# A study to evaluate sepsis and its markers in renal failure patients on hemodialysis: an observational study

# Madhav Gopal Agarwal\*

Principal Specialist, M.D. (General Medicine), J.L.N. Medical college and associated group of Hospitals Ajmer, Rajasthan, India

\*Corresponding author: Madhav Gopal Agarwal, Principal Specialist, M.D. (General Medicine), J.L.N. Medical college and associated group of Hospitals Ajmer, Rajasthan, India

#### **ABSTRACT**

Aim: To evaluate sepsis and its markers in renal failure patients on hemodialysisnand its correlation of hematological abnormalities.

**Material & Methods:** The present study was conducted for the period of one year and total of 50 patients of both sexes who were diagnosed as case of renal failure which include both acute kidney injury (AKI) and CKD on basis of clinical history, examination, biochemical markers and were advised for hemodialysis were included in the study.

**Results:** In our study among 50 patients of renal failure on hemodialysis the mean age in our study was 46.64±12.48 years with 32 male patients. Out of 50 patients 10 (20%) had positive blood and catheter tip culture and 40 (80%) of patients had negative blood and catheter tip culture. Out of 10 patients with sepsis 2 (20%) were in the age group between 15–25 years, 2 (20%) were in the age group between 26–35 years, 1 (10%) were in the age group 36–45 years and 5 (50%) were above 45 years of age. All 10 (100%) patients had episode of fever with chills and rigor, 4 (40%) patients had redness and pain at hemodialysis catheter site, 3 patients (30%) were confused, disoriented or comatose and 2 (20%) patients had hypotension. Among 10 patients of renal failure with sepsis, none had TLC less than 4.8/cumm (leucopenia), 2 (20%) patients had count between 4.8–10.8/cumm and 8 (80%) patients had TLC more than 10.8/cumm.

**Conclusion:** Patients requiring hemodialysis, who are having non modifiable risk factors like age, sex other risk factors for infection should be controlled to reduce incidence of infection.

Keywords: Sepsis, Chronic kidney disease, Hemodialysis, Blood stream infection

#### 1. INTRODUCTION

Severe bacterial infection leading to sepsis is an important cause of hospitalization and loss of health worldwide. <sup>1,2</sup> While the case fatality rate of sepsis is decreasing, the incidence is rising and will further rise as our population gets older. <sup>1,3,4</sup> To reduce the disease burden of sepsis and ensure proper disease risk stratification, it is important to identify targetable risk factors.

The diagnosis of AKI is currently based on an increase serum creatinine concentration and/or a decrease in urine output.<sup>5,6</sup> As in other forms of AKI, serum creatinine can be an insensitive indicator of kidney injury, and oliguria can be nonspecific in S-AKI. One of the most serious and life-threatening infections in dialysis patients is septicemia. The sepsis is complicated by the use of immunosuppressive drugs to treat and control underlying diseases and exacerbated

by nutritional deficiencies, the dialysis procedure and the disruption of cutaneous or mucosal barriers to infection.<sup>7</sup>

It accounts for over three fourths of deaths caused by infections. The annual percentage of mortality secondary to sepsis is approximately 100 to 300 fold higher in dialysis patients. Gram-negative bacteria were previously the most common cause of sepsis, in the last decade, gram-positive bacteria, most commonly staphylococci cause more than 50% of cases of sepsis. The annual percentage of mortality secondary to sepsis is approximately 100 to 300 fold higher in dialysis patients. Gram-negative bacteria were previously the most common cause of sepsis, in the last decade, gram-positive bacteria, most commonly staphylococci cause more than 50% of cases of sepsis. Uremia often results in immune deficiency. Malnourishment and older age may interact with uremia to impair the immune system. Risk may also vary according to the presence of comorbid conditions such as diabetes mellitus (DM) and disruptions of dermal barriers to gain access for dialysis. 14,15

The present study was conducted to study the presence of bacteremia, markers of sepsis and Inflammation in renal failure patients on hemodialysis, along with correlation of hematological abnormalities with sepsis in such patients.

#### 2. MATERIAL & METHODS

The present study was conducted for the period of one year and total of 50 patients of both sexes who were diagnosed as case of renal failure which include both acute kidney injury (AKI) and CKD on basis of clinical history, examination, biochemical markers and were advised for hemodialysis were included in the study.

