Diuretics)
- [C] [HydroCHLORothiazide](#) (Diuretics)
- [C] [Magnesium Sulfate](#) (Magnesium Sulfate)
- [C] [Oxazepam](#) (CNS Depressants)
- [B] [Paracetamol \(INT\)](#) (Acetaminophen)



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Micromedex Drugs Interactions

Drugs Interactions ⓘ

Major

- S** Aspirin : Fluoxetine Hydrochloride >
- S** Aspirin : Warfarin Sodium >
- S** Fluoxetine Hydrochloride : Ibuprofen >
- S** Fluoxetine Hydrochloride : Tramadol Hydrochloride >
- S** Fluoxetine Hydrochloride : Warfarin Sodium >

Moderate

- !** Acetaminophen : Warfarin Sodium >

 Drug Interactions  Information

Fluoxetine + tramadol

Interactions Interaction

Fluoxetine Hydrochloride : Tramadol Hydrochloride

Severity: Major
Onset: Rapid
Documentation: Good

Interaction Effect:
Concurrent use of FLUOXETINE and TRAMADOL may result in an increased risk of seizures, opioid toxicity, and serotonin syndrome (hypertension, hyperthermia, myoclonus, mental status changes), and increased concentrations of tramadol and decreased concentrations of tramadol active metabolite, M1.

Clinical Management:
Use caution with concomitant use of fluoxetine and tramadol. Concomitant use of tramadol with SSRIs, serotonin norepinephrine reuptake inhibitors

 Drug Interactions  Information

Aspirin + ibuprofen

Interactions Interaction

Aspirin : Ibuprofen

Severity: Moderate
Onset: Not Specified
Documentation: Good

Interaction Effect:
Concurrent use of ASPIRIN and IBUPROFEN may result in decreased antiplatelet effect of aspirin and additive risk of bleeding.

Clinical Management:
Ibuprofen can interfere with the irreversible platelet inhibition effects of aspirin via competition at the platelet cyclooxygenase-1 (COX-1) binding site. This interaction may increase the risk of adverse thrombotic events in patients who receive daily low-dose aspirin as preventive therapy. If regular combined

 Drug Interactions  Information



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Drug – Drug Interactions

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole 40 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- Diltiazem extended-release 90 mg BID
- Fluoxetine 20 mg QD
- Furosemide 20 mg QD
- Oxazepam 15 mg QD
- OTC : paracetamol, **ibuprofen**

► Added at admission

- KCl 40 mmol TID IV 1h-infusion (solvent: NaCl 0.9% 250 ml)
- Magnesium sulfate 2 g IV continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- **~~Tramadol 25 mg QID PO, then 50 mg QID, poorly effective~~**
- **Buprenorphine 0.1 mg TID + 0.1 mg BID if needed**

DOSES

Doses: Sources

► Software and Apps



- ▶ Lexi Comp's: www.lexi.com
- ▶ Micromedex (Drugdex): www.micromedexsolutions.com
- ▶ European Medicines Agency: www.ema.europa.eu
- ▶ U.S. Food and Drug Administration: www.accessdata.fda.gov/scripts/cder/drugsatfda/index.cfm
- ▶ National drug databases, SmPCs

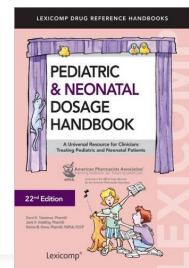
► Guidelines

- ▶ National and international scientific societies and regulatory agencies



► Specific tools

- ▶ Renal failure
ICAR : www.sitegrpr.com
The renal drug database:
renaldrugdatabase.com
- ▶ Pediatrics:
Pediatric & Neonatal dosage handbook (Lexicomp) www.lexi.com



Doses?

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole 40 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- Diltiazem extended-release 90 mg BID
- Fluoxetine 20 mg QD
- Furosemide 20 mg QD
- Oxazepam 15 mg QD
- OTC : paracetamol

► Added at admission

- KCl 40 mmol TID IV 1h-infusion (solvent: NaCl 0.9% 250 ml)
- Magnesium sulfate 2 g IV continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- Buprenorphine 0.1 mg TID + 0.1 mg BID if needed

Doses?

