



Comparison of Anti-Human T-Lymphocyte Immunoglobulin (r-ATLG - Grafalon) vs Anti-Thymocyte Globulin (r-ATG - Thymoglobulin) in Kidney transplant: A 5-year follow-up study

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Introduction

Induction agent in renal transplant is an essential element. In Living Donor Kidney Transplantation its practice is varied in India from no induction to different agents.

T cell depletion agents, Thymoglobulin (ATG) and Grafalon (ATLG), are commonly employed as induction agents in kidney transplantation

This study aims to assess the short- and long-term outcomes of kidney transplants at our centre focusing on the use of these agents.



<u>AIM</u>

 To look into graft outcome of patients who received Thymoglobulin (ATG) and Grafalon (ATLG) as an induction agent for Living Donor Kidney Transplant.



Objective

- To compare the graft function between the two T Cell depleting agents.
- To compare the Reoccurrence of Denovo Diseases between two groups.

METHODOLOGY

This retrospective study analysed data from Ktx patients at a tertiary care hospital in INDIA, between 2017 and 2024. Of the 581 patients included, 375 received Thymoglobulin and 206 received Grafalon as induction therapy.

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All the patients received standard triple immunosuppressant: Tac, EC-MPS and Prednisolone for maintenance.



Follow-up included monitoring for graft rejections, infections, denovo diseases, and overall graft survival.

Results - Demographic

| Parameters | Thymoglobulin (n=375) | Grafalon (n=206) | |
|---------------------------------|-----------------------|------------------|--|
| Age (years) | 41.4 ± 13 | 39.13 ± 12.56 | |
| Age range (years) | 10 to 68 | 10 to 68 | |
| Gender | | | |
| Male | 279(74.4%) | 180 (87.3%) | |
| Female | 96(25.6%) | 26 (13.4%) | |
| Body weight (Kg) | 61.83 ± 9.4 | 57.89 ± 10 | |
| | Native kidney disease | | |
| Hypertensive Nephrosclerosis | 75(20%) | 60 (29.1%) | |
| Diabetic nephropathy | 56(14.9%) | 35 (16.9%) | |
| CGN | 234(62.4%) | 98(47.5%) | |
| CIN | 14(3.7%) | 10(2.6%) | |
| Anti GBM | 1 | 2 | |
| CKDu | 0 | 1 | |

Results - Demographic

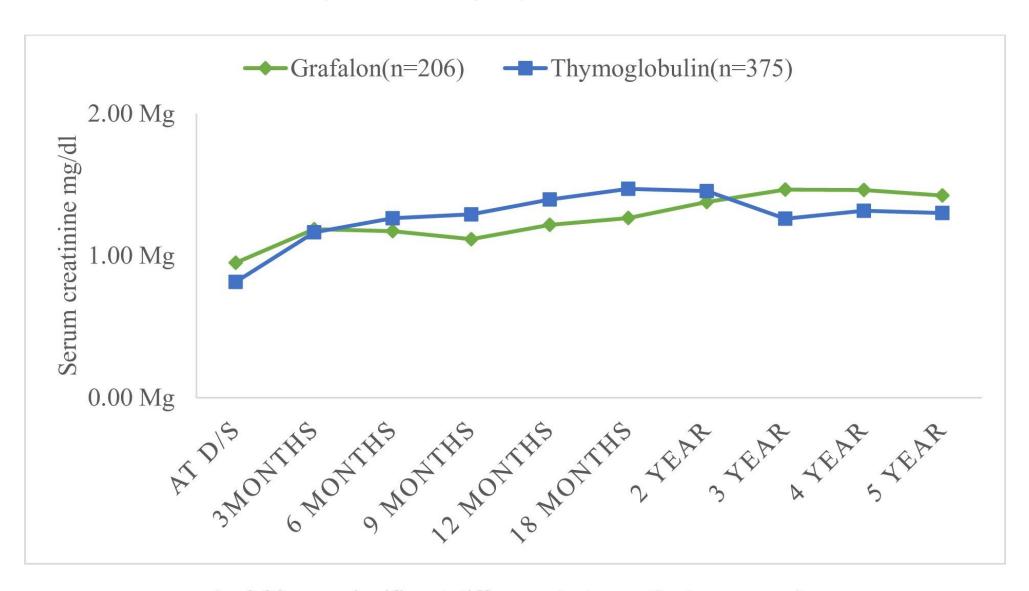
| Parameters | Thymoglobulin (n=375) | Grafalon (n=206) |
|------------------------------|-----------------------|------------------|
| ABOi Transplant | 40(10.6%) | 11(5.3%) |
| HIV Positive | 1 | 0 |
| HBsAg Positive | 3 | 12 |
| HCV Positive | 10 | 14 |
| Donor details | | |
| Donor Age (years) | 39.78±13.6 | 41±13.7 |
| Donor gender | | |
| Male | 135(36%) | 79 (38.3%) |
| Female | 240(64%) | 127(61.6%) |
| Mean Cumulative Dose (mg/kg) | 146.63 ± 35.5 | 305.1 ± 61.8 |
| Mean±SD (mg/kg) | 2.37±1.2 | 5.4±1.4 |

Results: Rejection Analysis

| Parameters | Thymoglobulin (n=375) | Grafalon (n=206) |
|-------------------------------------|-----------------------|------------------|
| Rejections | 37(9.8%) | 24(11.6%) |
| Acute cell mediated rejection (ACR) | 26 | 20 |
| Within 1 month | 6 | 6 |
| • 1 to 6 months | 11 | 8 |
| • >6 months | 5 | 5 |
| Acute on Chronic Rejection | 4 | 1 |
| Chronic ACR | 0 | 1 |
| Antibody mediated rejection (ABMR) | 1 | 1 |
| Hyper acute Rejection (1 Day) | 1 | - |
| Chronic ABMR | 9 | 2 |
| • >1 year | 3 7 | 2 |
| • <1 Year | | - |
| Acute tubular injury | 25(6.7%) | 15(7.2%) |
| Diffuse | 16 | 12 |
| Patchy | 9 | 3 |
| De-novo disease recurrence | 12(3.2%) | 3(1.4%) |
| IgA recurrence | 6 | 1 |
| FSGS recurrence | 6 | 2 |

(On rejection parameter Non-significant difference between two groups (p Value =0.54)

Result: Graft Function



(p=0.90, non-significant difference between the two groups)

Conclusion:

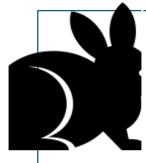


Both the short-term and long-term outcomes of kidney transplant recipients receiving Thymoglobulin or Grafalon as induction therapy were similar in terms of graft rejection incidence.





However, Thymoglobulin was associated with a higher occurrence of de novo diseases (this may be an incidental finding or due to some unexplained reasons).



Mechanism action of ATG and ATLG are different. ATG acts as a broader immunosuppression and T Cell depletion compared to ATLG which preferentially acts on the activated T cell.



Good compliance with Oral immunosuppressant can improve graft survival as per the current study.

