

Contribution of Different Patient Information Sources to Create the Best Possible Medication History

Contribuição de Diferentes Fontes de Informação para Obter a Melhor História Farmacoterapêutica Possível



Joelizy OLIVEIRA^{1,2}, Ana Cristina CABRAL^{2,3}, Marta LAVRADOR^{2,3}, Filipa A. COSTA^{4,5}, Filipe Félix ALMEIDA⁶, António MACEDO⁶, Carlos SARAIVA⁷, Margarida CASTEL-BRANCO^{2,3}, Margarida CARAMONA², Fernando FERNANDEZ-LLIMOS⁸, Isabel Vitória FIGUEIREDO✉^{2,3}
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ABSTRACT

Introduction: Obtaining the best possible medication history is the crucial step in medication reconciliation. Our aim was to evaluate the potential contributions of the main data sources available – patient/caregiver, hospital medical records, and shared electronic health records – to obtain an accurate ‘best possible medication history’.

Material and Methods: An observational cross-sectional study was conducted. Adult patients taking at least one medicine were included. Patient interview was performed upon admission and this information was reconciled with hospital medical records and shared electronic health records, assessed retrospectively. Concordance between sources was assessed. In the shared electronic health records, information was collected for four time-periods: the preceding three, six, nine and 12-months. The proportion of omitted data between time-periods was analysed.

Results: A total of 148 patients were admitted, with a mean age of 54.6 ± 16.3 years. A total of 1639 medicines were retrieved. Only 29% were collected simultaneously in the three sources of information, 40% were only obtained in shared electronic health records and only 5% were obtained exclusively from patients. The total number of medicines gathered in shared electronic health records considering the different time frames were 778 (three-months), 1397 (six-months), 1748 (nine-months), and 1933 (12-months).

Discussion: The use of shared electronic health records provides data that were omitted in the other data sources available and retrieving the information at six months is the most efficient procedure to establish the basis of the best possible medication history.

Conclusion: Shared electronic health records should be the preferred source of information to supplement the patient or caregiver interview in order to increase the accuracy of best possible medication history of the patient, particularly if collected within the prior six months.

Keywords: Electronic Health Records; Medical History Taking; Medication Reconciliation

RESUMO

Introdução: A obtenção da melhor história farmacoterapêutica possível é uma etapa crucial da reconciliação da medicação. O objetivo foi avaliar as potenciais contribuições das principais fontes de informação disponíveis – doente/cuidador, Processo Único, Plataforma de Dados da Saúde e – para obter uma mais exacta melhor história farmacoterapêutica possível.

Material e Métodos: Foi realizado um estudo transversal observacional. Incluíram-se doentes adultos a tomar pelo menos um medicamento. A entrevista com o doente foi realizada na admissão e os dados do Processo Único e da Plataforma de Dados da Saúde recolhidos retrospectivamente. A concordância entre as fontes de informação foi avaliada. Na plataforma de dados da saúde, os dados foram recolhidos em quatro janelas temporais: os últimos três, seis, nove e 12- meses. Os dados omitidos entre os diferentes tempos foram analisados.

Resultados: Participaram 148 doentes, com uma idade média de $54,6 \pm 16,3$ anos. Foram recolhidos 1639 medicamentos. Destes, 29% foram obtidos simultaneamente nas três fontes de informação, 40% foram obtidos apenas na Plataforma de Dados da Saúde e 5% foram obtidos exclusivamente a partir do doente. O número total de fármacos recolhidos na Plataforma de Dados da Saúde nos diferentes tempos foi 778 (três meses), 1397 (seis meses), 1748 (nove meses) e 1933 (12 meses).

Discussão: A consulta da Plataforma de Dados da Saúde permite obter dados omitidos nas outras fontes de informação e a recolha dos seis meses precedentes é o procedimento mais eficiente para constituir a base da melhor história farmacoterapêutica possível.

