Published online 2022 June 22.

Research Article

Utility of Citrate Dialysis in Patients with Contraindication for Heparin in a Limited-Resource Setting

Rajaram Jagdale^{1, 2, 3, *}, Alan Almeida¹, Jatin Kothari¹, Rasika Sirsat¹, Supriya Surwase⁴ and Dixon Thomas⁵

¹Department of Nephrology, PD Hinduja National Hospital and Medical Research Centre, Mumbai, India

²Department of Nephrology, Thumbay University Hospital, Ajman, United Arab Emirates

³College of Medicine, Gulf Medical University, Ajman, UAE

⁴Microbiology, Thumbay Hospital, Dubai, United Arab Emirates

⁵College of Pharmacy, Gulf Medical University, Ajman, United Arab Emirates

Corresponding author: Nephrology Department, Thumbay University Hospital, Ajman, UAE. Tel: +971-568547247, Email: rajaramjagdale@gmail.com

Received 2022 March 20; Accepted 2022 April 18.

Abstract

Background: Hemodialysis among critical care patients with acute kidney injury (AKI) is challenging, especially if heparin is contraindicated.

Objectives: This study assessed the utility of citrate dialysis for such patients in a limited-resource setting.

Methods: In this prospective study, patients were divided into group A (heparin-free saline flush dialysis), group B (heparin-free citrate dialysis without flushing), and group C (heparin-free citrate dialysis with flushing). The subjects underwent completed sustained low-efficiency daily dialysis (blood flow = 150 mL/minute, dialysate = 300 mL/minute) or intermittent hemodialysis (blood flow = 250 mL/minute, dialysate flow = 500 mL/minute). Statistical tests using SPSS software (version 26) were used to determine safety and effectiveness differences.

Results: Among 25 patients studied with multiple hemodialysis sessions, blood flow and dialysate flow were observed to be better in heparin-free citrate dialysis with flushing. There were further advantages of lesser dialyzer clotting and more reuse of dialyzers. Metabolic differences were insignificant. Heparin-free citrate dialysis with or without flushing was equally effective and safe, compared to heparin-free saline flush dialysis, in patients with or without liver impairment.

Conclusions: Citrate dialysis is observed to be a safe and effective alternative to heparin-free saline flushing dialysis in intensive care unit patients with AKI. More such studies are required in limited-resource settings to utilize citrate dialysis in patients with heparin contraindication.

Keywords: Citrate Dialysis, Heparin Contraindication, Bleeding, Acute Kidney Injury

1. Background

Acute kidney injury (AKI) patients in the intensive care unit (ICU) are complex to manage (1). Despite advances in renal replacement therapy (RRT) and critical care in recent years, the mortality rate remained high (2, 3). The risk, injury, failure, loss of kidney function, and end-stage renal failure (RIFLE) criteria consist of three graded levels of injury (i.e., risk, injury, and failure) based upon either the magnitude of the rise in serum creatinine or fall in urine output and two outcome measures (i.e., loss of kidney function and end-stage kidney disease [ESKD]) (4).

Heparin is the conventional anticoagulant used during hemodialysis. When there are contraindications for heparin use, namely perioperative period, active bleeding, deranged coagulation profile, or heparin-induced thrombocytopenia, it is necessary to resort to heparin-free dialysis in AKI patients admitted to the ICU. Dialyzer clotting is common in patients undergoing heparin-free dialysis (5, 6).

Citrate dialysis, where citric acid functions as an anticoagulant, is an alternative to heparin-free dialysis with advantages, such as less dialyzer clotting, more dialyzer reuse, and a better-delivered dose of dialysis. Citrate dialysate has been successfully used in regional citrate anticoagulation (RCA) and has been more effective than repeated saline flushing of the extracorporeal circuit (7). Citrate accumulation/toxicity due to failure to metabolize citrate, especially in chronic liver disease patients, is a rare problem due to the low concentration of citrate used in the dialysate. Citrate dialysate has been used for sustained

Copyright © 2022, Author(s). This is an open-access article distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 International License (http://creativecommons.org/licenses/by-nc/4.0/) which permits copy and redistribute the material just in noncommercial usages, provided the original work is properly cited.

low-efficiency daily dialysis (SLEDD) without any evidence of citrate accumulation or development of hypocalcemia in the presence of severe liver dysfunction (8, 9).

2. Objectives

The current study aimed to assess the safety and effectiveness of citrate dialysis as an alternative to heparin-free saline flushing dialysis in ICU patients with AKI.

