Evaluation of Effects of Medical Reconciliation in Renal Failure Patients in a Tertiary Care Hospital

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ABSTRACT

Renal failure is one of the major health problems in the world. Any gaps in the patient care may affect the patient adherence and quality of life. As part of a multi-disciplinary patient care strategy, clinical pharmacy service has led to improvement in patient care. The purpose of the study is to evaluate the role of medication reconciliation in renal failure patients and to observe the effect of medication alert card and patient counseling in medication reconciliation. In control, Drug Related Problems (DRPs) were identified only by chart review. In test group, one arm containing patients with DRPs were identified using medication reconciliation alone and in second arm, patient counseling and medication alert card was used in addition to reconciliation. Patient data were recorded on a standard template obtained from The Manitoba Renal Program (Canada). DRPs were recorded. Knowledge was assessed with a questionnaire before and after counseling and it was quantified. More number of DRPs was identified by means of medication reconciliation. Patient counseling and medication alert card had a significant role in medication reconciliation. The study was statistically significant with P < 0.05. Medication reconciliation in renal failure helps to identify more number of drug related problems. Patient counseling and medication alert card have effectively improved the effect of reconciliation in renal failure patients and thus could improve clinical outcomes and quality of life.

Key words: Medication reconciliation, Patient counselling, Medication alert card, Renal failure.

INTRODUCTION

Renal failure also known as renal insufficiency is the reduced ability of the Kidney to filter the waste materials mainly, the nitrogenous products. Chronic kidney disease (CKD) and End Stage Renal Disease (ESRD) are emerging public health problems in developing countries thereby needing changes in health-care policy. When patients having intercurrent diseases receive multiple medications, their chances of Drug Related Problems (DRPs) like druginteractions and adverse drug reactions tend to increase.1 Drug-related problems include medication errors (involving an error in the process of prescribing, dispensing, or administering a drug, whether there are adverse consequences or not) and adverse drug reactions.2 To reduce the burdens of DRPs caused on the patient as well as the health care provider, it is essential to implement methods to identify and measure such DRPs and tools to rectify the same. Some of the tools available are medication reconciliation, patient counselling and medication alert card. Medical reconciliation according to Joint Committee Accreditation Health Care Organizations (JCAHO) is the "process of comparing a patient's medication orders to all of the medications that the patient has been taking". This reconciliation is done to avoid medication errors such as omissions, duplications, dosing errors or drug interactions and Adverse Drug Events (ADEs). ESRD patients on an average are prescribed with 10-12 medications which increases the potential for Drug Related problems (DRPs) including inappropriate dose or indication for a medication and Adverse Drug Events (ADEs). Medication discrepancies are

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bound to occur unintentionally due to medication use before admission and after discharge. These are points of care where patient is more exposed to medication discrepancies as the responsibilities of a patient are transferred among health professionals.⁴

The significance of medication reconciliation includes mainly assessing whether the desired therapeutic outcomes are being achieved, monitoring for Drug Related Problems (DRPs), monitor for toxicity, ensure rational and quality use of medicines, assess patient's compliance (medication adherence), assess the completeness of medication chart, compare the inhospital and discharge medication and judge whether medicines should be included and omitted or if better alternatives could be suggested.⁵

Components majorly include medication order review (MOR)/ Treatment Chart Review (TCR), Clinical review/ Daily Progress Review, detection and management of Adverse Drug Reactions (ADRs).⁵

Steps involved in medication reconciliation process includes the following.

- Verification- The current (in-hospital) medication list is assembled by using 1 or more sources of information (e.g.: pharmacy records, general practitioner medical records, medication vials brought by the patient, information provided by the patient and his/her family in patient counselling).⁶
- 2. Clarification- The medication and dosages are checked for appropriateness.⁶
- Reconciliation- Newly prescribed medications are compared against the old ones and changes to pharmacotherapy are documented.⁶
- 4. Transmission-The updated and verified list is communicated with the next provider of care.⁶