Omeprazole

Omeprazole: Drug information Lexicomp®

[Access Lexicomp Online here.](#)

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(For additional information [see "Omeprazole: Patient drug information"](#) and [see "Omeprazole: Pediatric drug information"](#))

NSAID-induced ulcer prophylaxis (off-label use): Oral: 20 mg once daily for up to 6 months
(Cullen 1998)

Furosemide

Table 7.3 Doses of diuretics commonly used in patients with heart failure

Diuretics	Initial dose (mg)	Usual daily dose (mg)		
Loop diuretics^a				
Furosemide	20–40	40–240		
Bumetanide	0.5–1.0	1–5		
Torsemide	5–10	10–20		
Thiazides^b				
Bendroflumethiazide	2.5	2.5–10		
Hydrochlorothiazide	25	12.5–100		
Metolazone	2.5	2.5–10		
Indapamide ^c	2.5	2.5–5		
Potassium-sparing diuretics^d				
	+ACE-I/ ARB	-ACE-I/ ARB	+ACE-I/ ARB	-ACE-I/ ARB
Spironolactone/ eplerenone	12.5–25	50	50	100– 200
Amiloride	2.5	5	5–10	10–20
Triamterene	25	50	100	200

ACE-I = angiotensin-converting enzyme inhibitor, ARB = angiotensin receptor blocker.

^aOral or intravenous; dose might need to be adjusted according to volume status/weight; excessive doses may cause renal impairment and ototoxicity.

^bDo not use thiazides if estimated glomerular filtration rate <30 mL/min/1.73 m², except when prescribed synergistically with loop diuretics.

^cIndapamide is a non-thiazide sulfonamide.

^dA mineralocorticoid antagonist (MRA) i.e. spironolactone/eplerenone is always preferred. Amiloride and triamterene should not be combined with an MRA.



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Ponikowski P, et al. 2016 ESC Guidelines for the diagnosis and treatment of acute and chronic heart failure European Heart Journal (2016) 37, 2129–2200

Doses?

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole **40 mg 20 mg** QD
- Acenocoumarol QD dose-adjusted according to INR
- Diltiazem extended-release 90 mg BID
- Fluoxetine 20 mg QD
- Furosemide **20 mg 40 mg** QD
- Oxazepam 15 mg QD
- OTC : paracetamol

► Added at admission

- KCl 40 mmol TID IV 1h-infusion (solvent: NaCl 0.9% 250 ml)
- Magnesium sulfate 2 g IV continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- Buprenorphine 0.2 mg TID + 0.1 mg TID if needed

DRUG ADMINISTRATION

Drug administration?

- ▶ Parenteral administration
 - ▶ Which solvent (compatibility) ?
 - ▶ Which concentration (vascular irritation) ?
 - ▶ Which infusion rate (systemic intolerance) ?
 - ▶ Y-site compatibility ?
- ▶ Switch IV – PO
- ▶ Oral and enteral route
 - ▶ Scored/unscored tablets
 - ▶ Crush/chew tablets/ Open capsules ?
 - ▶ Oral suspension available
 - ▶ Administration through Naso/gastric tube

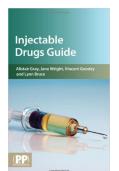
Drug administration: some sources

- ▶ Software and Apps
- ▶ Previous drug databases
 - ▶ Lexi Comp's: www.lexi.com
 - ▶ Micromedex: www.micromedexsolutions.com

Books



- ▶ Injectable Medicines administration guide
- ▶ Injectable Drugs Guide
ssu.ac.ir/cms/fileadmin/user_upload/bimarestanha/shahid_sadooghi/paraclinic/daokhane/file/Injectable_Drugs_Guide.pdf



- ▶ Handbook of Drug Administration via Enteral Feeding Tubes
www.pharmacy.cmu.ac.th/unit/unit_files/files_download/2012-03-26HandbkOfDrugAdminiViaEnteralFeedingTubes%201stEd_WhiteAndBradn.pdf



Drug administration?

