Role of Ruxolitinib in Steroid-Refractory Graft versus Host Disease in Patients Undergoing Allogeneic Stem Cell Transplant

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ABSTRACT

The objective of this study was to evaluate the role of Ruxolitinib in steroid-refractory graft versus host disease. This retrospective descriptive study was conducted from January 2018 to December 2021. A total of 157 patients underwent allogeneic stem cell transplants during the study period. Of these, 20 patients having steroid-refractory GVHD treated with Ruxolitinib were selected for the study. The primary endpoint was the overall response rate to Ruxolitinib measured at 4 weeks and 24 weeks for acute and chronic GVHD, respectively. The secondary endpoints were overall survival and failure-free survival.

Of these 20 patients, 7 (35%) had acute GVHD, and 13 (65%) had chronic GVHD. Of acute GVHD, 2 (10%) had grade II, 4 (20%) had grade III, and 1 (5%) had grade IV acute GVHD. Of 13 patients with chronic GVHD, 7 (35%) had moderate and 6 (30%) had severe chronic GVHD.

In steroid-refractory acute GVHD, the overall response rate to Ruxolitinib was 85.7%, and in chronic GVHD, it was 84.6%. The failure-free survival was 80% and overall survival was 85%. Adverse events of any grade occurred in 16 (80%) patients with grade III/IV adverse events in 4 (20%) patients only. The study showed that Ruxolitinib is a safe and effective second-line therapy for acute and chronic steroid-refractory GVHD.

Key Words: Ruxolitinib, GVHD, Allogeneic stem cell transplant.

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2021. Acute GVHD was defined and classified as per the Mount Sinai Acute GVHD International Consortium (MAGIC) criteria. Chronic GVHD was defined and graded as per the National Institute of Health (NIH) 2014 criteria. Steroid refractoriness was defined as per the NIH-EBMT-CIBMTR Task Force criteria for acute and chronic GVHD.1 Inclusion criteria were patients with steroid-refractory acute or chronic GVHD receiving Ruxolitinib for at least 4 weeks in acute and 12 weeks in chronic GVHD, respectively. Exclusion criteria was intolerance to Ruxolitinib causing discontinuation of therapy.

Table 1: Patient characteristics. Overall survival was calculated using Kaplan-Meier estimate with log-rank test, and categorical variables were analysed using chi-sq. test. MSD: Matched sibling donor, BMH: Bone marrow harvest, PBSC: Peripheral blood stem cells, CSA: Cyclosporine, MTX: Methotrexate, MMF: Mycophenolate mofetil, GVHD: Graft versus host disease.

The primary endpoint was overall response rate (ORR) defined as complete or partial response and was measured at 28 days of the Ruxolitinib treatment for acute GVHD and at 24 weeks for chronic GVHD. The secondary endpoints included overall survival (OS defined as time from transplant till death) and failure-free survival (FFS defined as occurrence of death, relapse of GVHD or need for the new systemic treatment for GVHD, whichever occurred first).3 Complete response (CR) was defined as resolution of all the manifestations of GVHD and partial response was defined as improvement in GVHD stage in all initial GVHD target organs without complete resolution or worsening. Data were analysed by the Statistical Package for Social Sciences (SPSS) version 20 using chi-square test for analysing the impact of categorical variables on ORR, and survival was calculated using Kaplan-Meier estimate with log-rank test, categorical variables were expressed as counts and percentages.

A total of 153 patients underwent allogeneic HSCT during the study period, out of which 20 were included in the study as per inclusion criteria. Patient characteristics, primary diagnosis, and study outcomes are shown in Table 1.

The overall response rate in acute GVHD measured at 28 days was 85.7%, and in chronic GVHD was 84.5% measured at 24 weeks. In acute GVHD, 6 out of the 7 patients responded on 28th day. Two patients had a complete response (CR) and 4 patients had a partial response (PR). In chronic GVHD, 11 out of 13 patients responded, 2 (15.3%) patients had a complete response and 9 (69.2%) had a partial response (Figure 1). In both acute and chronic GVHD, the failure-free survival was 80% and overall survival was 85%. The overall survival for acute GVHD was 62.5%, and for chronic GVHD was 100% (p = 0.004).

The median duration of therapy in the acute group was 60 days, and the median duration of therapy in the chronic group was 180 days. Adverse events of any grade occurred in 16 (80%) patients with grade III/IV adverse events in 4 (20%) patients only. The most common adverse effects were thrombocytopenia (30%), anaemia (20%), CMV reactivation (20%), fungal infection (10%), and tuberculosis (5%). However, there was no treatment discontinuation due to the adverse events related to the Ruxolitinib therapy.

This was a retrospective analysis on the use of Ruxolitinib in steroid-refractory acute and chronic GVHD. Ruxolitinib was the first JAK1/2 inhibitor given FDA approval in 2011 for primary myelofibrosis based on COMFORT-I and COMFORT-II trials and later for polycythemia vera in 2014 for patients intolerant or resistant to hydroxyurea. The role of Ruxolitinib was further explored in GVHD considering its myriad effects in signal transduction leading to cell proliferation and inflammation.
The youngest age in this study was 2 years, and there were no safety concerns; as shown by Khandelwal et al. in their study of Ruxolitinib in pediatric steroid-refractory acute GVHD.\(^6\)

Several retrospective studies have shown variable response rates for Ruxolitinib in acute GVHD.\(^6\) The overall response rate in these retrospective studies ranged from 45% to 82% for acute GVHD. In this study, the authors found an overall response rate of 85.7% at 28 days, which is comparable to the available studies. Similarly for chronic GVHD, the overall response rate ranged from 85 to 100%.

Ruxolitinib was awarded FDA approval for acute GVHD after REACH2 trial which was a phase 3 randomised multi-centre open-label trial. The trial included 309 patients randomised to receive either Ruxolitinib or the best available therapy for at least 28 days, and the overall response rate was found to be much higher in the former group as compared to the later (62.3% versus 39.4%, \(p < 0.001\)). The ORR for acute SR-GVHD was 85.7% which included CR of 28.5% and PR of 57.1%. This is much superior to that seen in REACH2 trial yet because of the smaller sample size, no definitive conclusion can be drawn. The ORR for chronic SR-GVHD was 84.6% including CR of 15.3% and PR of 69.2%. This is much superior to REACH3 trial which was a multi-centre, RCT showing an ORR of 50% at 24 weeks.

The record regarding adverse events showed that Ruxolitinib has an acceptable safety profile. Although this study has shown much superior ORR compared to the REACH studies, the sample size was small, and follow-up was shorter. Further evaluation in a randomised trial, with the larger number of patients is needed to confirm these findings.

The data shows that Ruxolitinib is an effective and safe oral therapy for acute and chronic GVHD. It can be used as a second-line therapy in steroid-refractory disease as it improves overall response and avoids complications of long-term steroid use.

**PATIENTS’ CONSENT:**
This is a retrospective study; therefore, patients’ consent is not required.

**COMPETING INTEREST:**
The authors declared no competing interest.

**AUTHORS CONTRIBUTION:**
HJ: Drafting the study, data analysis, and interpretation.
QUNC: Conception of work, critical revision, and final approval.
NS, MAK: Design and methodology of the study.
RI: Revising the study, data analysis, and interpretation.
MY: Data acquisition and analysis.
All the authors have approved the final version of the manuscript to be published.

**REFERENCES**