Patient counselling refers to the process of providing information, advice and assistance to help patients use their medications appropriately. The information and advice is given by the pharmacist directly to the patient or to the patient's representative and may also include information about the patient's illness and recommended lifestyle changes. The contents of typical counselling include Correct storage, minimum duration required to show therapeutic benefit, what to do if a dose It includes name and strength of the medication, the reason why it has been prescribed (if known), or how it works, how to take the medication? (how much and how often), expected duration of treatment, expected benefits of

treatment, possible adverse effects, possible medication or dietary interactions, advice on is missed? special monitoring requirements, for example: blood tests and arrangements for obtaining further supplies.⁷

MATERIALS AND METHODS

Study design and location

A prospective interventional study was conducted in the department of nephrology PSG Hospitals, Coimbatore. The study was approved by Institutional Ethics Committee (IHEC, PSG IMS and R) on 29/01/2016.

Study subjects and duration

All adults discharged with at least one prescribed drug from the department of Nephrology were included. The study was conducted for a period of six months.

Inclusion criteria

- Above 18 years of age.
- Patient willing to give consent.
- Renal failure and dialysis patients.

Exclusion criteria

- Pregnancy and lactation
- Psychiatric disturbances
- Death
- Transferred to another ward or hospital

Discharge within 24 h or out of office h and patients who could not be counseled (as stated by hospital physician due to physical/mental constraints, language restrictions or terminal illness) in the year 2016.¹¹

PROCEDURE

Medication review and reconciliation was conducted and documented using a standard template from the Manitoba Renal Program (Canada).⁸ Medication alert card and its questionnaire were obtained from Medipal. Patient counseling questionnaire was obtained from World journal of pharmacy and pharmaceutical sciences for assessing the knowledge before and after counselling in renal failure patients. Medication reconciliation intervention chart was obtained from PCNE version-4 used by the physicians to make the interventions with the medication list.¹⁰

The study contained one test group and control group. A control group was created in which medical reconciliation,

patient counselling and medication alert card were not provided. In the test group, medication reconciliation was performed. This group was divided into two to determine the effect of patient counselling and medication alert card on medication reconciliation. More number of DRPs was identified by means of medication reconciliation. Student *t*-test was used to compare the Test and the Control. Paired-*t*-test was used to compare Group-1 and Group-2 under the Test. Patient counseling and medication alert card had significant role in medication reconciliation. The study was statistically significant with *P*<0.05.

RESULTS AND DISCUSSION

Patient demographics of study patients

Age wise distribution

A total of 152 subjects were included in this study. Subjects were classified into 3 age groups less than 40 years, 40-60 years and greater than 60 years. There were 21.05% subjects in the age group of less than 40 years, 44.73% in 40-60 years and 34.21% in the age group greater than 60 years. As per the results, maximum subjects (44.73%) were in the age group of 40 to 60 years, followed by greater than 60 years (34.21%). Table 1.

Gender wise distribution

Among total population of 152 patients, 65.78% (n=100) were males and 34.21% (n=52) were females. In this group, male subjects were more compared to female subjects Table 2.

Comorbidities in CKD patients

In CKD patients, the most common co morbidity identified was Systemic hypertension (44.26%) followed by diabetes mellitus (24.50%), congestive heart failure and sepsis (5.13%). Table 3.

Table 1: Frequency Distribution of Age.				
Age group	No. of subjects (n=152)	Percentage (%)		
<40 years	32	21.05		
40-60 years	68	44.73		
>60 years	52	34.21		
Total	152	100.00		

Table 2: Gender Distribution in CKD Patients.				
Gender	No. of subjects(n=152)	Percentage (%)		
Male	100	65.78		
Female	52	34.21		
Total	152	100.00		

Commonly prescribed drugs

The table provides data regarding the commonly prescribed drugs in the CKD patients. Among these, the most commonly prescribed drugs are hemantinics (18.16%) and vitamin supplements (18.16%). Table 4.