Conclusão: A Plataforma de Dados da Saúde deve ser a fonte de informação preferencial para complementar a entrevista do doente/cuidador de forma a aumentar a exatidão da melhor história farmacoterapêutica possível, particularmente se a informação for recolhida em relação aos seis meses precedentes.

Palavras-chave: Anamnese; Reconciliação de Medicamentos; Registos Eletrónicos em Saúde

INTRODUCTION

Unsafe medication practices and medication errors are a leading cause of injury and avoidable harm in health care systems across the world. The transition of care is a critical point in patient safety, increasing the risk of medication-

1. CAPES Foundation. Ministry of Education. Brasília. Brazil.

2. Departamento de Farmacologia e Cuidados Farmacêuticos. Faculdade de Farmácia. Universidade de Coimbra. Coimbra. Portugal.

3. Coimbra Institute for Clinical and Biomedical Research. Faculdade de Medicina. Universidade de Coimbra. Coimbra. Portugal.

4. Centro de Investigação Interdisciplinar Egas Moniz. Instituto Universitário Egas Moniz. Monte de Caparica. Portugal.

5. Grupo de Farmacoepidemiologia. Research Institute for Medicines - iMed. ULisboa. Universidade de Lisboa. Lisboa. Portugal.

6. Serviço de Psiquiatria. Centro Hospitalar e Universitário de Coimbra. Coimbra. Portugal.

7. Departamento de Psiquiatria. Faculdade de Medicina. Universidade de Coimbra. Coimbra. Portugal.

8. Laboratório de Farmacologia. Departamento de Ciências do Medicamento. Universidade do Porto. Porto. Portugal.

✉ Autor correspondente: Isabel Vitória Figueiredo. isabel.vitoria@netcabo.pt

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related incidents and preventable hospital admissions.^{1,2} Therefore, within the framework of the general plan for patient safety, in 2016, the Portuguese General Directorate of Health [Direção Geral da Saúde (DGS)] published a clinical practice guideline on medication reconciliation: Norma 018/2016.³ The purpose of this guideline was to encourage the implementation of medication reconciliation – the formal process in which health care professionals partner with patients to ensure accurate and complete medication information transfer at interfaces of care.⁴ Medication reconciliation has demonstrated to improve safety and efficacy of medication prescribed between transitions of care, and reducing the risk of adverse events and consequently patients' medication-related morbidity and mortality.⁵⁻⁷

Medication reconciliation is a three-stage process. The first stage consists of gathering the best possible medication history (BPMH), which represents a compilation of different sources of information to obtain a list with the current medications (prescribed and non-prescribed) the patient was using before the transfer of interface of care; the second stage involves comparing the BPMH with a list of the current prescribed medication at the new interface of care, with the aim of identifying discrepancies; and finally in the third stage, these discrepancies are 'reconciled' with the medical team by differentiating intentional and unintentional ones.³ Thus, obtaining the BPMH is a crucial step to initiate the reconciliation process, since incomplete or inaccurate medication histories can increase the risk of medication-related errors and potential harm.⁸⁻¹¹ Unresolved discrepancies may lead to discontinuation of clinically important medicines, or introduction of inappropriate or interacting medicines, which can potentially lead to hospital admissions, readmissions, or increase hospital stays.¹²

The World Health Organization (WHO) defends the use of a systematic process to obtain the BPMH, considering the patient/caregiver as the main source of information.⁴ However, since patient's information may not always be reliable, other sources should be used to enhance the accuracy of medication history, such as hospital medical records (HMR) and shared electronic health records (SEHR). SEHRs are increasingly used worldwide and are leading to decreases in the number of medication errors and their role in the provision of health care is progressively recognized.¹³⁻¹⁶ In Portugal, in June 2012, an universal and centralised patient electronic health record platform [Plataforma de Dados da Saúde (PDS)] – was developed, enabling health professionals to have access to all the patient's clinical information,¹⁷ becoming one of the main sources to consider in medication reconciliation to obtain supplementary information to patient reports. The Portuguese PDS connects more than 370 health care institutions, including all the public hospitals in Portugal, compiling their medical records into five central data bases.