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole 20 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- Diltiazem extended-release 90 mg BID
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PARENTERAL ADMINISTRATION: INJECTABLE DRUGS GUIDE

Potassium chloride

The UK NPSA Patient Safety Alert (Oct 2002)

Potassium chloride concentrate solution can be fatal if given inappropriately, therefore:

- Commercially prepared ready to use dilute solutions containing potassium chloride should be used wherever possible.
- If a suitable solution is not available commercially, dilutions should be made in the hospital pharmacy wherever possible.
- Storage of concentrated potassium chloride solutions should be restricted to pharmacy departments and to those critical care areas where concentrated solutions are needed for urgent use.
- Receipt and use of these solutions should be recorded in a similar way to Controlled Drugs and stocks should be kept in a separate locked cupboard away from commonly used diluting solutions.
- Potassium chloride concentrate solution should not be transferred between clinical areas.

Dose

Treatment of hypokalaemia: the dose is dependent upon the biochemistry and clinical condition of the patient.

Mild to moderate ↓K ($K > 2.5 \text{ mmol/L}$): oral KCl supplements are usually adequate. If the oral route is temporarily unavailable, a single dose of 20–40 mmol by IV infusion over 6–8 hours may be sufficient to treat acute deficiency.

Severe ↓K ($K < 2.5 \text{ mmol/L}$): 20–40 mmol (as a 40 mmol/L solution) by IV infusion at a maximum rate of 20 mmol/hour (10 mmol/hour is the usual maximum in a general ward area) governed by the clinical status and hydration of the patient. Repeat as necessary according to biochemistry results.

More concentrated solutions may be given in a critical care area with ECG monitoring, particularly where fluid overload is problematic.

NB: There is a high risk of asystole if the administration rate reaches 40 mmol/hour.



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Drug administration?

► Medication history

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- Hydrochlorothiazide 25 mg QD
- Omeprazole 20 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- Diltiazem extended-release 90 mg BID
- Fluoxetine 20 mg QD
- Furosemide 40 mg QD
- Oxazepam 15 mg QD
- OTC : paracetamol

► Added at admission

- KCl 40 mmol TID once IV 1h infusion over 6-8 h (solvent: NaCl 0.9% 250 ml) then 16 mmol TID PO
- Magnesium sulfate 2 g IV-continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
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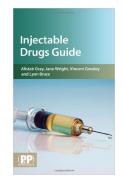
Drug administration: some sources

- ▶ Software and Apps
- ▶ Previous drug databases
 - ▶ Lexi Comp's: www.lexi.com
 - ▶ Micromedex: www.micromedexsolutions.com

Books



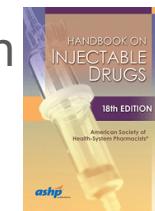
- ▶ Injectable Medicines administration guide
- ▶ Injectable Drugs Guide
ssu.ac.ir/cms/fileadmin/user_upload/bimarestanha/shahid_sadooghi/paraclinic/daokhane/file/Injectable_Drugs_Guide.pdf
- ▶ Handbook of Drug Administration via Enteral Feeding Tubes
www.pharmacy.cmu.ac.th/unit/unit_files/files_download/2012-03-26HandbkOfDrugAdminiViaEnteralFeedingTubes%201stEd_WhiteAndBradn.pdf



- ▶ Compatibility
- ▶ Software and Apps
 - ▶ King Guide : www.kingguide.com
 - ▶ Micromedex IV Compatibility: www.micromedexsolutions.com

Books

- ▶ Trissel's Handbook on Injectable Drugs
- ▶ Neofax



Tables

- ▶ CHUV table
files.chuv.ch/internet-docs/pha/medicaments/pha_phatab_compatibilitesy.pdf
- ▶ HUG table
pharmacie.hug-ge.ch/infomedic/utilismedic/HUG_CompatAdm_DC1.pdf

Compatibility: CHUV tables

CHUV

COMPATIBILITES DES MEDICAMENTS INJECTABLES ADMINISTRES EN Y - 05.2012 - v.6.2

Code couleur :

- Vert = compatible (tests physiques ou physico-chimiques)
- Rouge = incompatible
- Jaune = à vérifier aux notes ci-dessous
- Blanc = aucune donnée publique (contacter la hotline)
- Grise = référence employée:

Compatibility:

PERALGAN : ne pas mélanger avec d'autres médicaments (compatible avec: NaCl 0.9%, Dornic 1 mg/ml, Morphine 1 mg/ml, Potassium Chlorure 0.05 mmol/ml)*

1 Acrapid : absorption sur le verre et le plastique (au-delà 20-30% de perds, surtout lors de dilutions importantes, puis saturation des sites d'absorption).

