Clinical pharmacy in Belgium: from the bedside to the basement (and back to the bedside?)

#### EAHP SEMINAR – COMBINING CENTRALISED AND BEDSIDE CLINICAL PHARMACY

EAHP 2021 - Isabel Spriet

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• Nothing to disclose



- Can back-office clinical validation contribute to hospital-wide medication safety?
- Is it interesting to organise clinical pharmacy-services in a risk-based way?
- Can back-office clinical validation replace front-office clinical pharmacy services?



- Discuss the potential contribution of back-office clinical validation to overall clinical pharmacy services
- Understand the risk-based approach of back-office clinical validation
- Reflect about the ideal way of organising clinical pharmacy services in your (financial) context

# A "momentum" for the startup of clinical pharmacy





- Institute of Medicine, 1999
- 44,000-98,000 deaths/yr
- Medical errors 8th leading cause of death in the US
- Makary & Daniel, BMJ 2016
- Medical errors 3th leading cause of death in the US

Key element to improve quality of care: optimization of patients' pharmacotherapy

- Other elements creating the "momentum":
  - A large (ageing) population characterized by polypharmacy
  - Complex medication entering the market: NOAC, oral anticancer drugs, ...
  - Shift to shorter LOS
  - Initiation of fixed drug budget for hospitalized patients in 2006 in Belgium

# Shift in the role of the hospital pharmacist



#### • USA, Canada, UK, Australia

 Bedside clinical pharmacy implemented on patient wards since the '80s

 Development of clinical pharmacy services in Belgium since 2000

# Development and implementation of CDSS

- Since 2000: digitalization with EHR including electronic prescribing
- Active contribution of clinical pharmacists to the development and implementation of CDSS



#### **CDSS - basic**

(Drug-drug) interactions Maximum doses Drug use during pregnancy/lactation Therapeutic duplication Drug allergy

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TRADONAL (TABL ODIS 50 MG) # bij pijn om de 6 uur	🖉 РО	4*50 mg	4*50 mg	4*50 mg							

# Development of bedside 'front-office' clinical pharmacy services

 Partially funded by the Belgian government, first as "pilot projects" followed by structural funding ACTA CLINICA BELGICA 2019, VOL. 74, NO. 2, 75–81 https://doi.org/10.1080/17843286.2018.1462877

ORIGINAL PAPER

Development of clinical pharmacy in Belgian hospitals through pilot projects funded by the government

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- Front-office services
  - Attending ward rounds
  - Medication reconciliation on admission
  - Medication review
  - Medication counseling at discharge
  - Projects focusing on high-risk drugs, DDI, antibiotic stewardship...



Taylor & Francis

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Taylor & Francis Grou



CPS on each ward/consultation are provided by teams of 3 pharmacists – continuity of care

## Front-office services in Leuven – KPIs 2019

- Geriatric ward:
  - 1386 pharmacotherapeutic advices:
  - Acceptance rate: 86%
- Pediatric ward:
  - 1129 pharmacotherapeutic advices:
  - Acceptance rate: 92%
- Case discussions MID meeting trauma ward/septic orthopedic ward
  - 429 pharmacotherapeutic advices
  - Acceptance rate: 96%

Highly experienced bedside clinical pharmacist can contribute significantly (number of advices) and advice is highly appreciated and implemented (acceptance rate)

Development of bedside 'front-office' services

• FTE hospital pharmacists/ 100 beds



→ Implementation of bedside CP only on "high risk"-wards
 → CDSS & limited bedside CP do not cover all medication-related problems/risks



# **Back-office clinical pharmacy**

#### Clinical Validation Check of Medication Appropriateness (CMA)

Screening of patients at risk for potentially inappropriate prescriptions based on explicit clinical rules

Potentially inappropriate prescriptions are screened by clinical pharmacist

- Triggered by digitalization
- High-risk vs. all prescriptions
- Before or independent from distribution
- Stimulated by JCI "Check of medication appropriateness"
- "Low" staff investment
- Popular in Belgium

#### 2.6. Verdeling van de activiteiten tussen Back Office en Front Office

De klinisch-farmaceutische activiteiten kunnen zich afspelen in een specifieke zorgeenheid (bv. : intensieve zorgen), of transversaal met een doelgroep van patiënten in heel het ziekenhuis. Die doelgroep wordt bepaald door een gemeenschappelijk kenmerk (pathologie, farmacotherapie, leeftijd,...). Die activiteiten gebeuren dus op een gedecentraliseerde manier (in de zorgeenheid, aan het bed van de patiënt = *Front Office*), en/of op een gecentraliseerde manier (in de ziekenhuisapotheek = *Back Office*).