Our aim was to evaluate the potential contributions of the three main data sources available in Portugal – patient/caregiver, HMR, and SEHR (the Portuguese PDS) at different time frames – to obtain an accurate and reliable BPMH.

MATERIAL AND METHODS

An observational study was conducted in an acute care unit of the Center for Integrated Responsibility of Psychiatry and Mental Health, Coimbra University Hospital (CHUC) (January 2015 – February 2016). The study was approved by the Ethics Committees of Hospital University Center of Coimbra (CHUC-008-15) and University of Coimbra Faculty of Medicine (CE 109/2014). All participants signed an informed consent prior to their inclusion in the study.

Population and setting

This acute care unit consists of 27 beds in each of the two wards. The median length of hospitalization is 1.5 months, which means that the hospital unit receives on average 432 patients per year. Patients older than 18 years of age and taking at least one medicine at the moment of admission in the unit were invited to participate in the study. Exclusion criteria comprised pregnancy, patients controlled without medication, or patients unable to communicate by themselves or through a caregiver (due to cognitive impairment or language barriers).

Data sources

Data were obtained through three sources of information:

- Patient/caregiver – A standardized face-to-face interview was conducted by a trained pharmacist, using a data collection form, with the patient or the caregiver, within 72 hours after admission. When it was not possible to establish an interview with the patient due to serious mental impairment or other situations compromising the reliability of the information collected, the caregiver was invited. The main variables of interest collected were: medications currently taken, including prescription and non-prescription medicines, patient's medical conditions, allergies, and information about previous adverse drug reactions.
- Hospital medical records – HMR, the electronic medical record commonly used at the hospital, were used to collect information on the clinical information pertaining to the moment of admission at the mental unit, including admission diagnoses, and medicines prescribed at admission, and medicines identified during admission process as currently used (t0). This data source was assessed retrospectively.
- Shared electronic health record – SEHR (the Portuguese PDS) was accessed online from the acute care unit and was used to obtain the patient's medication history from the preceding year. This data source contains all the medicines prescribed to the patient, sequentially ordered, since 2012, which creates the need to define the best time cut-off to enable an efficient and feasible consultation in clinical practice. Information was collected for four time-periods: the preceding three months (t3), six months (t6), nine months (t9), and 12-months (t12).

Data analysis

Medicines were classified using the anatomical therapeutic chemical classification (ATC), with detail at the first level (ATC 1) and second level (ATC 2).¹⁸ Descriptive statistics were performed. To compare the four time cut-offs of the SEHR [the preceding three months (t3), six months (t6), nine months (t9), and 12-months (t12)], information was assessed using the proportion of omitted data in each of these moments and then sub-analysed by the ATC group. The selection of the best retrospective cut-off was made based on two criteria: a) proportion of medicines lost when reducing the period of time analysed, and b) clinical relevance of the potential treatment omission and the potential substitutive effects of another prescribed medicine to the patient (e.g., the omission of sertraline was considered irrelevant if Mirtazapine was newly prescribed). Medicines omitted from six to nine months analysis and from nine to 12-months analysis were classified by analysing the medical records as: medicines for acute conditions, changes in prescribed medicine, dose treatment changes, end of treatment, or real omissions.

RESULTS

During the study period, 148 patients were admitted to the acute mental health unit, with a mean age of 54.6 years (SD = 16.3) and 75 (50.7%) were females. Patient primary diagnoses are presented in Table 1.

SEHR best retrospective time cut-off

Table 2 provides the proportion of medicines omitted when considering one retrospective cut-off compared to the immediately longer period of time. A total of 778 medicines were retrieved considering the three months' time frame, 1397 medicines in the six months' time frame, 1748 medicines in the nine months' time frame, and 1933 medicines considering the 12-months' time frame of the SEHR. Comparing the three months cut-off versus the six months cut-off, the information obtained decreased 44.3%, while the decreases were lower in the remaining cut-offs compari-

sons (20.1% t6 vs t9, and 9.6% t9 vs t12). Table 3 presents the causes ascertained for medication omissions from six to nine months and from nine to 12-months analyses, classifying 55 as real omissions from six to nine months cut-off, and 19 from nine to 12-months.