2 ATTENTION : le calcium est incompatible avec les phosphates (et les préparations qui en contiennent) en fonction des concentrations.

3 Les TPN avec lipides sont incompatibles avec des solutions d'électrolytes concentrées (hors de l'utilisation du TPN, surtout en présence d'héparine).

4 Cefazidime: compatible avec dilutionne selon les concentrations de diluantes utilisées, se référer à la pharmacie.

5 Ciprofloxacin (ciprofloxacin): incompatible avec les solutions à pH acide.

6 Carbopenem (amikacine) : absorption sur les tubules en PVC.

Risque de précipitation si concentration < 0,6 mg/ml.

7 Flouxapen (flucloxacilline) compatible avec héparine <0,01M, au delà pas de données.

8 Flouxapen (flucloxacilline) compatible avec tamoxifène <0,5mg/ml, au delà pas de données.

9 Invers (antipyrine): compatible en Y avec potassium chlorure dans NaCl 0.9% uniquement.

10 Compatible avec diluants suivants: morphine 0,2mg/ml, sodium bicarbonate 2,1mg/ml.

11 Morphine : compatible en Y avec des TPN contenant des lipides si la concentration de morphine est < 1 mg/ml.

12 Nipras (nifuropratate sodique): doit être protégé de la lumière.

13 Phenylén (phénylénol): si possible administrer seul. Phenylén concentré pour perfusion 750 mg/50 ml compatible avec NaCl 0.9%, G10, Glucosin, Ringer-Lactate. Phenylén solution injectable 250 mg/5ml compatible uniquement avec NaCl 0.9%.

14 ATTENTION : le potassium dihydrophosphate est incompatible avec le calcium (et les préparations qui en contiennent) en fonction des concentrations.

15 Rocophine (ceftriaxone) : incompatible avec le calcium (et les préparations qui en contiennent). La Rocophine doit être si possible administrée seule.

16 Sodium bicarbonate : incompatible avec les acids (élimination de CO₂), précipitation de carbonates avec Ca et Mg.

Notes:

Les données de compatibilité de ce tableau sont tirées de tests en laboratoire de médicaments en association par deux; il existe très rarement des associations de plus de deux médicaments.

En cas de doutes ou de questions, veuillez appeler le (079 55 67 380 (Hotline Assistance pharmaceutique).

La responsabilité du CHUV n'est pas engagée lors de l'emploi de ce tableau hors de l'institution.

Compatibility: tools evaluation



- **Evaluation by 2 pharmacists**
- **40 pairs frequently used in pediatrics intensive care units**
- **Trissel's handbook = gold standard**

Table 4 Tool-evaluation summary

Tool	Accuracy score ^a	Completeness score ^a	Comprehensiveness score ^a	Applicability score ^a	Global score ^a
Ref	250	250	250	250	1000
Thé	234	200	218	188	840
pH	175	200	134	298	807
CHUV	213	150	174	266	803
Perf	230	138	218	191	776
NeoF	190	181	116	191	678
King	192	131	108	211	642
Stab	179	144	149	112	584
KIK	105	156	157	105	523

Ref Trissel's Handbook

Thé Thériaque database, *Perf* Perfysi database, *CHUV* CHUV's cross-table *King* King cross-table wall chart, *NeoF* Neofax handbook, *Stab* Stabilis database, *pH* pH cross-table, *KIK* software

Incompatibility

Micromedex IV Compatibility

Y-Site Test Detail			Rating
Ceftriaxone sodium - Magnesium sulfate		-	Not Tested
Ceftriaxone sodium - Potassium chloride		✓	Compatible
Magnesium sulfate - Potassium chloride		✓	Compatible

