Journal of Institute of Medicine Nepal Institute of Medicine, Kathmandu, Nepal



Original Article

JIOM Nepal. 2023 Apr;45(1):45-49.

Outcome of ABO-Incompatible Living Donor Kidney Transplantation: A Single Center Observational Study from Nepal

Saurav S Hamal¹, Rabin Nepali¹, Sujeet Poudyal², Dibya S Shah¹

Author(s) affiliation

¹Department of Nephrology and Transplant Medicine, Maharajgunj Medical Campus, Tribhuvan University Teaching Hospital, Institute of Medicine, Maharajgunj Kathmandu, Nepal

²Department of Urology and Kidney Transplant Surgery, Maharajgunj Medical Campus, Tribhuvan University Teaching Hospital, Institute of Medicine, Maharajgunj Kathmandu, Nepal

Corresponding author

Saurav S Hamal, MBBS, MD sauravkoid@gmail.com

Submitted

Jan 28, 2023

Accepted

Mar 30, 2023

ABSTRACT

Introduction

Kidney transplantation is the most effective treatment for end-stage renal disease (ESRD) patients. However, the increasing number of ESRD patients and the limited availability of living and cadaveric donors has led to a growing waiting list for kidney transplantation. ABO-incompatible transplantation has emerged as an alternative for these patients in Nepal.

Methods

This was an observational study of all the patients who underwent kidney transplantation from March 2017-Feb 2019. Data on demography of recipients and donor, blood group, human leukocyte antigen mismatch, induction agent, post-operative complications and creatinine clearance at discharge and one-year post-transplant were collected and analysed. Then we compared patient and kidney graft survival at one year between ABO incompatible and ABO compatible recipients.

Results

During the study period, there were total of 124 kidney transplant recipients among them 12 were ABO incompatible and 112 were ABO compatible recipients. The study showed slightly lower patient and graft survival in ABO-incompatible recipients than ABO compatible recipients (83.3% vs 99.2%, p<0.001). However, death-censored graft survival were similar in both groups (100%,p<0.001). The graft function at one year measured by creatinine clearance was better in ABO-incompatible recipients than ABO-compatible recipients (67.1±11.75 vs 61.87±12.82, p<0.001). Postoperative complications were slightly higher, however, complications at one year were lower in ABO-incompatible recipients.

Conclusion

The study showed that living donor ABO-incompatible kidney transplant recipients have a good outcome at one year but is associated with greater risks of patient and graft loss at early transplant period compared to ABO-compatible recipients.

Keywords

ABO incompatible; graft survival; infection; kidney transplantation; patient survival; rejection

© JIOM Nepal 45

INTRODUCTION

idney transplantation is the most effective treatment available for end-stage renal disease (ESRD).¹ Due to a growing number of ESRD patients, the waiting list of kidney transplantation is increasing. There is a strict living-related kidney donor program and limited availability of cadaveric donor program, hence ABO incompatible transplantation has emerged as an alternative to these patients in Nepal.²

ABO-incompatibility was once considered a contraindication to kidney transplantation.³ Until a breakthrough study formulated a desensitization protocol.⁴ This led to wider utilization of the procedure in Japan in the late 1980s, USA in the mid 1990s, and Europe in the early 2000s.⁴⁻⁶

ABO incompatible kidney transplantation was started in Nepal in 2017.² Few data are available about the long term survival of the recipients with ABO incompatible transplantation.⁷⁻⁹ The study aimed to evaluate the outcome of ABO incompatible transplantation at one year in terms of patient and graft survival. The subgroup analysis include perioperative complications and complications at one year, induction agent, mode of dialysis, native kidney disease and human leukocyte antigen (HLA) mismatch.

METHODS

It was a retrospective observational study which analysed all the living donor kidney transplantation at Department of Nephrology and Transplantation Medicine Tribhuvan University Teaching Hospital from March 2017 to Feb 2019. Prior approval was taken from the Institutional Review Committee (IRC) of Institute of Medicine. Data recorded included donor and recipient age and gender, blood group, HLA mismatch, induction agent, post-operative complications and creatinine clearance at discharge

and one year post transplant. Patients survival, refers to time from kidney transplant to patients death. Graft survival refers to time from transplant to kidney graft failure. Graft failure refers to need to resume the dialysis in the kidney transplant recipient, and was calculated as creatinine clearance measured by Cockcroft-Gault equation.