Contribution of each information source to BPMH

After aggregating the medicines obtained from the three sources (i.e., patient/caregiver, HMR, and SEHR at six months) a total of 1639 medicines were found. Only 476 medicines (29%) were included in the three sources analysed. Ignoring one of these three sources to create the BPMH produced substantially different results. A total of 653 (39.8%) medicines would be lost if the 6-months retrospective cut-off of the SEHR is not used, eight (0.5%) when the HMR is not used, and 77 (4.7%) when patient/caregiver is not interviewed (Table 4). A total of 157 medicines omitted when using only the SEHR would be recuperated using any of the two other sources (Fig. 1).

DISCUSSION

In our study we evaluated the potential contributions of the three main available data sources (patient/caregiver, HMR, and SEHR) to obtain an accurate BPMH. Considering the large amount of information contained in SEHR, it was necessary to define the best time cut-off, especially because the literature is not consistent in establishing the best time frame to an SEHR retrospective consultation, with different authors suggesting different time cut-offs: Kalb *et al*¹⁹ used the information of the previous 14 months; Lau *et al*⁹ used the information of the previous 12 months, while Prins *et al*²⁰ and Soler-Giner *et al*²¹ used the information of the previous six months. In our study, nearly 45% of the medicines retrieved in the six month analysis would be ignored if we had retrieved information only from the three more recent months. Medication for the nervous system (group N) presented the most similar profile when comparing the information at three and six months.

The number of medicines lost reduced to about 20%

Table 1 – Patient primary diagnosis coded by the International Statistical Classification of Diseases and Related Health Problems 10th Revision (ICD 10)

Patient primary diagnosis	n (%)
F30-F39 Mood [affective] disorders	59 (39.9)
F00-F09 Organic, including symptomatic, mental disorders	42 (28.4)
F10- F19 Mental and behavioral disorders due to psychoactive substance use	15 (10.1)
F50- F59 Behavioral syndromes associated with physiological disturbances and physical factors	9 (6.1)
F20-F29 Schizophrenia, schizotypal and delusional disorders	8 (5.4)
F60-F69 Disorders of adult personality and behavior	6 (4.1)
F70-F79 Mental retardation	3 (2.0)
F40-F48 Neurotic, stress-related and somatoform disorders	2 (1.4)
R40-R46 Symptoms and signs involving cognition, perception, emotional state and behavior	2 (1.4)
F80-F89 Disorders of psychological development	1 (0.7)
F99-F99 Unspecified mental disorder	1 (0.7)

Table 2 – Distribution of medicines registered at the SEHR at the four time points by ATC code

ATC 1 st level	Number (%) of medicines recorded in SEHR in the preceding months												Medicines omitted					
	3 m		6 m		9 m		12 m		3 m vs 6 m		6 m vs 9 m		9 m vs 12 m					
	n	%	n	%	n	%	n	%	n	%	n	%	n	%				
A: Gastrointestinal tract and metabolism	83	10.7	171	12.2	214	12.2	230	11.9	88	51.5	43	20.1	16	7.0				
B: Blood and blood forming organs	24	3.1	55	3.9	68	3.9	69	3.6	31	56.4	13	19.1	1	1.4				
C: Cardiovascular system	70	9.0	155	11.1	194	11.1	209	10.8	85	54.8	39	20.1	15	7.2				
D: Dermatologicals	11	1.4	31	2.2	46	2.6	57	2.9	20	64.5	15	32.6	11	19.3				
G: Genito urinary system and sex hormones	11	1.4	18	1.3	24	1.4	30	1.6	7	38.9	6	25.0	6	20.0				
H: Systemic hormonal preparations, excl. sex hormones and insulins	6	0.8	11	0.8	12	0.7	13	0.7	5	45.5	1	8.3	1	7.7				
J: Antineoplastic and immunomodulating agents	21	2.7	53	3.8	67	3.8	86	4.4	32	60.4	14	20.9	19	22.1				
L: Antineoplastic and immunomodulating agents	1	0.1	2	0.1	2	0.1	2	0.1	1	50.0	0	0	0	0				
M: Musculo-skeletal system	28	3.6	60	4.3	87	5.0	107	5.5	32	53.3	27	31.0	20	18.7				
N: Nervous system	479	61.6	752	53.8	908	51.9	976	50.5	273	36.3	156	17.2	68	7.0				
P: Antiparasitic products, insecticides and repellents	1	0.1	2	0.1	2	0.1	2	0.1	1	50.0	0	0	0	0				
R: Respiratory system	26	3.3	53	3.8	75	4.3	97	5.0	27	50.9	22	29.3	22	22.7				
S: Sensory organs	16	2.1	33	2.4	41	2.3	47	2.4	17	51.5	8	19.5	6	12.8				
V: Various	1	0.1	1	0.1	8	0.5	8	0.4	0	0	7	87.5	0	0				
TOTAL	778	100	1397	100	1748	100	1933	100	619	44.3	351	20.1	185	9.6				