The criteria used for AKI in the study was risk, injury, failure, loss of kidney function, and end-stage kidney disease (RIFLE) criteria. The kidney disease outcomes quality initiative (KDOQI) defines CKD as either kidney damage or a decreased glomerular filtration rate (GFR) of less than 60 ml/min/1.73 m2 for 3 or more months. The Criteria for the systemic inflammatory response syndrome, adapted from the American college of chest physicians/society of critical care medicine consensus conference. The conference of the systemic inflammatory response syndromes adapted from the American college of chest physicians/society of critical care medicine consensus conference.

#### Inclusion Criteria:

Patients of renal failure with newly inserted hemodialysis catheter subclavian venous catheter, internal jugular venous catheter or femoral catheter who developed systemic signs and symptom of sepsis e.g. fever, chills and rigor, tachycardia, tachypnea, hypotension, confusion, disorientation, and agitation after hemodialysis catheter insertion and hemodialysis and patients with local swelling, redness, pain or pus discharge at the site of hemodialysis catheter. Exclusion Criteria;

Those patients who had renal failure due to septicemia or post-operative renal failure, had history of hemodialysis in past, had known source of infection e.g. diabetic foot, pyelonephritis, bedsore, or had A-V fistula.

After recruiting patient for study, clinical history and relevant blood and radiological investigation (hemoglobin, total leucocyte count (TLC), differential leucocyte count (DLC), and platelet count), renal function test (RFT) (serum creatinine, blood urea, and serum electrolyte), serum phosphorus, C-reactive protein, liver function test (LFT) (serum bilurubin, serum total protein, serum albumin, alkaline phosphatase), thyroid function test – TFT (T3, T4, and thyroid stimulating hormone-TSH), urine routine and microscopy, urine culture and sensitivity. blood culture, central line catheter tip culture sensitivity, chest X-ray (CXR) P/A view, ultrasonography (USG) abdomen and kidney, ureter and bladder (KUB) were performed. Leukocyte count and blood culture were done prior to catheter insertion and a single sample

was collected from the peripheral vein before insertion of the catheter to rule out any existing bacteremia. If positive, the patient was excluded from the study. Secondly, after 72 hours of the insertion, two 5 ml samples of blood were collected, one from the peripheral vein and the other from the catheters; the latter being collected after at least 12 hours of hemodialysis.

In the laboratory, subcultures were done from Hartley's broth onto blood agar (BA) and MacConkey medium after overnight incubation at 37 0C and also on the 2nd, 4th and 7th days and were then discarded, if negative. Aseptically collected mid-stream urine sample in sterile bottle containing boric acid was transported to microbiology laboratory. Bacterial culture was performed by streaking 0.002 ml of mid-stream collected urine with a standard calibrated loop on MacConkey agar and 5% sheep blood agar plates which was incubated at 37 °C for 24 hours, under aerobic conditions and the colonies was counted by a colony counter. Sample that yielded pure bacterial growth of ≥105 cfu/ml was regarded as significant bacteriuria. Counts between 104 and 105 cfu/ml repeated while counts ≤104 cfu/ml considered as negative. Catheter tip was collected only from patients who had their catheters removed on completion of their HD sessions or in case they showed any signs of infection. It was cultured by Maki's standard semi quantitative method on blood agar and then put in trypticase soy broth (TSB).

#### STATISTICAL ANALYSIS

A colony count of  $\geq 15$  was considered significant for cultures done by Maki's method. <sup>19</sup> If the same organisms grew from both peripheral and central venous catherer (CVC) blood cultures confirmation was done by the pour-plate quantitative method. <sup>21</sup> Association and correlation assessment were done by statistical package for the social sciences (SPSS).

#### 3. RESULTS

Table 1: Patients on hemodialysis with sepsis and gender distribution

Parameter	Renal failure patients onhemodialysis with symptoms of sepsis		
	N=50	%	
Positive blood/cathetertip culture	10	20	
Negative blood/cathetertip culture	40	80	
Total	50		
Gender			
Male	32	64	
Female	18	36	

In our study among 50 patients of renal failure on hemodialysis the mean age in our study was 46.64±12.48 years with 32 male patients. Out of 50 patients 10 (20%) had positive blood and catheter tip culture and 40 (80%) of patients had negative blood and catheter tip culture.