IV COMPATIBILITY DETAIL				
Drug 1	Drug 2	Status	Information	Test Parameters
Ceftriaxone sodium 165mg/mL in' D5W-Dextrose 5% Roche Pharmaceuticals	Potassium chloride 1mEq/mL in' D5W-Dextrose 5% Abbott Laboratories	✓ Compatible	<p>Physical Compatibility: Physically compatible. No changes in measured haze or turbidity, particulates, or color were found.</p> <p>Storage: Ambient room temperature near 23 °C exposed to normal fluorescent light.</p> <p>Container: Simulated Y-site administration using glass test tubes.</p>	Reference: 8869 Study Period: 4 hours. Method: Visual observation and electronic assessment.
Ceftriaxone sodium 165mg/mL in' Normal saline- Sodium chloride 0.9% Roche Pharmaceuticals	Potassium chloride 1mEq/mL in' Normal saline- Sodium chloride 0.9% Abbott Laboratories	✓ Compatible	<p>Physical Compatibility: Physically compatible. No changes in measured haze or turbidity, particulates, or color were found.</p> <p>Storage: Ambient room temperature near 23 °C exposed to normal fluorescent light.</p> <p>Container: Simulated Y-site administration using glass test tubes.</p>	Reference: 8869 Study Period: 4 hours. Method: Visual observation and electronic assessment.

King Guide

Search IV Drugs (up to 10):

Search for another drug to add it to the multiple drug compatibility results.

Click any drug to remove it from the results.

C=Compatible X=Incompatible Ø=Conflicting Reports NA=No Data

Click any symbol to view the fluids, dosages, and details. Click any drug to view the complete monograph for that drug.

Y-Site			CEFTRIAXONE SODIUM	MAGNESIUM SULFATE	POTASSIUM CHLORIDE
CEFTRIAXONE SODIUM	X	Ø			
MAGNESIUM SULFATE	X		C		
POTASSIUM CHLORIDE	Ø	C			

Syringe			CEFTRIAXONE SODIUM	MAGNESIUM SULFATE	POTASSIUM CHLORIDE
CEFTRIAXONE SODIUM	X	X			
MAGNESIUM SULFATE	X			NA	
POTASSIUM CHLORIDE	X	NA			

Admixture			CEFTRIAXONE SODIUM	MAGNESIUM SULFATE	POTASSIUM CHLORIDE
CEFTRIAXONE SODIUM	X	Ø			
MAGNESIUM SULFATE	X		C		
POTASSIUM CHLORIDE	Ø	C			

Caution: The compatibility of two or more drugs in the same IV line/container should not be inferred from information presented here, unless data on that specific combination has actually been reported within the text of the monograph.

Drug administration?

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole 20 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- Diltiazem extended-release 90 mg BID
- Fluoxetine 20 mg QD
- Furosemide 40 mg QD
- Oxazepam 15 mg QD
- OTC : paracetamol

► Added at admission

- KCl 40 mmol once IV infusion over 6-8 h (solvent: NaCl 0.9% 500 ml) then 16 mmol TID PO
- Magnesium sulfate 2 g IV-continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- Buprenorphine 0.2 mg TID + 0.1 mg TID if needed

**2 different veins or rinse the infusion line between administration
and as soon as the clinical condition allows it, switch antibiotic IV-PO**



ADVERSE DRUG REACTIONS

Adverse drug reactions: some sources

► Software and Apps



- Lexi Comp's: www.lexi.com
- Micromedex (Drugdex): www.micromedexsolutions.com
- European Medicines Agency: www.ema.europa.eu
- U.S. Food and Drug Administration: www.accessdata.fda.gov/scripts/cder/drugsatfda/index.cfm
- National drug databases, SmPCs



► Hepatic disorders



LiverTox

- LiverTox: livertox.nlm.nih.gov



► G6PD deficiency

- G6PD Deficiency favism association: www.g6pd.org/en/G6PDDeficiency/SafeUnsafe/DaEvitare_ISS-it



► QT prolongation

- CredibleMeds: crediblemeds.org



► Combination of apps

- PIM-Check: pimcheck.org



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Adverse drug reactions: key symptoms