3. Methods

This prospective study was carried out at a tertiary care hospital in Mumbai, India. The patients were selected from those admitted to the ICU with AKI and the need for heparin-free dialysis.

3.1. Inclusion Criteria

(1) Age >18 years

(2) Patients with AKI or acute, chronic kidney disease requiring heparin-free dialysis in the ICU with one or more of the following criteria:

(a) Patients with active bleeding

(b) Platelet count < 50,000 and/or international normalized ratio > 1.5 or partial thromboplastin time > 50% above the control value

(c) Pericarditis

(d) Perioperative patients for whom heparin could not be used.

3.2. Exclusion Criteria

(1) Patients requiring heparin-free dialysis but for whom heparin or other anticoagulant was used to treat other conditions (e.g., probable or proven deep vein thrombosis, pulmonary embolism, and unstable angina).

(2) Patients requiring continuous renal replacement therapy

(3) Hepatitis B, C, or human immunodeficiency viruspositive patients

(4) Patients who were unwilling to participate in the study

3.3. Criteria for Data Determination

After obtaining written, valid, informed consent and fulfilling selection criteria, the patients were enrolled in the study and randomized into one of the three groups. The data were collected for 6 months in 2011. Group A (heparin-free saline flush dialysis, 100 mL saline flushing every 15 minutes), group B (heparin-free citrate dialysis

without flushing), and group C (heparin-free citrate dialysis with flushing, 200 mL saline flush every hour) underwent completed SLEDD (blood flow = 150 mL/minute, dialysate = 300 mL/minute) or intermittent hemodialysis (IHD) (blood flow = 250 mL/minute, dialysate flow = 500 mL/minute).

A Citrasate® solution was used in the present study. It was similar to the bicarbonate dialysis solution, except for having 2.4 mEq/L citrate and acetic acid of 0.3 mEq/L instead of 4 mEq/L in the bicarbonate dialysis solution. A small amount of citrate (0.8 mmol/L, i.e., 2.4 mEq/L) was used in comparison to that of RCA (4 mmol/L). This solution was used instead of acetic acid as the acidifying agent. This citrate binds with calcium and inhibits blood coagulation at the dialyzer membrane surface, resulting in better dialyzer clearance, less dialyzer clotting, and increased reuse. Citrasate® concentrate is available in all standard concentrations and formulations, requiring it to be poured into a standard A concentrate container and attached to the dialysis system. Therefore, no additional patient or system monitoring is needed beyond those typically employed in standard dialysate formulations treatment (10).

All dialysis sessions were carried out with Fresenius 4008S machines, and F6 dialyzers were used for all patients (11). The fiber bundle volume (FBV) was calculated (by machine) after each dialysis session, and a dialyzer with FBV of > 80% of the baseline value of that session was reused for the next session. The present study was approved by the Ethics Committee of PD Hinduja National Hospital and Medical Research Centre, Mumbai, India.

Completed treatment was defined as completing treatment duration as ordered (4 or 6 hours). The clotting of the dialyzer was defined as the early discontinuation of dialysis, beyond 30 minutes, prior to the prescribed time due to circuit clotting in either the lines, chambers, or dialyzer itself (45, 46). The Simplified Acute Physiology Score II [SAPS II] was calculated for each patient after 24 hours of ICU stay (12).

3.4. Data Analysis

Statistical methods used in this study were Pearson's chi-square test, student t-test, analysis of variance (ANOVA), and Mann-Whitney U test. Quantitative data were represented in the form of mean \pm standard deviation or median (interquartile range) as per the distribution of the variable, with the former used in cases of data with normality and the latter in the absence of the same. Accordingly, depending upon the normality status, a comparison of quantitative variables by variables with two subgroups was made using an unpaired t-test or Mann-Whitney U test, respectively. The comparison of quantitative data between more than two subgroups of a nominal variable was made using one-way ANOVA. If a significant difference was observed, an appropriate posthoc test was applied for pairwise comparison. The data were analyzed using SPSS software (version 26).

4. Results

As shown in Table 1, critically ill patients in the ICU with AKI requiring heparin-free dialysis were included in this study (n = 25). Moreover, 17 (68%) and 8 (32%) patients were male and female, respectively. Deranged coagulation profile (laboratory abnormalities), external or internal bleeding and perioperative period were the indications for inclusion in the study in 21 (80%), 3 (12%), and 1 (4%) patients, respectively. Based on the results, 24 (96%) and 1 (4%) patients were in the failure and injury stages, as per the RI-FLE criterion for AKI, respectively. Medical, surgical, and obstetric causes of AKI were present in 20 (80%), 4 (16%), and 1 (4%) patients, respectively. Acute tubular necrosis (ischemic or toxic) was the most common cause of AKI (48%), followed by sepsis (44%) and prerenal (8%) issues.