Table 2: Distribution of patients according to age groups, symptoms and TLC

Age groups	N%	
15-25 years	2 (20)	
26-35 years	2 (20)	
36-45 years	1 (10)	
>45 years	5 (50)	
Symptoms		

Fever with chills and rigor	10 (100)	
Redness and Pain at hemodialysis catheter site	4 (40)	
Confused, Disoriented or comatose	3 (30)	
Hypotension	2 (20)	
TLC		
Less than 4.8/cumm (leucopenia),	0	
Between 4.8–10.8/cumm	2 (20)	
More than 10.8/cumm	8 (80)	

Out of 10 patients with sepsis 2 (20%) were in the age group between 15–25 years, 2 (20%) were in the age group between 26–35 years, 1 (10%) were in the age group 36–45 years and 5 (50%) were above 45 years of age. All 10 (100%) patients had episode of fever with chills and rigor, 4 (40%) patients had redness and pain at hemodialysis catheter site, 3 patients (30%) were confused, disoriented or comatose and 2 (20%) patients had hypotension. Among 10 patients of renal failure with sepsis, none had TLC less than 4.8/cumm (leucopenia), 2 (20%) patients had count between 4.8–10.8/cumm and 8 (80%) patients had TLC more than 10.8/cumm.

Table 3: Bacteria found on patients with sepsis

	Renal failure patients on hemodialysis with sepsis	
Type of bacteria	N=10	%
S. aureus	7	70
E. coli	1	10
Acinectobacter	1	10
Candida	1	10
Total	10	100

7 (70%) patients' blood culture was positive for S. aureus, and E. coli found in blood culture 1 (10%) patient, Acinectobacter in 1 (10%) patient and Candida in 1 (10%) patient.

Table 4: Most common catheter site associated with infection

	Renal failure patients onhemodialysis with	
Site of hemodialysis catheter	sepsis	
	N=10	%
Internal jugular venous catheter	2	20
Femoral catheter	7	70
Subclavian catheter	1	10
Total	10	100

Among 10 patients of renal failure on hemodialysis with sepsis 2 (20%) patients had internal jugular line for hemodialysis, 1 (10%) had subclavian line and 7 (70%) had femoral line for hemodialysis.

aiduinin levels		
Catheter duration	N%	
7-14 days	1 (10)	
14-21 days	1 (10)	
>21 days	8 (80)	
Serum phosphate levels		
Less than 3.5 mg/dl	0	
Between 3.5–5.5 mg/dl	2 (20)	
>5.5 mg/dl	8 (80)	
Serum albumin levels		
Less than 3.4 gm/dl	6 (60)	
More than 3.4 gm/dl	4 (40)	

Table 5: Distribution of patients according to catheter duration and serum phosphate and albumin levels

Catheter duration of 7-14 days was found in 1 (10%), 1 (10%) patients had central line between 14–21 days, and 8 (80%) patients had central line >21 days. None had serum phosphate level less than 3.5 mg/dl, 2 (20%) had serum phosphorus level between 3.5–5.5 mg/dl and 8 (80%) patients had serum phosphorus level >5.5 mg/dl. Albumin level less than 3.4 gm/dl was found in 6 (60) patients, 4 (40%) had serum albumin level more than 3.4 gm/dl.

### 4. DISCUSSION

Sepsis-associated acute kidney injury (S-AKI) is a common complication in hospitalized and critically ill patients, which increases the risk of developing chronic comorbidities and is associated with extremely high mortality. <sup>22-24</sup> As individual syndromes, sepsis and acute kidney injury (AKI) render the host susceptible to each other. Although sepsis is the most common contributing factor for developing AKI, AKI of any origin is associated with higher risk of developing sepsis. Sepsis has a complex and unique pathophysiology, which makes S-AKI a distinct syndrome from any other phenotype of AKI. <sup>25</sup>