- ▶ Hepatic disorders (abnormal liver function tests)
- ▶ Hematologic disorders (Neutropenia/ Agranulocytosis, leukopenia or lymphopenia, haemolytic anaemia, thrombocytopenia)
- ▶ Haemorrhage
- ▶ Skin reactions
- ▶ Anaphylactic reactions
- ▶ Acute renal failure
- ▶ Digestive disorders (dysphagia, epigastric pain, constipation, diarrhoea)
- ▶ Cough
- ▶ Hypo/Hyperglycaemia
- ▶ Electrolyte disorders (hyponatremia, hypo/hyperkalemia, hypocalcemia ...)
- ▶ Neurologic or musculoskeletal disorders (dizziness, seizure, myalgia)
- ▶ Cardiovascular disorders (cardiac rhythm disorders, heart failure, hypo/hypertension)

Adverse drug reactions?

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole 20 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- Diltiazem extended-release 90 mg BID
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- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- Buprenorphine 0.1 mg TID + 0.1 mg BID if needed

INAPPROPRIATE PRESCRIBING

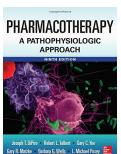
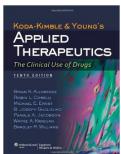
**Under-prescriptions
Over-prescriptions
Mis-prescriptions**

Inappropriate prescribing: sources

► Guidelines

- Uptodate: www.uptodate.com

► Books



- Koda-Kimble & Young's Applied Therapeutics: The Clinical Use of Drugs
- DiPiro Pharmacotherapy : A Pathophysiologic Approach. 2014
- National and international scientific societies and regulatory agencies

► Explicit criteria

- STOPP/START version 2:

Screening tool of older people's prescriptions / Screening tool to alert to right treatment

O'Mahony D et al. STOPP/START criteria for potentially inappropriate prescribing in older people:version 2. Age Ageing. 2015;44:213-8. PMID: 25324330

- PIM-Check:

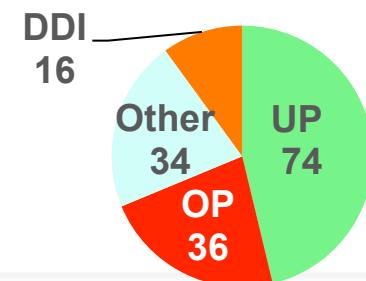
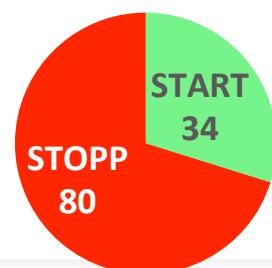
Potentially inappropriate prescribing for patients in internal medicine checklist

pimcheck.org

Desnoyer et al. PIM-Check: An International Prescription-Screening Checklist Designed by a Delphi Method for Adult Internal Medicine Patients. In revision

Explicit criteria: STOPP/START vs PIM-Check

	STOPP/START version 2	PIM-Check
▶ Explicit criteria validated using a 2 round web-based Delphi method involving	19 experts in geriatrics from 13 european countries	39 experts in internal medicine from Be, Fr, QC, CH
▶ Target	Older people	Adult in internal medicine unit
▶ Number of items	114 items	160 items
▶ Types of PIM	Over- (STOPP) /under- (START) prescriptions	Over- (OP) / under- (UP) / mis-prescriptions (Other) / drug-drug interactions (DDI)



Explicit criteria: STOPP/START vs PIM-Check

STOPP/START version 2

Classification

Classified according to pathophysiological systems

Presentation

2 tables (paper)

Appendix 3: Screening Tool of Older Persons' Prescriptions (STOPP) version 2.

The following prescriptions are potentially inappropriate to use in patients aged 65 years and older.

Section A: Indication of medication

1. Any drug prescribed without an evidence-based clinical indication.
2. Any drug prescribed beyond the recommended duration, where treatment duration is well defined.
3. Any duplicate drug class prescription e.g. two concurrent NSAIDs, SSRIs, loop diuretics, ACE inhibitors, anticoagulants (optimisation of monotherapy within a single drug class should be observed prior to considering a new agent).

