Clinical services with benefits: medicines optimisation



Conflict of interest: We have nothing to disclose



Goals for the workshop

Teaching goals

- 1. To introduce methods used in medicines optimisation
- To discuss the effects of medicines optimisation and how to measure these effects
- To show the advantages and disadvantages of applying these different tools

Learning objectives

After the workshop the participant should be able to:

- describe medicines optimisation and its effects
- evaluate different tools used in medicines optimisation

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- 1. Are you able to describe what medicines optimisation is and why it should be done? YES / NO
- 2. Can you describe why a medication reconciliation should be performed? YES / NO
- 3. Can you mention two tools used for evaluation of the quality of prescribing? YES / NO

Outline of workshop

Brief introduction

Medication reconciliation

- the what's, why's, when's, who's and how's
- practical example

Medication review

- to perform a medication review
- to assess the effects of a medication review

Discussion & summary

Medicines optimisation



"A person-centered approach to safe and effective medicines use, to ensure people obtain the best possible outcomes from their medicines."



Today we are going to focus on two of the tools for medicines optimisation



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Discussion & summary

A patient typically moves in-between different levels of health-care



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Many medication lists are not in accordance with what patients are actually taking



Inappropriate use Adverse drug reactions Suboptimal drug therapy Prolonged hospitalisation Re-hospitalisations Deaths

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Medication reconciliation



WHAT's

Medication reconciliation – definition

«The process of creating the most accurate list possible of all medications a patient is taking — including drug name, dosage, frequency, and route — and comparing that list against the physician's admission, transfer, and/or discharge orders, with the goal of providing correct medications to the patient at all transition points within the hospital»

International Healthcare Institute (IHI)

http://www.ihi.org/topics/adesmedicationreconcili ation/Pages/default.aspx



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WHAT CAN GO WRONG?



Fig. 2 Medication discrepancies identified by pharmacists (n=235 77% 180 in 99 patients) and nurses (n=222 identified medication discrepancies, n in 94 patients) * P-value for cells 160 63% < 5 has not been calculated. PG; 140 pharmacist group, NG; nurse 120 group 100 80 60 11%9% 40 8%10% 9% 20 3%1% 3%0% 2% 3%1% HORE EX16Z 0 Wrong Comitted Wrong Wrong Wrong Omitted Wrong dosage drug drug strenght generic dosage dosage interval (P=0,002) (P=0,005) time* (NS) name* form* (NS) 149 21 19 7 27 6 6 NG 170 20 6 23 2 1 0

> Aag T, Garcia BH, Viktil K Eur J Clin Pharmacol (2014) 70:1325–1332



Medicine Reconciliation A Practice Guide



When should it happen?

Medicines should be reconciled at the transfer of care between different settings e.g. hospital admission (planned and emergency)

hospital discharge

Movement between settings step up step down and ward/department transfer Entry into residential/nursing care

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A patient typically moves in-between different levels of health-care





Medicine Reconciliation A Practice Guide



When should it happen?

Medicines should be reconciled at the transfer of care between different settings e.g. hospital admission (planned and emergency)

hospital discharge Movement between settings step up step down and ward/department transfer Entry into residential/nursing care

1.3.1 In an acute setting, accurately list all of the person's medicines (including prescribed, over-the-counter and complementary medicines) and carry out medicines reconciliation within 24 hours or sooner if clinically necessary, when the person moves from one care setting to another – for example, if they are admitted to hospital.

NICE guideline Published: 4 March₂2015 nice.org.uk/guidance/ng5



Medicine Reconciliatio A Practice Guide



Who Should Carry Out Medicine Reconciliation?

The responsibility for medicine reconciliation rests with all individuals involved with the transfer of care between different settings.