SEHR: shared electronic health record

when comparing six and nine months and to nearly 10% when comparing nine and 12-months analyses. This may be a consequence of the repeat prescribing system in Portugal, where a patient can obtain a prescription order with up to six months validity. Of these 351 medicines lost when comparing the six months and the nine month cut-offs, only 55 (15.7%) were real omissions, with 15 medicines (4.3%) considered as potentially serious omissions (anticoagulant, antiplatelet and antihypertensive medicines). These findings suggest that retrieving the information at six months is the most efficient procedure to constitute the basis of the BPMH.

The use of SEHR to obtain BPMH allows not only identifying inaccurate doses and frequencies of prescribed medicines²² but also increases the accuracy of BPMH by providing data that were omitted in the other data sources available. In agreement with the WHO recommendations,⁴ we found that the SEHR should be supplemented with a face-to-face interview with the patient or the patient's caregiver. When supplementing the SEHR with the interview, the two drug classes more frequently added to the BPMH were the gastrointestinal tract drugs (ATC group A), followed by nervous and musculoskeletal system drugs (groups N and M, respectively). This way, the use of SEHR allows a reduction in the frequency of the main errors of the medication history identified at admission to hospital.²³ Similar results were obtained in other studies made with National databases.²⁴ Our study also demonstrated that the HMR does not add relevant data to that obtained with the other two information sources. In summary, the three sources present different strengths: while the SEHR revealed as the source providing more and more accurate information, patient or the patient's caregiver interview added information about non-prescription medicines and shared prescription medicines,²⁵ and finally HMR complemented the BPMH with hospital-based prescriptions.

Although the SEHR demonstrated the highest potential to feed the BPMH, several limitations should be considered and overcome. The existence of different computerized prescribing systems among hospitals requires the use of information transfer processes to ensure that information is not lost when submitted to central repositories like the Portuguese Plataforma de Dados da Saúde – PDS. Technology-based solutions are being pointed as an important way to improve health care of patients with polypharmacy and multiple long-term conditions.²⁶ Consequently, in the future, shared medication information repositories could simplify the reconciliation during the transitions of care, and increase patient safety.²⁷

Table 3 – Causes of medicines omission between t6 vs t9 and between t9 vs t12

Omissions	t6 vs t9	t9 vs t12
Types of omission	n (%)	n (%)
Medicines for acute conditions	165 (47.0)	93 (50.3)
Medicines switches (i.e., changes in prescribed medicines)	93 (26.5)	56 (30.3)
Changes in treatment dose	24 (6.8)	10 (5.4)
End of treatment	14 (4.0)	7 (3.8)
Real omissions	55 (15.7)	19 (10.3)
Potentially serious real omissions	15 (4.3)	7 (3.8)
TOTAL	351 (100)	185 (100)