Section B: Cardiovascular System

1. ~~Digoxin~~ for heart failure with normal systolic ventricular function (no clear evidence of benefit)
2. ~~Verapamil~~ or ~~diltiazem~~ with NYHA Class III or IV heart failure (may worsen heart failure).
3. Beta-blocker in combination with ~~verapamil~~ or ~~diltiazem~~ (risk of heart block).
4. Beta-blocker with bradycardia (< 50/min), type II heart block or complete heart block (risk of complete heart block, asystole).
5. ~~Amiodarone~~ as first-line antiarrhythmic therapy in supraventricular tachyarrhythmias (higher risk of side-effects than beta-blockers, digoxin, verapamil or diltiazem)

PIM-Check

Website and Webmobile application



Screening

Favorites

Items list

PIMcheck.org

Display ALL items Display ALREADY read items Display UNREAD items



ALL items

CARDIOLOGY

Heart failure

UP Heart failure: Start ACEI or ARB

UP Heart failure: Start beta-blocker* treatment

UP Heart failure: Start aldosterone antagonist when LVEF ≤ 35% despite optimal treatment

OP Heart failure: Drugs that may exacerbate HF

Dyslipidaemia and hypolipidemics

UP Dyslipidaemia and high cardiovascular risk: start statins



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Explicit criteria: STOPP/START vs PIM-Check

	STOPP/START version 2	PIM-Check
Medical specialties, pathologies and domains included		
► Cardiovascular System	✓	✓
► Coagulation System	✓	✓
► CNS and Psychiatry	✓	✓
► Renal/ urogenital System	✓	✓
► Gastrointestinal System	✓	✓
► Respiratory System	✓	✓
► Musculoskeletal System	✓	✓
► Urogenital System	✓	✓
► Endocrine System	✓	✓
► Pain and Analgesic Drugs	✓	✓
► Ophthalmology	✓	✓
► Vaccination	✓	✓
► Other	<ul style="list-style-type: none">► Drug indication► Drugs with increased risk of falls in older people► Antimuscarinic/ anticholinergic drugs	<ul style="list-style-type: none">► Dependencies► Infectious diseases► Obesity► Pharmacology & toxicology► Transplantation

Inappropriate prescribing?

► Medication history

- Aspirin 100 mg QD
- Hydrochlorothiazide 25 mg QD
- Omeprazole 20 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- Diltiazem extended-release 90 mg BID
- Fluoxetine 20 mg QD
- Furosemide 40 mg QD
- Oxazepam 15 mg QD
- OTC : paracetamol

► Added at admission

- KCl 40 mmol TID IV 1h-infusion (solvent: NaCl 0.9% 500 ml) then 16 mmol TID PO
- Magnesium sulfate 2 g IV continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- Buprenorphine 0.1 mg TID + 0.1 mg BID if needed

Over-prescriptions

STOPP criteria

Section B: Cardiovascular System criteria

2. Verapamil or diltiazem with NYHA Class III or IV heart failure (may worsen heart failure)

Section C: Antiplatelet/Anticoagulant Drugs

5. Aspirin in combination with vitamin K antagonist, direct thrombin inhibitor or factor Xa inhibitors in patients with chronic atrial fibrillation (no added benefit from aspirin)

PIM-Check over-prescription criteria

- OP Heart failure: Drugs that may exacerbate HF
triggering factors: HEART FAILURE + Diltiazem
- OP Proper use of antibiotics: reevaluate the duration of therapy
triggering factors: ceftriaxone
- OP QT prolongation: drugs that prolong the QT interval
triggering factors: STEMI, NSTEMI, Heart failure, Hypokaliémia + fluoxetine + furosemide + hydrochlorothiazide

Under-prescriptions

START criteria

Section A: Cardiovascular System criteria

5. Statin therapy with a documented history of coronary, cerebral or peripheral vascular disease, unless the patient's status is end-of-life or age is > 85 years.
6. Angiotensin Converting Enzyme (ACE) inhibitor with systolic heart failure and/or documented coronary artery disease.
7. Beta-blocker with ischaemic heart disease.
8. Appropriate beta-blocker (bisoprolol, nebivolol, metoprolol or carvedilol) with stable systolic heart failure.