Physician







Pharmacist technician





METHODOLOGY

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Top of Medication reconciliation form – LIMM model

Depart	ment	Bed	l Name						Date of	birth
		-			Medicat	ion recon	ciliati	on form (Iı	itegrated	Medicin
Inkl.nr			Rnd.nr:		Gender	Age	🛛 Int	erview	Performed	l (date, sign
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	administers medic I □Partly	ation h	im/herself		Multidosage di N V, dat Pill organizer f	·	-	e: ON OY		Dosages*
Date IN 1	Medication, adm. 1	Porm, s	trenght	Dosage	Comments			Discontinuation date	Р	Ph
-> N	Metoprolol,	depo	t 100 mg	1-0-0-0	P: Not sure ab Ph: 50 mg	out dosage			1-0-0-0	1-0-0-0
$\overline{}$		~		~~	$\overline{\mathbf{x}}$			\sim	$\overline{}$	\sim

"Infomation from: patient (P), next of him (NC), general practitioner (GP), specialist (S), Community health cure (CH), multidosage dispensing plasmacy (M), Plasmacy system (P5), electronic patient journal (EP))

Patient knowledge and adherence is also identified and documented



Bottom of Medication reconciliation form – LIMM model



Outline of workshop

Brief introduction

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- the what's, why's, when's, who's and how's

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Medication review

- to perform a medication review
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Discussion & summary

Mary 87 years old

Diagnoses: Systolic heart failure, atrial fibrillation

Problems with: Unsteadiness, irritated skin, fatigue, sleeping difficulties, shortness of breath.

Case work – medication reconciliation Instructions

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- 1. Use the medication list & tool in front of you
- 2. Listen to the interview
- Note discrepancies between the medication list and what the patient tells the pharmacist
- 4. Summary which discrepancies did you identify?

Renal function: 70 ml/min, S-digoxin: 0.6 nmol/L. BP 120/60, HR 60/min



Medication list (bef	ore medicat	tion reconciliation)
Lisinopril tab	5 mg	1 morning
Metoprolol PR tab	50 mg	1 morning
Digoxin tab	0,25 mg	1 morning
Warfarin tab	2,5 mg	as indicated in list
Zopiclone tab	10 mg	1 evening
Furosemide tab	40 mg	1 morning 32

What did you identify?

Was the medication list we started out with correct?

Depa	riment	Bed	Name: M	ARV					Date of birth: 192	9	
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->	Lisinopril, t	ab, 5 mg		1-0-0-0			-		Θ		
->	Metoprolol,	PR tab,	50 mg	1-0-0-0					1x1	V	V
-0	Digoxin, ral	n. 0,25 m	e	1-0-0-0	Path	unch ti	me	0	0-1/2-0	V	V
>	Warfarin, ta	lb, 2,5 mj	t.	As indicated on the list					1×1	v	V
->	Zopiclone,	tab, 10 m	e.	0-0-0-1	P' neve	orus	ect		-		
~	Furosemide,	cab, 40 r	ng	1-0-0-0	P: Stay				E		
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Information from patient records Reasons for admission: Unstead	iness, fatigue, shortness of breath - worsening of heart failure?	Information during modication	re recoeccilation
Perina denies Heart failure, atrial fibrillation		A little bit of the morning	trowsy in
Social information			
Are you taking any other drugs?	Fais Minari Quinnach (Helabera (Hristoporoni) Quinep/arceny lation diags (Horiscians Weinan/naches (Hristoporoni)/agtenza-Bro		Patient's pharmac
Eye-/eardrops/natalspray 🖨 who	terral and de programme a construction of regions and / re		39 0

Outline of workshop

Brief introduction

Medication reconciliation

* practical example

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Discussion & summary

The medication review aims to optimize the prescribing of medicines and the use of prescribed medicines

For each patient, ask...