In our study among 50 patients of renal failure on hemodialysis the mean age in our study was 46.64±12.48 years with 32 male patients. Out of 50 patients 10 (20%) had positive blood and catheter tip culture and 40 (80%) of patients had negative blood and catheter tip culture. Out of 10 patients with sepsis 2 (20%) were in the age group between 15–25 years, 2 (20%) were in the age group between 26–35 years, 1 (10%) were in the age group 36–45 years and 5 (50%) were above 45 years of age. All 10 (100%) patients had episode of fever with chills and rigor, 4 (40%) patients had redness and pain at hemodialysis catheter site, 3 patients (30%) were confused, disoriented or comatose and 2 (20%) patients had hypotension. We noted the incidence of sepsis was more in patients of age group greater than 45 years of age. Longitudinal cohort study conducted by Powe et al showed that sepsis was more common in older age group. <sup>26</sup> In 2013 a study conducted by Gupta in 45 patients of CKD showed that the prevalence of CRBSI was 17.78% in patients above 65 years of age.<sup>27</sup> So, our study conforms with other studies, who had shown that advanced age is risk factor for CRBSI. Robinson et al found that was fever was the most consistent symptom at onset of CRBSI (28 of 32 cases). 28 Kairaitis et al conducted a study of 105 haemodialysis catheters in 52 patients in order to identify patient outcomes and to analyse the effect of patient and catheter factors on the incidence of infectious complications, they found that exit-site infection was the cause for removal in 8% and most common clinical symptom was fever.<sup>29</sup>

Among 10 patients of renal failure with sepsis, none had TLC less than 4.8/cumm (leucopenia), 2 (20%) patients had count between 4.8–10.8/cumm and 8 (80%) patients had TLC more than 10.8/cumm. A study conducted by Gupta on 45 CKD patient on haemodialysis, catheter related infections were correlated with TLC.<sup>27</sup> 7 (70%) patients' blood culture was positive for S. aureus, and E. coli found in blood culture 1 (10%) patient, Acinectobacter in 1 (10%) patient and Candida in 1 (10%) patient. Nagarika et al in 2006-2007 conducted a study in 210 patients and found that bacteremia occurred in 17 (47.22%) patients with femoral catheter, 8 (22.22%) patients with subclavian catheter and 11 (30.55%) patients with jugular hemodialysis catheter.<sup>30</sup> Among 10 patients of renal failure on hemodialysis with sepsis 2 (20%) patients had internal jugular line for hemodialysis, 1 (10%) had subclavian line and 7 (70%) had femoral line for hemodialysis. Oliver et al had shown that incidence of bacteremia was 5.4% after three weeks of placement in internal jugular vein and 10.7% after one week in femoral vein.<sup>31</sup> Catheter duration of 7-14 days was found in 1 (10%), 1 (10%) patients had central line between 14–21 days, and 8 (80%) patients had central line >21 days. None had serum phosphate level less than 3.5 mg/dl. 2 (20%) had serum phosphorus level between 3.5–5.5 mg/dl and 8 (80%)

Catheter duration of 7-14 days was found in 1 (10%), 1 (10%) patients had central line between 14–21 days, and 8 (80%) patients had central line >21 days. None had serum phosphate level less than 3.5 mg/dl, 2 (20%) had serum phosphorus level between 3.5–5.5 mg/dl and 8 (80%) patients had serum phosphorus level >5.5 mg/dl. Albumin level less than 3.4 gm/dl was found in 6 (60) patients, 4 (40%) had serum albumin level more than 3.4 gm/dl. The study conducted by Plantinga had shown high phosphorus level was associated with infection in dialysis patients which supports our finding too.<sup>32</sup> We noted hypoalbuminemia is contributing to increased risk of catheter related infection matches with studies of Powe et al.<sup>26</sup> He suggested hypoalbuminemia was common in catheter related blood stream infection.

#### 5. CONCLUSION

Incidence of renal failure requiring hemodialysis has increased and accordingly use of vascular access to deliver haemodialysis therapy has increased. The patient requiring haemodialysis are prone to infections because of risk factors like advanced age, male sex, diabetes, anemia, hypoalbuminemia, hyperphosphatemia and prolonged duration of hemodialysis. The site of vascular access is an important risk factor for development of sepsis. GPC (S. aureus) is the commonest cause of sepsis.