■ FO gemiddeld ■ BO gemiddeld

Figuur 13: gemiddelde tijdsverdeling tussen FO en BO



Gemiddeld wordt 62% van de klinischfarmaceutische activiteiten uitgevoerd vanuit de ziekenhuisapotheek (BO) tegenover 38% die plaatsvinden in de zorgeenheden (FO). Het aandeel van de activiteiten in *Front Office* is met 9% toegenomen ten opzichte van 2015.

Bij nader inzien, merken we een zekere variabiliteit in de verdeling. Drieëntwintig ziekenhuizen passen de klinische farmacie enkel gecentraliseerd toe, terwijl in 10 ziekenhuizen meer dan 75% van de activiteiten plaatsvinden binnen de diensten. Anderzijds voeren iets minder van de helft van de ziekenhuizen de klinische farmacie tussen 26 en 75% van de tijd gedecentraliseerd uit.

# Back-office clinical pharmacy - credits to the Netherlands!

#### Original research

#### Development of a computerised alert system, ADEAS, to identify patients at risk for an adverse drug event

M K Rommers,<sup>1</sup> M H Zegers,<sup>1</sup> P A De Clercq,<sup>2</sup> M L Bouvy,<sup>3</sup> P H E M de Meijer,<sup>4</sup> I M Teepe-Twiss,<sup>1</sup> H-J Guchelaar<sup>1</sup>

#### ABSTRACT

Introduction Adverse drug events (ADEs) are frequent and pose an important risk for patients treated with drugs. Fortunately, a substantial part of ADEs is preventable, and computerised physician order entry with a sophisticated clinical decision support system may be used to reach this goal.

Objective To develop a new automated system that could improve the quality of medication surveillance. The system should focus on detecting patients at risk for an ADE by combining data from the hospital information system and computerised physician order entry (drug prescription data, drug—drug interaction alerts, clinical chemical laboratory parameters, demographic features), using clinical rules. Methods The clinical rules were formulated in pharmacists to take corrective actions before harm occurs. In our hospital, the CPOE system in use provides online drug-drug interaction checks and drug-

dosing checks but is not integrated with a sophisticated CDSS.<sup>6</sup> This current surveillance system has some limitations: (1) there is no possibility to adapt medication surveillance to categories of patients and/or to medical specialities,<sup>7</sup> (2) the system does not take into account relevant laboratory values, and (3) most alerts are of low clinical relevance with the consequence of "alert fatigue".<sup>8</sup>

Given these limitations and the wish for more clinical pharmacy activities, we aimed to develop a new pharmacy decision support system that could improve the quality of our medication



richtlijneditor. Het laat de computer alle benodigde stappen uitvoeren. Deze DCS communiceert met alle informatiesystemen rond de patiënt, zoals het EPD, het huisartsinformatiesysteem (HIS), het apotheeksysteem (AIS), het ziekenhuisinformatiesysteem (ZIS), het anesthesietoestel of de patiënten monitor, maar ook met het elektronische systeem van de stadsapotheek of het streeklaboratorium (LIS).

Rommers MK et al. Artif Intell Med. 2013, 59:15-21 Rommers MK et al. Drug Saf. 2011, 34:233-42 Rommers MK et al. Qual Saf Health Care. 2010, 19(6):e35

Wasylewicz ATM et al. J Clin Pharm Ther. 2019 Dec 24



**Risk analysis** to identify patients at risk for PIM Formulating advanced clinical rules to screen for high risk patients

![](_page_16_Figure_4.jpeg)

Daily hospitalwide screening of all EHR generating a worklist

![](_page_17_Figure_3.jpeg)