✓ Indication?	Is there an indication for each drug?Are there any untreated indications?
✓ Effect?	 Is each drug effective for the condition? Are the dosages correct (or too low)?
✓ Safety?	 Are there any drug-drug interactions or drug-disease interactions? Does the patient have any adverse drug reaction(s)? Are the doses correct (or too high)?
✓ Compliance?	Does the patient know how to use the drugs correctly?Is the patient able to use the drugs correctly?
	37

Mary 87 years old

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Renal function: 70 ml/min, Sdigoxin: 0.6 nmol/L. BP 120/60, HR 60/min



Case work: Medication review Instructions

- 1. Work in groups of two or three
- Perfom a medication review by using the information you have about Mary (and the checklist on the handout)

Medication list review)	(before me	dication
T. Warfarin	2,5 mg	as indicated on the list
T. Digoxin	0,125 mg	1x1 (at lunch
T. Omeprazole	20 mg	time) 1x1 (since Dec 2015)
T. Metoprolol	50 mg	1x1
T. Nitrazepam	5 mg	1x1 prn
Cream Canoder	m 5 %	when needed

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T. Warfarin	2,5 mg	as indicated on the list	T. Warfarin	2,5 mg	as indicated on the list
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T. Omeprazole	20 mg	1x1 (since Dec 2015)	T. Omeprazole	-20 mg	1x1 (since Dec 2015)
T. Metoprolol	50 mg	1x1	T. Metoprolol	50 mg	1x1
T. Nitrazepam	5 mg	1x1 prn	T. Nitrazepam	5 mg-	1x1 prn
Cream Canoder	m 5 %	when needed	Cream Canoder	m 5 %	when needed
			Enalapril	5 mg	1x1

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Discussion & summary

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The benefit of the medication review can be evaluated by assessing the quality of prescribing

- Frequency of "drug-related problems"
- Standardized and validated tools:
 - "Implicit", judgement-based criteria
 - Medication Appropriateness Index (MAI)
 - "Explicit", checklist-based criteria
 - Eg. Beers' criteria, PRISCUS, STOPP & START, ...







- Medication Appropriateness Index (MAI)
- Ten questions about each drug
- When an answer indicates inappropriateness, a score is assigned
- Scores are weighted and summated

- 1. Is there an indication for the drug?
- 2. Is the medication effective for the condition?
- 3. Is the dosage correct?
- 4. Are the directions correct?
- 5. Are the directions practical?
- 6. Are there clinically significant drugdrug interactions?
- 7. Are there clinically significant drugdisease/condition interactions?
- Is there unnecessary duplication with other drug(s)?
- 9. Is the duration of therapy acceptable?
- 10. Is this drug the least expensive alternative compared to others of equal utility?

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The STOPP and START criteria

- Checklist-based criteria
- Based on literature review and expert opinion
- Can be used as a tool for evaluation of prescribed medications but also when performing medication reviews
- STOPP: Screening Tool Of Older People's potentially inappropriate Prescriptions
 - 80 criteria
- START: Screening Tool to Alert doctors to Right (i.e. appropriate, indicated) Treatment
 - 34 criteria

The STOPP and START criteria



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T. Metoprolol	50 mg	1x1
T. Nitrazepam	5 mg	1x1 prn
Cream Canoder	m 5 %	when needed

45

46

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T. Digoxin	0,125 mg	1x1 (at lunch time)	T. Digoxin	0,125 mg	1x1 (in the morning)
T. Omeprazole	20 mg	1x1 (since Dec 2015)	T. Omeprazole	- 20 mg	1x1 (since Dec 2015)
T. Metoprolol	50 mg	1x1	T. Metoprolol	50 mg	1x1
T. Nitrazepam	5 mg	1x1 prn	T. Nitrazepam	5 mg-	1x1 prn
Cream Canoder	m 5 %	when needed	Cream Canoder	m 5 %	when needed
			Enalapril	5 mg	1x1

MAI assessment



Diagnoses: Systolic heart failure, atrial fibrillation

Mary

87 years old

Problems with: Unsteadiness, irritated skin, fatigue, sleeping difficulties, shortness of breath. Renal function: 70 ml/min, S-digoxin: 0.6 nmol/L. BP 120/60, HR 60/min