Table 3: Comorbidities in CKD Patients.						
Comorbidities	No. of patients	Percentage (%)				
Systemic hypertension	112	44.26				
Diabetes mellitus	62	24.50				
Congestive heart failure	13	5.13				
Cerebrovascular accident	7	2.76				
SLE	7	2.76				
IHD	5	1.97				
Hypothyroidism	6	2.37				
Sepsis	13	5.13				
Others	28	11.06				
Total	253	100.00				

Table 4: Commonly Pr	escribed Drug	s.
Category of drugs	No. of drugs	Percentage (%)
Beta blocker	33	2.22
Alpha blocker	31	2.09
CCB	74	4.99
$\underline{\alpha}$ -2 agonist	29	1.96
ACE I/ARB	40	2.70
Diuretics	26	1.75
Insulin	39	2.63
Hypoglycaemic agents	3	0.20
Vasodilators	40	2.70
Antiplatelet agents	44	2.97
Antibiotics	88	5.94
Antifungal	2	0.13
Inotropic agents	17	1.14
Haematinics	269	18.16
Antithyroid agents	13	0.87
Vitamin	269	18.16
Sodium bicarbonate	7	0.47
Potassium supplements	3	0.20
Calcium	81	5.46
Electrolytes	18	1.21
Phosphate binder	33	2.22
PPI	119	8.03
H2 receptor blocker	3	0.20
Antianginal	15	1.01
Statins	23	1.55
Steroids	12	0.81
Immunosuppressants	7	0.47
Probiotics	5	0.33
Others	138	9.31
Total	1481	100.00

Types of errors identified

In the control group, 72 total errors were identified. Among these transcribing errors was more (34.72%) followed by wrong dose (19.44%) and administering error (18.05%). Table 5. In test group 1, 44 total errors were identified. Among these, omission and administering errors (27.27%) were found to be more followed by wrong dose and wrong dosage form errors (11.36%). Table 7. In test group 2, 73 total errors were identified. Among these, administering error (24.65%) were found to be more, followed by transcribing error (19.17%), wrong dose and omission error (15.06%). Table 9. As per the study conducted by Landon, lack of information on discharge prescriptions about the medication prescribed, re-prescribed without any changes or discontinued during hospital stay was the main source of discrepancies. Findings from our study support the need for the multiple disciplinary practice of discharge and admission medication reconciliation. This process should clarify the status of medication the patient was taking prior to admission and it must also account for all new medications started in hospital, discontinued medications and adjusted medications to their dose.

Interactions identified

In the control group, total number of interactions identified was 196. Among these, significant interactions were found to be more (61.22%), followed by minor (27.55%) and serious (11.22%). Table 6. In test group 1, total number of interactions identified was 69. Among these, significant interactions were found to be more (53.62%), followed by minor (39.13%) and serious (7.24%) Table 8. In test group 2, total number of interactions identified was 77. Among the identified interactions, significant interactions were found to be more (62.33%), followed by minor (29.87%) and serious (7.79%) Table 10. According to a study by Varkey et al, which was split into two phases I: with only admission medication review and II: in hospital and discharge medication review, it was found that phase I had 4% serious, 47.7% significant and 48.4% minor drug interaction and it was found that in phase II the percentages of the same were 1.1%, 20.9% and 78% respectively. 12 As per Raghavan et.al study among 156 patients, 474 drug interactions were identified with 2.7 interactions per patient. Among 474 potential interactions, when classified according to resulting effect, the frequency of outcomes of drug-disease interactions was found that 150 (31.65%) may lead to hypotension or hypertension related to beta blockers and calcium channel blockers, 67 (19.14%) hypoglycaemia or hyperglycaemia associated insulin, and 25 (5.27%) hyperkalaemia associated with ACEI. This shows that medication reconciliation plays a significant role in identification of

drug related problems in hospitalised patients.

Medication reconciliation interventions

Medication reconciliation is a resource intensive process that targets efforts in ways that maximize impact on patient safety. Some studies show that errors during discharge are more, even though drug related problems didn't occur only due to lack of medication reconciliation but also due to inadequate information about the pharmacotherapy of the patient.⁴ Many studies show that drug use by the patient and what has been recorded

Table 5: Types of Group.	Errors Identified	in the Control	
Types of errors	No. of errors	Percentage (%)	
Prescribing	5	6.94	
Transcribing	25	34.72	
Administering	13	18.05	
Wrong drug	3	4.16	
Wrong strength	4	5.55	
Wrong dose	14	19.44	
Wrong time	1	1.38	
Omission	4	5.55	
Wrong dosage form	3	4.16	
Total	72	100.00	