## 6. REFERENCES

- 1. Rudd KE, Johnson SC, Agesa KM, Shackelford KA, Tsoi D, Kievlan DR, Colombara DV, Ikuta KS, Kissoon N, Finfer S, Fleischmann-Struzek C, Machado FR, Reinhart KK, Rowan K, Seymour CW, Watson RS, West TE, Marinho F, Hay SI, Lozano R, Lopez AD, Angus DC, Murray CJL, Naghavi M. Global, regional, and national sepsis incidence and mortality, 1990-2017: analysis for the Global Burden of Disease Study. Lancet. 2020 Jan 18;395(10219):200-211.
- 2. Singer M, Deutschman CS, Seymour CW, Shankar-Hari M, Annane D, Bauer M, Bellomo R, Bernard GR, Chiche JD, Coopersmith CM, Hotchkiss RS, Levy MM, Marshall JC, Martin GS, Opal SM, Rubenfeld GD, van der Poll T, Vincent JL, Angus DC. The Third International Consensus Definitions for Sepsis and Septic Shock (Sepsis-3). JAMA. 2016 Feb 23;315(8):801-10.
- 3. Jawad I, Lukšić I, Rafnsson SB. Assessing available information on the burden of sepsis: global estimates of incidence, prevalence and mortality. J Glob Health. 2012 Jun;2(1):010404.

- 4. Liyanarachi KV, Solligård E, Mohus RM, Åsvold BO, Rogne T, Damås JK. Incidence, recurring admissions and mortality of severe bacterial infections and sepsis over a 22-year period in the population-based HUNT study. PLoS One. 2022 Jul 12;17(7):e0271263.
- 5. Bellomo R, Ronco C, Kellum JA, Mehta RL, Palevsky P, ADQI workgroup. Acute renal failure—definition, outcome measures, animal models, fluid therapy and information technology needs: the Second International Consensus Conference of the Acute Dialysis Quality Initiative (ADQI) Group. Critical care. 2004 Aug;8:1-9.
- 6. Kidney Disease: Improving Global Outcomes (KDIGO) Acute Kidney Injury Work Group. KDIGO clinical practice guideline for acute kidney injury. Kidney Int Suppl. 2012; 2:1–138.
- 7. Kessler M, Hoen B, Mayeux D, Hestin D, Fontenaille C. Bacteremia in patients on chronic hemodialysis: a multicenter prospective survey. Nephron. 1993 Dec 12;64(1):95-100.
- 8. U.S. Renal Data System: USRDS 1997 Annual Data Report. National Institutes of Health, NIDDK, Bethesda, April, 1997.
- 9. Sarnak MJ, Jaber BL. Mortality caused by sepsis in patients with end-stage renal disease compared with the general population. Kidney international. 2000 Oct 1;58(4):1758-64.
- 10. Hirasawa H, Oda S, Nakamura M. Blood glucose control in patients with severe sepsis and septic shock. World journal of gastroenterology: WJG. 2009 Sep9;15(33):4132.
- 11. Sarnak MJ, Jaber BL. Mortality caused by sepsis in patients with end-stage renal disease compared with the general population. Kidney international. 2000 Oct 1;58(4):1758-64.
- 12. Hirasawa H, Oda SH, Nakamura M. Blood glucose control in patients with severe sepsis and septic shock. World J Gastroenterol. 2000;15(33):4132-6
- 13. Powe NR, Jaar B, Furth SL, Hermann J, Briggs W. Septicemia in dialysis patients: incidence, risk factors, and prognosis. Kidney international. 1999 Mar 1;55(3):1081-90.
- 14. Carton JA, Maradona JA, Nuno FJ, Fernandez-Alvarez R, Perez-Gonzalez F, Asensi V. Diabetes mellitus and bacteraemia: a comparative study between diabetic and non-diabetic patients. The European journal of medicine. 1992 Sep 1;1(5):281-7.
- 15. Khan IH, Catto GR. Long-term complications of dialysis: infection. Kidney International Supplement. 1993 Jun 2(41).
- 16. Biesen WV, Vanholder R, Lameire N. Defining Acute Renal Failure: RIFLE and Beyond. CJASN. 2006;1(6):1314-9.
- 17. National Kidney Foundation: K/DOQI clinical practice guidelines for chronic kidney disease: Definition, identification, and prediction of CKD progression Kidney International Supplements. 2013;3:63-72.
- 18. American College of Chest Physicians/Society of Critical Care Medicine Consensus Conference. Definitions for sepsis and organ failure and guidelines for the use of innovative therapies in sepsis. Crit Care Med. 1992;20:864-74.
- 19. Maki DG, Weise CE, Sarafin HW. A semiquantitative culture method for identifying intravenous catheter related infection. N Eng J Med. 1997;296(23):1305-9.
- 20. Chukwu BF, Okafor HU, Ikefuna AN. Asymptomatic bacteriuria in children with sickle cell anemia at The University of Nigeria teaching hospital, Enugu, South East, Nigeria. Italian Journal of pediatrics. 2011 Dec;37(1):1-5.
- 21. Quilici N, Audibert G, Conroy MC, Bollaert PE, Guillemin F, Welfringer P, Garric J, Weber M, Laxenaire MC. Differential quantitative blood cultures in the diagnosis of catheter-related sepsis in intensive care units. Clinical infectious diseases. 1997 Nov 1;25(5):1066-70.