The age of the patients ranged from 26 to 86 years, with the elderly (media = 63) reported as the majority of the study population. The mean age of group A (63 years), group B (60 years), and group C (61 years) did not vary statistically, with a p-value of 0.936. The median SAPS II score among the study population was 63. The patients in group A had a lower SAPS II score than group B (P=0.0059); however, the difference was insignificant in other comparisons. In 198 dialysis sessions for groups A-C, 63% and 37% of the sessions were SLEDD and IHD, respectively. The IHD was the lowest in group B (6%), in which 94% of the patients received SLEDD.

The groups differed in blood flow and dialysate flow based on ANOVA. In addition, Tukey's posthoc analysis revealed differences between groups A and B and groups B and C for both blood and dialysate flow (more details in Table 2).

Table 3 shows that although there was no significant difference in dialyzer clotting among the three groups, the number of dialyzer clotting in group C was less than B and in B less than A.

As shown in Table 4, the average reuse of the dialyzer was higher in group B > A(P = 0.018); nevertheless, the difference was not significant in other comparisons.

The percentage of incomplete and complete treatments was not different in the three groups. Dialysis needed to be terminated before completion during 11 out of 198 sessions (5.6%), among which dialyzer clotting and persistent hypotension were the indications for the termination in 8 and 3 sessions, respectively. The groups did not differ significantly in fall in pre- to postdialysis FBV (P > 0.5).

The difference in the urea reduction ratio (URR) among the three groups was not significant, indicating that the delivered dose of dialysis (URR in this study) was not different in the three groups. However, the rise in pre- to postdialysis bicarbonate in all the three groups was significant (P < 0.01). Table 5 shows the differences in ionized calcium among the three groups.

The bicarbonate and ionized calcium in patients without liver dysfunction showed a significant improvement after dialysis; nonetheless, there was no significant difference in patients with liver dysfunction. The mortality rate of critically ill patients requiring dialysis for AKI in the ICU was 40% in the present study. There was no significant difference among the three groups in this regard.

5. Discussion

Numerous patients in the ICU with AKI need RRT, and the mortality rate is still high despite advances in RRT and critical care. Extracorporeal circuit clotting in patients who need heparin-free dialysis can be prevented by either RCA or a low concentration of citric acid. A bicarbonate dialysis solution with a low concentration (0.8 mmol/L) of citric acid is better than heparin-free saline flush dialysis.

In the present study, there were fewer dialyzer clotting episodes with citrate dialysis without flushing than saline flush dialysis (group B < A); however, it was not significant, which could be due to the smaller number of dialysis sessions in the study. Madison et al. demonstrated significantly fewer dialyzer clotting episodes in citrate without flushing than in saline flushing p (13). The present study showed that the average reuse of dialyzer was significantly higher in citrate without flushing than in saline flush (B > A) group, which could be explained by less dialyzer clotting in group B. Dialyzer reuse is recommended in limitedresource countries (14).

There was no difference in the percentage of completed treatments among the three groups. The delivered dose of the dialysis (URR in this study) was comparable in the three groups. The present study showed a significant rise in postdialysis bicarbonate levels in all three groups; nevertheless, the values remained normal. There was a postdialysis rise in ionized calcium in the citrate group (groups B and C) in the present study. No episode of clinically significant hypocalcemia was noted in the present study population, indicating the safety of citrate dialysis, precluding the need to monitor ionized calcium. Hypocalcemia with citrate dialysis is reported in other studies (15, 16). The present study demonstrated that citrate dialysis

Table 1. Baseline Characteristics of Patients (n = 25)								
Variables	No.	Mean \pm Standard Deviation	Minimum	Maximum	Median			
Age (y)	25	61.40 ± 14.47	26.00	86.00	63.00			
Systolic blood pressure (mmHg)	25	129.38 ± 22.23	86.00	180.00	126.00			
Diastolic blood pressure (mmHg)	25	66.50 ± 13.76	40.00	100.00	70.00			
Simplified Acute Physiology Score II	25	60.24 ± 17.71	29.00	97.00	63.00			
Number of inotropes	25	1.00 ± 0.91	0.00	3.00	1.00			
Bicarbonate (mEq/L)	25	18.47± 5.04	6.00	26.00	18.00			
Hemoglobin (gm%)	25	9.39 ± 2.84	6.00	16.00	8.85			
Platelets (cmm)	25	71807.69±66093.28	14000.00	264000.00	46500.00			
International normalized ratio	18	1.72 ± 0.62	1.20	3.80	1.46			