The data was recorded as per the working proforma, continuous data was presented as mean with standard deviation (SD) and compared using paired t-test, and the categorical data were compared using Chi-square test. Patient and graft survival was calculated by Kaplan-meier curve and compared between the groups by log-rank test. A p-value <0.05 was considered statistically significant. Analysis was performed using SPSS software ver 20.

RESULTS

A total of 124 patients were included in the study among them 12 were ABO incompatible (ABOi) recipient and 112 were ABO compatible (ABOc) recipients.

The ABOi donors had a mean age of 46.75±8.55SD, and nine of the 12 donors being female. Similarly the ABOi recipients had mean age of 32.5±12.2 SD and eight of them being males. ABOc donors had mean age of 46.9±10.84SD, 77 of them being females. Similarly in the ABOc recipients had mean age of 34.6±9.8 SD, 93 of them being males.

Among 12 ABOi recipients Antithymocyte globulin(ATG) was used as induction agent in seven patients, nine had HLA mismatch of less than three, and nine had blood group of A positive. Similarly in 112 ABOc recipients ATG was used in 108 patients, 80 patients had HLA mismatch of less than three as shown in Table 1.

Among 12 ABOi recipients, 10 patients were alive during the first year and the two patients died.

Table 1. Baseline profile of the recipients and donor

Parameters	ABO inco	ompatible	ABO compatible		
	Recipients	Donor	Recipients	Donor	
Age (Mean±SD)	32.5±12.2	46.75±8.55	34.6±9.86	46.9±10.84	
Sex	8				
Male	(66.6%)	3 (25%)	93 (83.03%)	35 (31.25%)	
Female	4 (33.3%)	9 (75%)	19 (16.9%)	77 (68.75%)	
HLA mismatch					
0-3	9 (75%)		80 (71.42%)		
4-6	3 (25%)		32 (28.58%)		
Induction agent					
ATG	7 (58.33%)		108 (96.4%)		
Basiliximab	5 (41.6%)		4 (3.56%)		

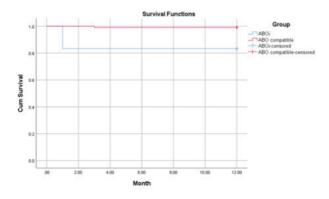


Figure 1. Patients survival at one year

Among the patients who did not survived during the first year the cause of death was ABMR followed by sepsis in one patient and shock in the other, and both of them died with the functioning graft in the first month of transplantation. Among 112 ABOc recipients 111 patients survived at one year, one patient died due to surgical complication. The patients survival at one year in ABOi and ABOc recipients was 83.3% and 99.2%.(P<0.001) as shown in Figure 1.

In the ABOi, graft survival and death censored graft survival was 83.3% and 100%, and in ABOc recipients the graft survival and death censored

Table 2. Creatinine clearance at discharge and one year

Creatinine clearance	ABOi recipients (n=12)	ABOc recipients (n=112)	p value	
Discharge	62.1±11.75	58.47±15.34	< 0.001	
One year	67.1±14.16	61.87±12.82	<0.001	

Table 3. Post-operative complications

	Number (%)			
Complications	ABOi recipients (n=12)	ABOc recipients (n=112)		
UTI	4 (33.3%)	10 (8.9%)		
Pneumonia	1 (8.3%)	4 (3.57%)		
Delay Graft Function	0	6 (5.3%)		
ACR	0	1 (0.89%)		
ABMR	1 (8.3%)	1 (0.89%)		
Hematoma	0	4 (3.57%)		
Dissection	1 (8.3%)	0		
Re-anastomosis	1 (8.3%)	2 (1.78%)		
Renal artery stenosis	1 (8.3%)	3 (2.67%)		

Graft Survival

Group

ABO compatible

ABO compatible

ABO compatible censored

ABO compatible censored

ABO compatible censored

Month

Figure 2. Graft survival, death censored graft survival at one year

graft survival was 99.2% and 100%. Overall graft survival was low in ABOi recipient however death censored graft survival was similar in both the groups(p<0.001) as shown in Figure 2.