- 22. Uchino S. Beginning and Ending Supportive Therapy for the Kidney (BEST Kidney) Investigators. Acute renal failure in critically ill patients: a multinational, multicenter study. JAMA. 2005;294:813-8.
- 23. Bagshaw SM, Uchino S, Bellomo R, Morimatsu H, Morgera S, Schetz M, Tan I, Bouman C, Macedo E, Gibney N, Tolwani A. Septic acute kidney injury in critically ill patients: clinical characteristics and outcomes. Clinical Journal of the American Society of Nephrology. 2007 May 1;2(3):431-9.
- 24. Bouchard J, Acharya A, Cerda J, Maccariello ER, Madarasu RC, Tolwani AJ, Liang X, Fu P, Liu ZH, Mehta RL. A prospective international multicenter study of AKI in the intensive care unit. Clinical Journal of the American Society of Nephrology. 2015 Aug 7;10(8):1324-31.
- 25. Mehta RL, Bouchard J, Soroko SB, Ikizler TA, Paganini EP, Chertow GM, Himmelfarb J, Program to Improve Care in Acute Renal Disease (PICARD) Study Group. Sepsis as a cause and consequence of acute kidney injury: Program to Improve Care in Acute Renal Disease. Intensive care medicine. 2011 Feb;37:241-8.
- 26. Powe NR, Jaar B, Furth SL, Hermann J, Briggs W. Septicemia in dialysis patients: incidence, risk factors, and prognosis. Kidney international. 1999 Mar 1;55(3):1081-90.
- 27. Punit G, Khunte P, Dubey P, Gupta GB. Catheter Related Infection In Geriatric Population On Hemodialysis, A Study From Central India. Rep Opinion. 2014;6(5):24-6.
- 28. Robinson JL, Casey LM, Huynh HQ, Spady DW. Prospective cohort study of the outcome of and risk factors for intravascular catheter-related bloodstream infections in children with intestinal failure. J Parenter Enteral Nutr. 2014;38(5):625-30.
- 29. Kairaitis LK, Gottlieb T. Outcome and complications of temporary haemodialysis catheters. Nephrology, dialysis, transplantation: official publication of the European Dialysis and Transplant Association-European Renal Association. 1999 Jul 1;14(7):1710-4
- 30. Nagarik AP, Soni S, Barnela S, Gondane S, Kishan AG. BACTEREMIA FOLLOWING TEMPORARY HEMODIALYSIS CATHETER INSERTION: A PROSPECTIVE STUDY. Indian Journal of Nephrology. 2007 Jul 1;17(3).
- 31. Oliver MJ, Callery SM, Thorpe KE, Schwab SJ, Churchill DN. Risk of bacteremia from temporary hemodialysis catheters by site of insertion and duration of use: a prospective study. Kidney international. 2000 Dec 1;58(6):2543-5.
- 32. PlevkovaJ. Systemic inflammatory response syndrome. JFMED. 2011;122-4.