Table 2. Comparison of Variables (Blood and Dialysate Flow) in Three Groups

Group	No.	$\textbf{Mean} \pm \textbf{Standard deviation}$	Median	P-Value	Pairwise Comparison ^{a, b}	P-Value
Blood flow (mL/min)				< 0.0		
А	72	191.67 ± 49.65	150.00		A vs. B	< 0.05
В	66	159.09 ± 28.97	150.0; 150.00; 100		A vs. C	> 0.05
С	60	213.33 ± 48.60	250.00		B vs. C	< 0.05
Dialysate flow (mL/min)				< 0.001		
А	72	383.33 ± 99.29	300.00		A vs. B	< 0.05
В	66	318.18 ± 57.94	300.00		A vs. C	> 0.05
C	60	426.67±97.19	500.00		B vs. C	< 0.05

Abbreviations: A, heparin-free saline flush; B, heparin-free citrate dialysis without flushing; C, heparin-free citrate dialysis with flushing. ^a One-way analysis of variance ^b Tukey's posthoc test for multiple comparisons

Dialycer Clotting			Total		
Dialyser clotting	-	Α	В	С	-
Yes	n	25 (34.7)	16 (24.2)	11 (18.3)	52 (26.3)
No	n	47(65.3)	50 (75.8)	49 (81.7)	146 (73.7)
Total	n	72 (100)	66 (100)	60 (100)	198 (100)
Pearson chi-square test	Value: 4.748	df: 2	P-value 0.093	No significant association	

Abbreviations: A, heparin-free saline flush; B, heparin-free citrate dialysis without flushing; C, heparin-free citrate dialysis with flushing. ^aValues are expressed as No. (SD).

Fable 4. Comparison of Number of Reuses of Dialyzers in Three Groups								
Variable	Group	n	Mean ± Standard Deviation	Median	Р	Pairwise Comparison ^{a, b}	P-Value	
	А	72	1.89 ± 0.99	2.00	0.02 (significant difference)	A vs. B	0.018	
Number of reuses	В	66	2.59 ± 2.02	2.00		A vs. C	0.146	
	С	60	2.38 ± 1.34	2.00		B vs. C	0.719	

Abbreviations: A, heparin-free saline flush; B, heparin-free citrate dialysis without flushing; C, heparin-free citrate dialysis with flushing.

^a One-way analysis of variance ^b Tukey's posthoc test for multiple comparisons

Table 5	rable 5. Comparison of Pre- to Posthemodialysis Ionized Calcium Differences (mmol/L) in Groups B and C							
Grou	ıp	No.	Mean \pm Standard Deviation	P-Value ^a				
В				0.005				
	Prehemodialysis ionized calcium (mmol/L)	13	4.04 ± 0.39					
	Posthemodialysis ionized calcium (mmol/L)	13	4.22 ± 0.40					
С				0.003				
	Prehemodialysis ionized calcium (mmol/L)	15	4.18 ± 0.47					
	Posthemodialysis ionized calcium (mmol/L)	15	4.43 ± 0.39					

^a Paired *t*-test

could be safely used in patients with or without liver failure with no excess risk of hypocalcemia or metabolic alkalosis. Such metabolic issues are reported with RCA (17-20).

The overall mortality of the study population was 40% in this study. Other studies showed a mortality rate of 30-50% in AKI patients requiring dialysis (21-23). The present study showed that citrate dialysis could be performed with either no or infrequent flushing, reducing nursing time and requiring less ultrafiltrate removal, which might be vital in patients with poor left ventricular function in comparison to the saline flushing group.

5.1. Conclusions

It can be concluded that citrate dialysis appears to be a safe and effective alternative to heparin-free saline flushing dialysis in ICU patients with AKI, even in patients with liver failure. However, a more extensive prospective study with a large sample size needs to be performed to validate the findings of this pilot study. In addition, a similar study also needs to be performed in patients with ESKD on maintenance hemodialysis to determine whether the results of the present study could be extrapolated to that population.