The creatinine clearance of ABOi recipients at discharge was 62.1 ± 11.75 and 67.1 ± 14.16 at one year. Similarly in the ABOc recipients the creatinine clearance at discharge was 58.47 ± 15.34 and at one year it was 61.87 ± 12.82 . The graft function was better in ABOi recipients at one year(p<0.001) as shown in Table 2.

In the ABOi recipients medical complication was seen in six patients with UTI being the common complication seen in four patients, antibody mediated rejection (ABMR) seen in one patients and three patients had surgical complications. Similarly in ABOc recipients medical complications were seen in 22 patients with UTI seen in 10 patients, acute cellular rejection(ACR) in one, antibody mediated rejection in one patient. Similarly nine patients had surgical complications as shown in Table 3.

Among the ABOi recipients, 3(30%) patients were hospitalized for pneumonia and cryptococcal

Table 4. Complications at one year

	Number (%)			
Complications	ABOi recipients (n=12)	ABOc recipients (n=112)		
Pneumonia	2 (16.66%)	4 (3.57%)		
PTDM	0	9 (8.03%)		
ABMR	0	4 (3.57%)		
ACR	0	8 (7.14%)		
FSGS recurrence	0	4 (3.57%)		
Sepsis	0	2 (1.78%)		
BK virus nephropathy	0	2 (1.78%)		
Cryptococcal meningitis	1 (8.33%)	0		
Pulmonary TB	0	2 (1.78%)		

meningitis, similarly 35 (31.25%) ABOc recipients had complications, among them Post transplant diabetes mellitus (PTDM) was seen in nine patients, ACR was seen in eight patients, ABMR, FSGS recurrence and pneumonia were seen in four patients. Sepsis, BK virus nephropathy and pulmonary tuberculosis were seen in two patients as shown in Table 4.

The most common cause of CKD among ABOi recipient as well as in ABOc recipients was chronic glomerulonephritis(CGN). There were seven preemptive transplantation in ABOc recipients and two patients in ABOc recipient were on peritoneal dialysis as shown in Table 5.

DISCUSSION

With the increasing number of patients with ESRD, obtaining a living-related compatible kidney transplantation has become more challenging, leading to ABO incompatible kidney transplant as a safe and alternative way to shorten the waiting time for a renal graft.

In this study a total of 12 ABO incompatible kidney transplant recipient along with 112 ABO compatible kidney transplant recipient were included. The initial studies had shown a graft survival of 79% and patient survival rate of 88% at 1 year, subsequent studies have shown improved patients and graft survival, due to improved immunosuppression protocols.

Study done in Japan found comparable patient and graft survival rates between ABOi and ABOc recipients, with most rejections occurring in ABOi recipients within the first month after transplantation, suggesting that successful graft function in ABOi recipient requires keeping antibody level at safe level in the initial post-operative period.¹¹

A meta-analysis found 96% and 98% patient survival rate of ABOi and ABOc recipients at one year post transplant, with infection being the most common cause of death. 9,12 Complications like infection, antibody mediated rejections were more common in ABO incompatible recipients during the period. 12

Studies have shown a graft survival of over 94% and patient survival of more than 95% in ABO incompatible recipients. 10-12 However domestic reports on ABO incompatible kidney transplantation are few. Our study of 12 ABO incompatible recipients performed since March 2017 showed a patient and graft survival rate of 83.3% at one year, and death censored graft survival was 100% in both the ABOi and ABOc recipients. Two ABO incompatible patients who died, one of them developed ABMR and died due to sepsis and the other died due to shock with the functioning graft. Post-operative complications were higher in ABO incompatible recipients, however complications at one year was lower than ABO compatible recipients.

CONCLUSION

The result from our studies shows that living donor ABO incompatible kidney transplant recipients have good outcome at one year but is associated with greater risk of patient and graft loss at early transplant period compared to ABO compatible recipients.