Table 6: Interactions Identified in the Control Group.				
Type of interactions No. of errors Percentage (
Significant	120	61.22		
Serious	22	11.22		
Minor	54	27.55		
Total	196	100.0		

Table 7: Types of Errors Identified in the Test Group 1.				
Types of errors	No. of errors	Percentage (%)		
Prescribing	3	6.81		
Transcribing	3	6.81		
Administering	12	27.27		
Wrong drug	3	6.81		
Wrong dose	5	11.36		
Wrong time	1	2.27		
Omission	12	27.27		
Wrong dosage form	5	11.36		
Total	44	100.00		

Table 8: Interactions Identified in the Test Group 1.				
Type of interactions	No. of errors	Percentage (%)		
Significant	37	53.62		
Serious	5	7.24		
Minor	27	39.13		
Total	69	100.00		

Table 9: Types of Errors Identified in the Test Group 2.					
Types of errors	No. of errors	Percentage (%)			
Prescribing	6	8.21			
Transcribing	14	19.17			
Administering	18	24.65			
Wrong drug	5	6.84			
Wrong dose	11	15.06			
Wrong time	1	1.36			
Omission	11	15.06			
Wrong dosage form	7	9.58			
Total	73	100.00			

Table 10: Interactions Identified in the Test Group 2.					
Type of interactions No. of errors Percentage%					
Significant	48	62.33			
Serious	6	7.79			
Minor	23	29.87			
Total	77	100.00			

in the medication list is completely different Therefore, patient participation in medication reconciliation is very much important. That is why in our study we included patient counselling. But some studies showed that the impact of patient counselling towards the reconciliation process is still unclear. In this study, we recorded unintended discrepancies between home medication list and admission medication in the patients admitted to the nephrology unit. Nearly all the patients left the hospital with changes in their medication regimen and didn't receive any written instructions highlighting the changes. An effective system must be implemented to ensure adequate patient education and communication between the providers. From the Annals of Pharmacotherapy, edition 2003 we have adopted PCNE version 4 medication reconciliation chart.6

Group 1 vs group 2

Comparing the two test groups, no significant results

Table 11: Medication Reconciliation Interventions of Control Group Vs Test Group 1.						
Classification	Control			Test Group 1		
	M	SD	M	SD	р	
Prescription-related interventions: correction of unintended discre	pancies					
Start	0.5395	0.70125	0.6842	0.61972	0.283	
Dosage/schemes	0.5789	0.59471	0.6316	0.54132	0.647	
Switch:-	0.000	0.000	0.6053	0.54720	0.000***	
Stop:-	0.2237	0.41948	0.6316	0.63335	0.000***	
Prescription-related interventions: optimization of pharmacotherap	ру					
Start:-	0.0132	0.11471	0.6316	0.48885	0.000***	
Dosage/schemes	1.1184	0.74775	1.0526	0.76925	0.662	
Switch	0.7105	0.64943	0.8421	0.59395	0.297	
Stop	0.000	0.000	0.4737	0.55687	0.000***	
Patient medication-handling interventions: improvement of medic	ation use by	the patient.				
Inappropriate medication use	0.000		0.9474	0.83658	0.000***	
Answering questions on medications	0.000	0.000	0.2895	0.56511	0.000***	
Medication supply	0.000	0.000	0.2895	0.56511	0.000***	

N=76Patients per control group. N=38 patients per test group 1, P<0.05=*, P<0.01=**, P<0.001=***, 95% CI, Calculated by using independent t-test.

were obtained with respect to prescription related interventions on correction of unintended discrepancies. In prescription related intervention for the optimization of pharmacotherapy, highly significant results were recorded in 'dosage/schemes'. With patient medication handling interventions, significant results were found in test group 2 compared to that of test group 1 (p=0.000, p<0.005)

Control vs test group

Prescription related interventions; correction of

unintended discrepancies by comparing the control and the test groups significant discrepancies were identified in test 'switch' and 'stop' p<0.05. With prescription related intervention; optimization of pharmacotherapy significant discrepancies was identified with 'start' and 'stop' in the test group compared to the control p=0.001 (p<0.05). With patient education handling interventions; improvement of medication use by the patient significant difference between the two groups were observed. The patients were adequately provided with appropriate use of medication instructions, medication supply and by