Section H: Analgesics

2. Laxatives in patients receiving opioids regularly.

Section I: Vaccines

1. Seasonal trivalent influenza vaccine annually
2. Pneumococcal vaccine at least once after age 65 according to national guidelines

PIM-Check under-prescription criteria

- UP Heart failure: Start ACEI or ARB
 - UP Heart failure: Start beta-blocker* treatment
 - UP STEMI / NSTEMI Secondary prevention: Start statins
 - UP STEMI / NSTEMI Secondary prevention: Start ACEI or ARB
 - UP RF: Calcium, vitamin D and/or phosphate-binding agents
 - UP Opioids: start prophylactic measures* to prevent constipation
 - UP Annual influenza vaccination
 - UP Pneumococcal vaccination: high-risk patients*
- triggering factors: Heart failure, STEMI/NSTEMI, Chronic kidney disease

Other and drug interaction

PIM-Check other criteria

OTH

HBP: Favour ACEI or ARB in patients with diabetes/CKD/HF/STEMI/NSTEMI and HBP

OTH

Proper use of antibiotics: reevaluate the route of administration

PIM-Check DDI criteria

DDI

Anticoagulation and DDI

triggering factors: acenocoumarol + diltiazem, omeprazole, ceftriaxone, fluoxetine

DDI

DDI: strong enzyme inducers and inhibitors*

triggering factors: diltiazem (inh CYP3A4, Pgp), fluoxetine (inh CYP2D6, Pgp)

Stop : Aspirin, diltiazem, hydrochlorothiazide

Start: Betablocker, ACEI, Statin, Calcium, Vitamin D, Phosphate biding agent (according to phosphate plasma concentrations), Laxative

Switch: antibiotic with good oral bioavailability (according to antibiotic resistance testing : ciprofloxacin?)

DDI: strengthen INR and ECG monitoring



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Inappropriate prescribing?

► Medication history

- ~~Aspirin 100 mg QD~~
- ~~Hydrochlorothiazide 25 mg QD~~
- Omeprazole 20 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- ~~Diltiazem extended-release 90 mg BID~~
- Fluoxetine 20 mg QD
- Furosemide 40 mg QD
- Oxazepam 15 mg QD
- **Calcium / colecalciferol 1000/800 QD**
- **Bisoprolol 1.25 mg QD (starting dose)**
- **Enalapril 2.5 mg BID (starting dose)**
- **Pravastatin 20 mg QD**
- **Macrogol 3350 QD**
- OTC : paracetamol

► Added at admission

- KCl 40 mmol TID IV 1h-infusion (solvent: NaCl 0.9% 250 ml) then 16 mmol TID PO
- Magnesium sulfate 2 g IV continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- Buprenorphine 0.1 mg TID + 0.1 mg BID if needed

And now?

► Medication history

- Omeprazole 20 mg QD
- Acenocoumarol QD dose-adjusted according to INR
- Fluoxetine 20 mg QD
- Furosemide 40 mg QD
- Oxazepam 15 mg QD
- Calcium / colecalciferol 1000/800 QD
- Bisoprolol 1.25 mg QD (starting dose)
- Enalapril 2.5 mg BID (starting dose)
- Pravastatin 20 mg QD
- Macrogol 3350 QD
- OTC : paracetamol

► Added at admission

- KCl 40 mmol TID IV 1h-infusion (solvent: NaCl 0.9% 250 ml) then 16 mmol TID PO
- Magnesium sulfate 2 g IV continuous infusion (solvent: NaCl 0.9% 250 ml)
- Ceftriaxone 2 g IV infusion over 30 min (solvent: NaCl 0.9% 100 ml)
- Buprenorphine 0.1 mg TID + 0.1 mg BID if needed



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You need to start again the approach

- ▶ Drug – Drug Interactions
- ▶ Doses
- ▶ Drug administration
- ▶ Adverse drug reactions
- ▶ Inappropriate prescribing:
 - ▶ Under-prescriptions
 - ▶ Over-prescriptions
 - ▶ Mis-prescriptions



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Take Home Messages

- ▶ **To detect DRPs you need to have:**
 - ▶ Clinical and therapeutic knowledge (e.g.: guidelines, interaction, dose,...)
 - ▶ Some clinical experience
- ▶ **But most of all:**
 - ▶ To be responsive
 - ▶ To know how to use appropriate tools



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THANK YOU FOR YOUR ATTENTION



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