Table 5. Native kidney disease and mode of dialysis in patients

Native kidney disease	ABOi recipients (n=12)		ABOc recipients (n=112)				
	HD	PD	Total	HD	PD	Preemptive	Total
CGN	9 (75%)	0	9	70 (62.5%)	1 (0.9%)	5 (4.5%)	76
IgAN	0	0	0	6 (5.4%)	0	0	6
Diabetes	1 (8.3%)	0	1	8 (7.1%)	1 (0.9%)	2 (1.8%)	11
SLE	0	0	0	3 (2.7%)	0	0	3
MPGN	0	0	0	3 (2.7%)	0	0	3
ADPKD	0	0	0	3 (2.7%)	0	0	3
Obstructive Uropathy	1 (8.3%)	0	1	3 (2.7%)	0	0	3
VUR	0	0	0	3 (2.7%)	0	0	3
FSGS	0	0	0	2 (1.8%)	0	0	2
Alport syn-drome	0	0	0	2 (1.8%)	0	0	2
Crescentic glomerulonephritis	1 (8.3%)	0	1	0	0	0	0
Total	12	0	12	103	2	7	112

Note:cgn-chronic glomerulonephritis, IgAN- IgA nephropathy, SLE-systemic lupus erythematosus, mpgn-membranoproliferative glomerulonephritis, ADPKD-autosomal dominant polycystic kidney disease, vurvesicoureteral reflux, fsgs-focal segmental glomerulosclerosis.

ACKNOWLEDGEMENT

We would like to thank the faculty and residents who helped us with the study.

FINANCIAL SUPPORT

The author(s) did not receive any financial support for the research and/or publication of this article.

CONFLICT OF INTEREST

The author(s) declare that they do not have any conflicts of interest with respect to the research, authorship, and/or publication of this article.

REFERENCES

- Colin RL, Jane CT. Clinical management of the adult kidney transplant recipient. In YU ASL, Chertow GM, Luyckx VA Editors. Brenner and Rector's The Kidney. 11th ed. 2020. p.2244-87.
- Rabin N, Shah DS. Experience of starting ABO incompatible renal transplant in Nepal. Transplant report 2019; 4: 100026.
- Tonelli M, Wiebe N, Knoll G et al. Systematic review: Kidney transplantation compared withdialysis in clinically relevant outcomes. Am J Transpl 2011; 11: 2093–109.
- 4. Alexandre GP, Squifflet JP, De Bruyère M et al. Present experiences

- in a series of 26 ABO incompatible living donor renal allografts. Transplant Proc 1987; 19: 538–42.
- Tanabe K. Japanese experience of ABO-incompatible living kidney transplantation. Transplantation 2007; 84: 4–7
- Montgomery RA, Locke JE, King KE et al. ABO incompatible renal transplantation:a paradigm ready for broad implementation. Transplant 2009; 87: 1246–55.
- Tyden G, Kumlien G, Fehrman I. Successful ABO-incompatible kidneytransplantations without splenectomy using antigen-specific immunoadsorption and rituximab. Transplantation 2003; 76: 730–31
- Zschiedrich S, Janigen B, Dimova D et al. One hundred ABO incompatible kidney transplantations between 2004 and 2014: A single-centre experience. Nephrol Dial Transplant 2016: 31; 663–671.
- Jha PK, Bansal SB, Sethi SK et al: ABO-incompatible renal transplantation in developing world -crossing the immunological (and mental) barrier. Ind J Nephrol 2016; 26: 113–18
- Hart A, Lentine KL, Smith JM, Miller JM, Skeans MA, Prentice M, et al.OPTN/SRTR 2019 annual data report: kidney. Am J Transpl 2021; 21:21–137..
- Takahasi K, Saito K, Takahara S et al. Excellent Long-term outcome of ABO-incompatible living donor kidney transplantation in Japan. Am J Transplant 2004; 4: 1089-96.
- Weerd DAE, Betjes MGH. ABO-incompatible kidney transplant outcomes: A Meta-analyses. Clin J Am Soc Nephrol 2018; 13:1234-43.