Table 12: Medication Reconciliation Interventions of Test Groups (Group 1 VS Group 2).					
Classification	TEST (group 1)		TEST (group 2)		
	М	SD	M	SD	р
Prescription-related interventions: correction of unintended dis-	crepancies				
Start	0.6842	0.61973	0.6842	0.90360	0.999
Dosage/schemes	0.6316	0.54132	0.5526	0.68566	0.579
Switch:-	0.6053	0.54720	0.7105	0.69391	0.468
Stop:-	0.6316	0.63335	0.4211	0.59872	0.141
Prescription-related interventions: optimization of pharmacothe	erapy				
Start:-	0.6316	0.48885	0.7895	0.77661	0.292
Dosage/schemes	1.0526	0.76925	1.5789	0.79293	0.004**
Switch	0.8421	0.59395	0.9211	0.63167	0.576
Stop	0.4737	0.55687	0.3684	0.48885	0.384
Patient medication–handling interventions: improvement of medication use by the patient.					
Inappropriate medication use	0.9474	0.8684	1.8421	1.17465	0.000***
Answering questions on medications	0.8684	0.87522	2.6053	1.22009	0.000***
Medication supply	0.2865	0.56511	1.9211	1.47743	0.000***

N=38 Patients per group. P<0.05=*, P<0.01=**, P<0.001=***, 95% CI, Calculated by using independent t-test.

Table 13: Patient Counselling Effect of Test Groups (Group 1 and Group 2).					
Classification	Baseline		After counselling		
	M	SD	M	SD	P
Patient counselling effect					
Group 1(no counselling done)	4.8684	1.77324	5.1579	2.03385	0.010**
Patient counselling effect					
Group 2	5.0263	1.95195	10.1842	2.52400	0.000***

N=38 Patients per group. P<0.01=**, P<0.001=***, 95% CI, Calculated by using paired t-test.

answering all the questions posed by the patients in the test group resulting in significant end results.

Patient counselling effect of test groups

(Group 1 and Group 2)

The knowledge of patients was assessed during baseline review and final review of both group 1 and group 2. Comparing mean knowledge between baseline and final review showed statistically significant (p<0.05). The significance was more in test group 2 when compared to group 1. This change in the significance is highly due to counseling done. Viktil *et al.* showed that, in hospitalized patients, a mean of 4.4 DRPs per patient were recorded due to patient counseling versus a mean of 2.4 DRPs due to using medication records and participating in multidisciplinary team discussions. Viktil *et al.* recorded significantly more interventions after patient counseling in the "need for additional drug" and "therapy discussion" categories, which is consistent with our study. These observations point out the impact

Table 14: Knowledge Assessment Scoring for each question after Patient Counselling- Group 1.

•			•	
Questions	No. of 'yes' answer	Percentage	No. of 'no' answer	Percentage
Qs1	30	78.94	8	21.05
Qs2	21	55.26	17	44.73
Qs3	18	47.36	20	52.63
Qs4	8	21.05	30	78.94
Qs5	7	18.42	31	81.57
Qs6	19	50.00	19	50.00
Qs7	21	55.26	17	44.73
Qs8	0	0	38	100.00
Qs9	13	34.21	25	65.78
Qs10	13	34.21	25	65.78
Qs11	0	0	38	100.00
Qs12	31	81.57	7	18.42
Qs13	4	10.52	34	89.47
Qs14	2	5.26	36	94.73
Qs15	22	57.89	16	42.10

Table 15: Knowledge Assessment Scoring for each question after Patient Counselling- Group 2.