Footnotes

Authors' Contribution: Study concept and design: RJ, AA, JK, and RS; Analysis and interpretation of the data: RJ, AA, JK, RS, and SS; Drafting of the manuscript: RJ and DT; Critical revision of the manuscript for important intellectual content: RJ, AA, JK, RS, SS, and DT; Statistical analysis: RJ and SS (with statisticians' help).

Conflict of Interests: The authors declare no conflict of interest.

Data Reproducibility: All the data collected are presented in this manuscript. Any additional information can be provided on request from the corresponding author by this journal representative at any time during submission or after publication. Otherwise, all the consequences of possible withdrawal or future retraction will be with the corresponding author.

Ethical Approval: This study was approved by the Institutional Review Board of PD Hinduja National Hospital and Medical Research Centre, Mumbai, India (ref no.: 2010/17). It was an MD thesis at the study site.

Funding/Support: No fund of any kind was received as part of this study or its publication.

Informed Consent: Informed consent was obtained from all the patients before enrolling in the study.

References

- Garzotto F, Piccinni P, Cruz D, Gramaticopolo S, Dal Santo M, Aneloni G, et al. RIFLE-based data collection/management system applied to a prospective cohort multicenter Italian study on the epidemiology of acute kidney injury in the intensive care unit. *Blood Purif*. 2011;31(1-3):159–71. [PubMed ID: 21228585]. https://doi.org/10.1159/000322161.
- Park WY, Hwang EA, Jang MH, Park SB, Kim HC. The risk factors and outcome of acute kidney injury in the intensive care units. *Korean J Intern Med*. 2010;25(2):181-7. [PubMed ID: 20526392]. [PubMed Central ID: PMC2880692]. https://doi.org/10.3904/kjim.2010.25.2.181.
- Yohannes S, Chawla LS. Evolving practices in the management of acute kidney injury in the ICU (Intensive Care Unit). *Clin Nephrol.* 2009;**71**(6):602–7. [PubMed ID: 19473627]. https://doi.org/10.5414/cnp71602.
- Ricci Z, Cruz D, Ronco C. The RIFLE criteria and mortality in acute kidney injury: A systematic review. *Kidney Int.* 2008;73(5):538–46. [PubMed ID: 18160961]. https://doi.org/10.1038/sj.ki.5002743.
- Vandenbosch I, Dejongh S, Claes K, Bammens B, De Vusser K, Van Craenenbroeck A, et al. Strategies for asymmetrical triacetate dialyser heparin-free effective haemodialysis: the SAFE study. *Clin Kidney J.* 2021;14(8):1901–7. [PubMed ID: 34345413]. [PubMed Central ID: PMC8323132]. https://doi.org/10.1093/ckj/sfaa228.
- Faguer S, Saint-Cricq M, Nogier MB, Labadens I, Lavayssiere L, Kamar N, et al. Heparin-Free Prolonged Intermittent Hemodialysis Using Calcium-Free Citrate Dialysate in Critically III Patients. Crit Care Med. 2017;45(11):1887-92. [PubMed ID: 28857854]. https://doi.org/10.1097/CCM.00000000002694.
- Kozik-Jaromin J, Nier V, Heemann U, Kreymann B, Bohler J. Citrate pharmacokinetics and calcium levels during high-flux dialysis with regional citrate anticoagulation. *Nephrol Dial Transplant*. 2009;24(7):2244–51. [PubMed ID: 19196824]. [PubMed Central ID: PMC2698091]. https://doi.org/10.1093/ndt/gfp017.