Mary MAI assessment 87 years old



Diagnoses: Systolic heart failure, atrial fibrillation

Problems with: Unsteadiness, irritated skin, fatigue, sleeping difficulties, shortness of breath. Renal function: 70 ml/min, S-digoxin: 0.6 nmol/L. BP 120/60, HR 60/min



Mary STOPP/START assessment



87 years old

Diagnoses: Systolic heart failure, atrial fibrillation

Problems with: Unsteadiness, irritated skin, fatigue, sleeping difficulties, shortness of breath. Renal function: 70 ml/min, S-digoxin: 0.6 nmol/L. BP 120/60, HR 60/min



Mary STOPP/START assessment 87 years old



Diagnoses: Systolic heart failure, atrial fibrillation

Problems with: Unsteadiness, irritated skin, fatigue, sleeping difficulties, shortness of breath. Renal function: 70 ml/min, S-digoxin: 0.6 nmol/L. BP 120/60, HR 60/min

Medication list (before)		STOPP	START
T. Warfarin 2,5 mg	as indicated on the list		
T. Digoxin 0,125 mg	1x1 (in the morning)		
T. Omeprazole 20 mg	-1x1 (since Dec 2015)	1	(
T. Metoprolol 50 mg	1x1		
T. Nitrazepam 5 mg	<u>1x1 prn</u>	1	
Cream Canoderm 5 %	when needed		
Enalapril 5 mg	1x1		1

OPEN @ ACCESS Fronty available online

Effects of Pharmacists' Interventions on Appropriateness of Prescribing and Evaluation of the Instruments' (MAI, STOPP and STARTs') Ability to Predict Hospitalization-Analyses from a Randomized Controlled Trial

Ulrika Gillespie¹⁺, Anna Alassaad², Margareta Hammarlund-Udenaes¹, Claes Mörlin³, Dan Henrohn^{9,2}, Maria Bertilsson⁵, Håkan Melhus³

Edvisor of Phermitoleutro and Drug Therapy Department of Phermisovecal Bosciesces, Uppade University and Uppade University Hospital, Uppade University Hospital, Uppade University Hospital, Uppade, Sweder, a Department of Medical Sciences, Uppade University Hospital, Uppade, Sweder, & Medical Practices, Uppade University Hospital, Uppade, Sweder, & Medical Practices, Uppade, University Hospital, Uppade, Sweder, & Medical Practices, Uppade, University Hospital, Uppade, University Hospital, Uppade, Sweder, & Medical Practices, Uppade, University Hospital, Uppade, University, Hospital, University, Hospital, University, Hospital, University, Hospital, University, Hospital, Uppade, University, Hospital, Uppade, University, Hospital, Uppade, University, Hospital, University, Hospital, Uppade, University, Hospital, University, Hospital, University, Hospital, Uppade, University, Hospital, University, Hospital, Uppade, University, Ho

Table 2. Scores on admission and at discharge and change from admission

Instrument		Intervention	Intervention group (in = 182)			Control group in = Tilds			
		Admission	Discharge	Change from administration?	Administer	Discharge	Onenge from administer*		
MW'	Million (SD)	63168	58 (42)	-83(83)	67.031	10.0 (7.9)	1.8 (8.1)	p-1100	
	Median (Mills, Max)	0.08-345	5 (8-30)	-21-26-88	7 (5-34)	8.5-(0-42)	11-7-10		
STORP*	Meani (SD)	1.4 (3.3)	0.11.9.0	+4.6 (10)	1.8 11:58	171181	4.2 (0.2)	p=10001	
	Median Stire Max	1.0-11	1.01-01	81-4-33	10-01	1 (0-4)	01-9-8		
TWIT!	Meany (SDC	0.4 (0.7)	8.7 (0.3)	0.3 (64)	8.4 (8.7)	5.5 (0.7)		p-3.801	
	Hedlari Diri, Maxi	0.01-41	8-10-21	10-4-0	0 (8-2)	1-(3-3)	01-3-D		

The clinical pharmacist intervention improved the quality of prescribing, as measured with MAI, STOPP and START

50, Senderd Alvision. "Searnaled Alvises per patient. "Number of MMI per patient. "Dander of mMI per patient. "Dander form antimotion collaberd as four at discharge four un administ. "Exologi four antimotion collaberd as four at discharge four un administr. "Second four well adaptive constraints for discharge four un administr. 25:01171/junnal.gove1005/41.002

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Pros and cons with the tools for appropriate prescribing ي.