Questions	No. of 'yes' answer	Percentage	No. of 'no' answer	Percentage
Qs1	37	97.36	1	2.63
Qs2	32	84.21	6	15.78
Qs3	21	81.57	17	44.73
Qs4	23	60.52	15	39.47
Qs5	23	60.52	15	39.47
Qs6	34	89.47	4	10.52
Qs7	29	76.31	9	23.68
Qs8	19	50.00	19	50.00
Qs9	26	68.42	12	31.57
Qs10	34	89.47	4	10.52
Qs11	15	39.47	23	60.52
Qs12	34	89.47	4	10.52
Qs13	25	65.78	13	34.21
Qs14	10	26.31	28	73.68
Qs15	35	92.10	3	7.89

Table 16: Know	ladaa /	Assessment Questionnaire.	
Table to: Know	iedae <i>i</i>	assessment Questionnaire.	

Table '	16: Knowledge Assessment Questionnaire.
Qs no	Questions
1	Do you know your medical condition or disorder?
2	Do you know the signs and symptoms of your medical condition, disease or disorder?
3	Do you know the total number of medications in your treatment chart?
4	Do you know which medication is used for what purpose in your medication treatment chart?
5	Do you know the dose of your medicines?
6	Do you know the frequency per day of your medications?
7	Do you know which medications should be taken before or after food?
8	Do you know what to do incase you missed a dose?
9	Do you know the storage conditions of your medicines?
10	Do you know which food to avoid?
11	Do you know what to do in case you take a double dose by mistake?
12	Do you know the route of administration of every medicine in your treatment chart?
13	Do you know the serious side effects of your medicines?
14	Do you know which medicines should not be taken together?
15	Do you know the personal care and preventions

of patient counseling at regular intervals along with the pharmacotherapy to improve the quality of life in CKD and HD patients.

regarding your disease?

Table 17: Medication Alert Card Questionnaire Scoring for Each Question-Group 2.

Questions	No. of 'yes' answer	Percentage	No. of 'no' answer	Percentage
Qs1	0	0	38	100
Qs3	37	97.37	1	2.63
Qs4	12	31.58	26	68.42
Qs5	26	68.42	12	31.58
Qs6	22	57.89	16	42.10
Qs7	33	86.84	5	13.16
Qs8	34	89.47	4	10.53
Qs9	31	81.58	7	18.42
Qs11	5	13.16	33	86.84

Table 18: Medication Alert Card Questionnaire.

Qs no	Questions
1	Have you used a medication alert card before?
3	Was this medication alert card cost saving?
4	Were additional tests done in spite of the medication alert card?
5	Has the medication alert card been life-saving during emergency?
6	Has the emergency number been contacted?
7	Were any updates made entered in the medication alert card?
8	Was the medication alert card checked during refilling of medications/change in medication?
9	Was the alert card portable?
11	Was it misplaced anytime in the past?

Table 19: Response Regarding Storage of Alert Card.

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Storage	Percentage
Home	21.05
Work	2.63
Purse	0
Vehicle	15.78
Dialysis note	60.52

Medication alert card

In this study, frequently updated medication alert card was provided to the second arm of the cases to in this check their involvement in reducing the potential errors and thus improving the quality of medical care. Medication alert card were distributed to all the 38 cases and the same were frequently updated. Among the results it was shown that all 38 cases (100%) had not used such alert cards any time in the past. 37 cases (97.37%) found it to be cost saving as no additional test was needed during outside admissions. 22 cases (57.89%) found the emergency number in the card to be effective. 33 cases (86.84%) claimed the alert card to be updated frequently

and 34 cases (89.47%) claimed its efficacy during refills. 31 cases (81.58%) found it to be portable. 23 cases (60.52%) stored their card in the dialysis note and 33 cases (86.84%) claimed that it was not misplaced in the past.

LIMITATIONS OF THE STUDY

The study was not focused to see the impact of interventions on patients even though the need was present. Duration of the study was limited to provide appropriate interventions to minimize the DRPs identified due to medication reconciliation. The interventions would have been more accurate if compared with interventions done by other healthcare professionals.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

ABBREVIATIONS USED

CKD: Chronic kidney disease, ESRD: End stage renal disease, DRP: drug related problems, JCAHO: Joint committee accreditation health care organization, ADE: Adverse drug events, MOR: Medication order review, TCR: Treatment chart review, ADR: Adverse drug reactions, PCNE: Pharmaceutical care network Europe, HD: Hemodialysis, IHD: Ishemic heart disease, SLE: Systemic lupus erythematosus.

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