- Zhang W, Bai M, Yu Y, Li L, Zhao L, Sun S, et al. Safety and efficacy of regional citrate anticoagulation for continuous renal replacement therapy in liver failure patients: a systematic review and metaanalysis. *Crit Care*. 2019;23(1):22. [PubMed ID: 30678706]. [PubMed Central ID: PMC6345001]. https://doi.org/10.1186/s13054-019-2317-9.
- Karkar A, Ronco C. Prescription of CRRT: A pathway to optimize therapy. Ann Intensive Care. 2020;10(1):32. [PubMed ID: 32144519]. [PubMed Central ID: PMC7060300]. https://doi.org/10.1186/s13613-020-0648-y.
- Cheng YL, Yu AW, Tsang KY, Shah DH, Kjellstrand CM, Wong SM, et al. Anticoagulation during haemodialysis using a citrate-enriched dialysate: A feasibility study. *Nephrol Dial Transplant*. 2011;26(2):641–6. [PubMed ID: 20615906]. https://doi.org/10.1093/ndt/gfq396.
- Zhou YL, Liu HL, Duan XF, Yao Y, Sun Y, Liu Q. Impact of sodium and ultrafiltration profiling on haemodialysis-related hypotension. *Nephrol Dial Transplant*. 2006;21(11):3231-7. [PubMed ID: 16954178]. https://doi.org/10.1093/ndt/gfl375.
- Lopes JA, Jorge S, Resina C, Santos C, Pereira A, Neves J, et al. Acute renal failure in patients with sepsis. *Crit Care*. 2007;11(2):411. [PubMed ID: 17466080]. [PubMed Central ID: PMC2206468]. https://doi.org/10.1186/cc5735.
- Madison J, Llumin M, Chin A. Citrate-containing dialysate is well tolerated during slow extended daily dialysis in the ICU. 38th Annual Renal Week Meeting. Philadelphia, Penn., USA. American Society of Nephrology; 2005.
- Manandhar DN, Chhetri PK, Tiwari R, Lamichhane S. Evaluation of dialysis adequacy in patients under hemodialysis and effectiveness of dialysers reuses. *Nepal Med Coll J.* 2009;**11**(2):107-10. [PubMed ID: 19968150].
- Durao MS, Monte JC, Batista MC, Oliveira M, Iizuka IJ, Santos BF, et al. The use of regional citrate anticoagulation for continuous venovenous hemodiafiltration in acute kidney injury. *Crit Care Med.* 2008;**36**(11):3024–9. [PubMed ID: 18824904]. https://doi.org/10.1097/CCM.0b013e31818b9100.
- 16. Gubensek J, Buturovic-Ponikvar J, Ponikvar R. Regional citrate anticoagulation for single-needle hemodialysis: a prospective clin-

ical study. Blood Purif. 2007;25(5-6):454-6. [PubMed ID: 18057876]. https://doi.org/10.1159/000111808.

- Stucker F, Ponte B, Tataw J, Martin PY, Wozniak H, Pugin J, et al. Efficacy and safety of citrate-based anticoagulation compared to heparin in patients with acute kidney injury requiring continuous renal replacement therapy: a randomized controlled trial. *Crit Care*. 2015;**19**:91. [PubMed ID: 25881975]. [PubMed Central ID: PMC4364313]. https://doi.org/10.1186/s13054-015-0822-z.
- Slowinski T, Morgera S, Joannidis M, Henneberg T, Stocker R, Helset E, et al. Safety and efficacy of regional citrate anticoagulation in continuous venovenous hemodialysis in the presence of liver failure: the Liver Citrate Anticoagulation Threshold (L-CAT) observational study. *Crit Care*. 2015;**19**:349. [PubMed ID: 26415638]. [PubMed Central ID: PMC4587580]. https://doi.org/10.1186/s13054-015-1066-7.
- Bianchi NA, Altarelli M, Eckert P, Schneider AG. Complications of Regional Citrate Anticoagulation for Continuous Renal Replacement Therapy: An Observational Study. *Blood Purif.* 2020;**49**(5):567-75. [PubMed ID: 32126564]. https://doi.org/10.1159/000506253.
- Lahmer T, Messer M, Rasch S, Beitz A, Schnappauf C, Schmid RM, et al. Sustained low-efficiency dialysis with regional citrate anticoagulation in medical intensive care unit patients with liver failure: A prospective study. J Crit Care. 2015;30(5):1096-100. [PubMed ID: 26254678]. https://doi.org/10.1016/j.jcrc.2015.06.006.
- Sakhuja A, Kumar G, Gupta S, Mittal T, Taneja A, Nanchal RS. Acute Kidney Injury Requiring Dialysis in Severe Sepsis. *Am J Respir Crit Care Med.* 2015;**192**(8):951-7. [PubMed ID: 26120892]. https://doi.org/10.1164/rccm.201502-0329OC.
- Kolhe NV, Muirhead AW, Wilkes SR, Fluck RJ, Taal MW. National trends in acute kidney injury requiring dialysis in England between 1998 and 2013. *Kidney Int.* 2015;88(5):1161–9. [PubMed ID: 26221750]. https://doi.org/10.1038/ki.2015.234.
- Levey AS, James MT. Acute Kidney Injury. Ann Intern Med. 2017;167(9):ITC66–80. [PubMed ID: 29114754]. https://doi.org/10.7326/AITC201711070.