	Implicit (judgement-based)	Explicit X (criterion-based) X
(+)	 Focus on the individual patient and are judgement-based and therefore more sensitive 	 Are easy to use Are not dependent on the experience and knowledge of the user Can be applied to large quantities of patients
\bigcirc	 Are time-consuming Require access to more extensive information about the patient 	 Don't account for the presence of co-morbidities or patient preferences The inclusion of drugs/criteria can be subject for controversy Need to be continuously updated

Are tools for evaluating appropriate prescribing associated with clinical outcomes?

Is a high quality of prescribing...



... linked to positive clinical outcomes?



Relationship between appropriate prescribing and clinical outcome

Effects of Pharmacists' Interventions on Appropriateness of Prescribing and Evaluation of the Instruments' (MAI, STOPP and STARTs') Ability to Predict Hospitalization— Analyses from a Randomized Controlled Trial

Ulrika Gillespie¹', Anna Alassaad¹, Margereta Hammarlund-Udenaus¹, Claus Mörlin¹, Dan Henrohm^{4,2}, Maria Bertilsson¹, Håkan Melhus²

Disease of Pharmanisers and Dog ParageDepartment of Pharmacount Residence, Sapana Liferenty, and Sapana Denserby Inspiral, Sapana, Seeting, 2015;patrices of Hindu Schwarz, Appendia Disease by A Sapana Schwarz, Vegana, David, Sapana Schwarz, Paraged Schwarz, Par

Table 3. Effect of WAL START, STOPP on number of total with to boubtal, number of readmissions and number of daug-telated
readmissions (N = 368).

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Aliated	1.8217-80-1.000, p=6.658-	100 1126-1841 (1-0000	1381-0384-0382 p+10801
STOPP			
Linutionet	1.05 (0.071.14, p=0.24	108 3098-1791 g-4037	1.0017-05-1000-0-1005
Adjunte	1.0518.07-1.156, p=0.22	108 ft/87-1.362 p=0.28	134(125-176 p-008
TRATE			
Deadlocked	1.03.2587-1.28. p=0.40	3.17 (019)-1.042 (1-00)8	1.4v (0.42-2.9%) p> 0.18
Adjusted	1.00 (0.80-1.10), p=8.29	118 805-1421 # +0.14	1.49 (0.81-0.45) (p=0.11)

798, Rate 1255, Cl. Canfidence interval doi:10.1971/journal.pone.5062481.0081 High MAI and STOPP scores at discharge were associated with a higher number of drugrelated readmissions

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 No statistically significant relationship was found between the scores and the total number of re-visits to hospital

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Discussion & summary

Medicines optimisation



NICE guideline Published: 4 March 2015 nice.org.uk/guidance/ng5 Medicines optimisation: the safe and effective use of medicines to enable the best possible outcomes

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Goals for the workshop

Teaching goals	Learning objectives	
1. To introduce methods used in medicines optimisation	After the workshop the participant should be able to:	
 To discuss the effects of medicines optimisation and how to measure these effects 	 describe medicines optimisation and its effects 	
 To show the advantages and disadvantages of applying these different tools 	 evaluate different tools used in medicines optimisation 	



NG 2 ssessment

- 1. Are you able to describe what medicines optimisation is and why it should be done? YES / NO
- 2. Can you describe why a medication reconciliation should be performed? YES / NO
- 3. Can you mention two tools used for evaluation of the quality of prescribing? YES / NO



Enjoy your congress !