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	CMA Interacties PCIA/PCEA (27)		57-05	-	0				041	GER	102/2	Zorgregistratie van FPgewicht { FPgewichtW = 61.9, FPgewichtBMI = true } voor patient 80740731 uitgevoerd door <u>uitvoerder: x258278</u> op 2019-03-03 11:22:11.			
	CMA NSAIDs zonder PPI (GI versie)														
	CMA Novalgine en agranulocytose/ne											<ul> <li>Voorschrift voor LIXIANA uit te voeren op 2019-03-08 08:00:00 en gevalideerd door validator;</li> </ul>			
	CMA Overrules van interacties (477)											<u>dmerte0</u> op 2019-03-02 22:37:33.			
	CMA Rivaroxaban bij verminderde nie											In de periode van 2019-03-08 00:00:00 tot 2019-03-08 23:59:59 waren er elke dag			
	CMA Screening NRS (opvolging) (20'											voorschritten			
	CMA TPN zonder konakion (34)											- Suggesties your bericht			
	CMA Temocilline GI (11)														
	CMA apixaban dosisaanpassing (20)											V / Interne nota voor apotheek			
	CMA cefazoline (12)														
	CMA edoxaban dosis (17)											Bekijk EMV			
	CMA hyperkalemie (45)											🖄 🥒 Bekijk laboresultaten			
	CMA hypokalemie (142)											Debilk zere			
	CMA metformine en nier (21)	-										Dekijk zorg			
	CMA verhoogde INR bij VKA (10)											Maak opvolgnota			
	Dabigatran (Pradaxa): dosisaanpassi											04 03 2019 jhias1 25962208 ADVIES APOTHEEK: De standaarddosis van Lixiana			
	Labogestuurde QTc verlengende mec											(edoxaban) bedraagt 60 mg 1x per dag. Dosisreductie is pas aangewezen bij een matige			
	NSAID bij verminderde nierfunctie (10											therapie			

Validation by trained clinical pharmacists using standardized flowcharts

![](_page_19_Figure_3.jpeg)

Interventions via electronic alerts or phone calls

# **Example: screening for incorrect dosing of edoxaban**

• Intervention: call + electronic alert

![](_page_20_Picture_4.jpeg)

	Afdeling	Template	Tekst	Contact type	Zender	Groep
	GER	opvolgnota	ADVIES APOTHEEK: Op basis van de nierfunctie (CrCl < 50 ml/min) wordt de dosis van Lixiana (edoxaban) aangepast naar 30mg 1x per dag. Graag nazicht therapie.	hospitalisatie		apotheek
1						

# CMA: what's in a name?

![](_page_21_Figure_1.jpeg)

### December 2018 – November 2020

Pharmacotherapeutic bundle	Clinical rules	Actions provided by	Acceptance rate by	
	implemented (n)	pharmacists (n)	physicians (%)	
Antibiotic stewardship	43	2710	79.9%	
Anticoagulant stewardship	13	2737	74.2%	
Pain management	15	2944	77.7%	
IV to oral switch therapy	2	1206	74.1%	
Varia	9	2248	75.8%	
Total	82	11845	77%	

- CMA serves as a bridge between CDSS and (limited) bedside clinical pharmacy
- CMA contributes in an important way to medication safety in our hospital
  - Pharmacotherapeutic support is now provided <u>hospital-wide</u>: also to patients admitted on wards on which a bedside pharmacist is not available
  - As clinical rules are screening continuously and in <u>'near-time'</u>, changing biochemical parameters impacting pharmacotherapy are rapidly taken up

## The basement vs. the bedside?

- Back-office clinical pharmacy services are complementary to bedside services
- Back-office clinical validation is an add-on but can not replace bedside services
  - Acceptance rate is higher for advice provided by bedside pharmacist
    - Bedside pharmacist has a more focused expertise
    - Orally provided advices have a higher impact than written ones
  - Type of clinical pharmacy services is not identical
    - Back-office: screening for potentially inappropriate prescriptions, ADEs
    - Front-office: multi-faceted approach, including
      - Med rec, medication review, discharge counseling, specific projects, discussion on complex cases
  - Experienced front-office clinical pharmacist provides many spontaneous advices, even when using explicit screening tools

![](_page_24_Picture_11.jpeg)

## From the bedside to the basement... and now back to the bedside?

Application of clinical rules to guide clinical pharmacists at te bedside

For example:

- clinical pharmacists counseling patients with chronic leukemia on DDIs –DHIs are available for 0.5 FTE at the consultation
- On average, 80 patients are daily visiting the consultation
- Clinical rule based screening can help and support the clinical pharmacist in prioritizing their work at the consultation

![](_page_25_Picture_6.jpeg)

Plan for the near future:

- Development of clinical rule based bedside clinical pharmacy services
- Trauma ward, emergency dpt, pediatric ward
- Comparison of clinical rule based advices with historical spontaneous, randomly given ones

![](_page_25_Picture_11.jpeg)

- Back-office clinical validation contributes to hospital-wide medication safety
- It is interesting to organise clinical pharmacy-services in a risk-based way
- Back-office clinical validation is complementary to front-office CP services