Table 4 – Distribution of medication on the BPMH by ATC code and relative loss of information from each of the three sources considered

ATC 1 st level	BPMH	Lost if SEHR unused	Lost if HMR unused	Lost if patient's interview unused
A: Gastrointestinal tract and metabolism	227 (13.8%)	107 (6.5%)	0	32 (1.9)
B: Blood and blood forming organs	70 (4.3%)	20 (1.2%)	1 (0.1%)	7 (0.4%)
C: Cardiovascular system	189 (11.5%)	30 (1.8%)	0	9 (0.5%)
D: Dermatologicals	35 (2.1%)	28 (1.7%)	0	1 (0.1%)
G: Genito urinary system and sex hormones	20 (1.2%)	11 (0.7%)	0	0
H: Systemic hormonal preparations, excl. sex hormones and insulins	11 (0.7%)	5 (0.3%)	0	0
J: Antiinfectives for systemic use	57 (3.5%)	52 (3.2%)	0	0
L: Antineoplastic and immunomodulating agents	2 (0.1%)	0	0	0
M: Musculo-skeletal system	72 (4.4%)	40 (2.4%)	0	11 (0.7%)
N: Nervous system	850 (51.9%)	281 (17.1%)	6 (0.4%)	16 (0.9%)
P: Antiparasitic products, insecticides and repellents	3 (0.2%)	2 (0.1%)	0	0
R: Respiratory system	60 (3.7%)	49 (2.9%)	1 (0.1%)	0
S: Sensory organs	38 (2.3%)	26 (1.6%)	0	0
V: Various	5 (0.3%)	2 (0.1%)	0	1 (0.1%)
TOTAL	1639 (100%)	653 (39.8%)	8 (0.5%)	77 (4.7%)

BPMH: best possible medication history; SEHR: shared electronic health record; HMR: hospital medical record

Study limitations

Our study has included patients admitted to an acute mental hospital unit. We cannot ensure that the results are similar to those that could be obtained with patients with other medical conditions or recruited in other hospital units in Portugal.

CONCLUSION

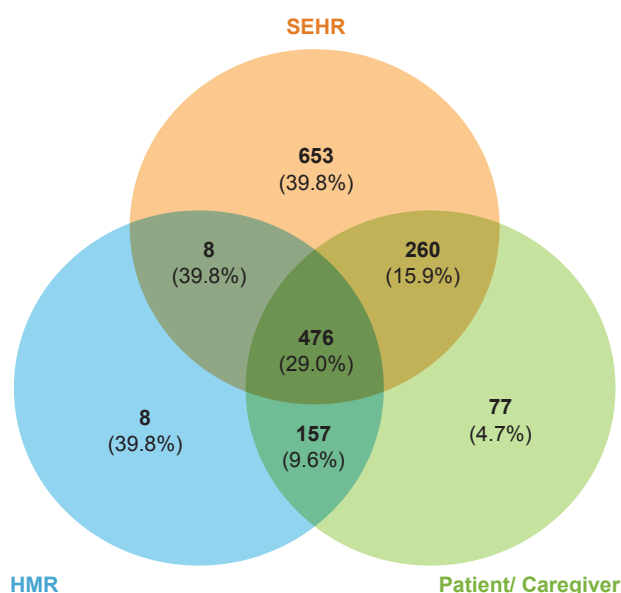
Electronical health records represent the most comprehensive source of information to create the patient's BPMH, although SEHR should be supplemented with a patient or patient's caregiver interview. The most efficient medication data retrieval process should consider a 6-month retrospective analysis of the SEHR.

PROTECTION OF HUMANS AND ANIMALS

The authors declare that the procedures were followed according to the regulations established by the Clinical Research and Ethics Committee and to the Helsinki Declaration of the World Medical Association.

DATA CONFIDENTIALITY

The authors declare having followed the protocols in

**Figure 1** – Venn diagram exhibiting the contribution of each data source for the BPMH

use at their working center regarding patients' data publication.

CONFLICTS OF INTEREST

The authors declare they have no conflict of interest.

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