Original Article

Efficacy of a Reduced-dose Rasburicase: Single-institution Experience in India

Abstract

Background: Tumor lysis syndrome (TLS) is an oncological emergency associated with life-threatening metabolic abnormalities. Hyperuricemia is a feature of TLS and is treated with hydration, urine alkalinization, and allopurinol. Rasburicase lowers uric acid (UA) rapidly at the labeled dose of 0.15–0.2 mg/kg/day for 5 days. In a developing country like India where affordability is one major limitation to medical care, the use of rasburicase at the dose recommended by the US Food and Drug Administration (FDA) is not always possible. There is no convincing data suggesting the efficacy of a lower dose of rasburicase (1.5 mg or 3 mg) in the treatment of TLS. We conducted a retrospective study from January 2015 to June 2016 to assess the efficacy of a reduced-dose rasburicase in patients with TLS. Materials and Methods: All the patients with TLS were given rasburicase (single dose of 1.5 mg) on day 1 of chemotherapy. Serum UA, potassium, creatinine, and calcium levels were monitored every 24 h. All the patients who did not achieve normalization of UA with one dose of rasburicase were given another 1.5 mg of rasburicase. **Results:** Out of 90 patients, 54 patients (60%) had normalization of UA levels after 1.5 mg of rasburicase and 16 (18%) patients required 3 mg of rasburicase for bringing down the UA level to normal. The low serum UA levels were maintained even on the 3rd day of rasburicase. Rasburicase was well tolerated, and there was no death due to TLS. Thirty-one patients (64%) had normalization in the serum creatinine levels after rasburicase. Conclusion: We conclude that a low dose of rasburicase (1.5 mg or 3 mg) is cost effective in reducing serum UA (especially for low-risk and intermediate-risk TLS) and the higher dose as recommended by the US FDA is required only for patients with high-risk TLS.

Keywords: Low dose, rasburicase, tumor lysis syndrome

like India where affordability is one of the major limitations to medical care, the use of rasburicase at the dose recommended by the US Food and Drug Administration (FDA) is not always possible. There is no convincing data from India suggesting the efficacy of a lower dose of rasburicase (1.5 mg or 3 mg)

Introduction

Tumor lysis syndrome (TLS) is an oncological emergency associated with life-threatening potentially metabolic abnormalities.[1] Hyperuricemia is a feature of TLS and is treated with hydration, urine alkalinization, and allopurinol. Allopurinol inhibits the conversion of hypoxanthine to xanthine and xanthine to uric acid (UA) by inhibiting xanthine oxidase. It has no direct effect on existing UA. Rasburicase being a recombinant urate oxidase is highly efficacious in TLS. Rasburicase lowers UA rapidly to very low levels at the labeled dose of 0.15-0.2 mg/kg daily for 5 days by converting UA to allantoin which is rapidly excreted. Despite this dramatic effect on UA, rasburicase has not been shown to have any beneficial impact on survival. There are various studies suggesting the effectiveness of a reduced dose of rasburicase (3 mg to 6 mg single dose). In a developing country

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Objectives

The objective is to study the efficacy of a reduced dose of rasburicase (1.5 mg or 3 mg) in adult patients with TLS.

Materials and Methods

in the treatment of TLS.

A retrospective review from January 2015 to June 2016 was conducted in adult oncology patients who received rasburicase. We evaluated the efficacy of a reduced dose of rasburicase (1.5 mg or 3 mg) in patients aged 18–72 years presenting with clinical or laboratory TLS^[1] [Table 1] to our institution. These patients were

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administered rasburicase, hydration (3 L/m²/day), and chemotherapy on day 1. Patient's biochemistry parameters such as serum UA, serum potassium, serum creatinine, and serum calcium were studied before and after giving rasburicase. All the patients with TLS [Bishop and Cairo definition of clinical and/or laboratory TLS is shown in Table 1] received 1.5 mg of rasburicase on the 1st day, and the response was studied in terms of decrease in UA levels or decrease in serum creatinine levels. Those patients who did not achieve normal UA level within 24 h of 1.5 mg of rasburicase were given one more dose of rasburicase (1.5 mg). All patients were also evaluated for the change in serum creatinine, serum calcium, and serum potassium levels, postrasburicase administration.

Results

The median UA level was 9.9 mg/dl (8.2–13.4 mg/dl). A total of 90 patients received low-dose rasburicase. Out of 90 patients, 54 patients (60%) had normalization of UA levels after 1.5 mg of rasburicase and 16 (18%) patients required 3 mg of rasburicase for bringing down the UA level to normal. The low serum UA levels were maintained even on the 3rd day of rasburicase. Twenty patients (22%) did not achieve normal UA levels even with 3 mg of rasburicase although they had more than 50% reduction in UA levels. All the patients who did not

Laboratory tumor lysis	Clinical tumor lysis
2 or more of the following criteria	Laboratory tumor lysis plus
within 3 days prior to or 7 days	1 or more of the following

Seizure

range

sudden death

Cardiac dysthymias or

Creatinine >1.5 times of

age adjusted reference

Table 1: Bishop and Cairo criteria for TLS

after initiation of chemotherapy
Uric acid: ≥ 8 mg/dl or 25%
increase from baseline
Potassium: ≥6 mEq/l or 25%

increase Phosphorus: ≥4.5 mg/dl or 25% increase from baseline

Calcium: ≤7 mg/dl or 25% decrease from baseline

acheive normal UA levels after 3 mg rasburicase had high risk^[2] [Table 2] of TLS. Rasburicase was well tolerated, and there was no death due to TLS among the patients studied. Out of 90 patients, 48 patients (53%) had elevated creatinine due to TLS. The median serum creatinine level was 3.8 mg/dl (1.9–5.4 mg/dl). Thirty-one patients (64%) had normalization in the serum creatinine levels after rasburicase. Two patients of Burkitt's lymphoma required hemodialysis due to acute renal failure. The cost of one dose (1.5 mg) of rasburicase was Rs. 6000 as compared to the usual FDA recommended dose (0.15 mg/kg/day) which comes out to be Rs. 36,000 per day.

Discussion

TLS and hyperuricemia are serious complications with significant morbidity and potential mortality in patients with hematologic malignancies undergoing anticancer therapy. Allopurinol has been used for many years in the prevention and management of TLS-related hyperuricemia. However, allopurinol should be administered for ≥ 3 days for the achievement of significant reduction in UA levels. Rasburicase offers a potential advantage over allopurinol by its rapid onset of action, reducing preexisting pool of UA within few hours.[3] The results of our study demonstrate that a fixed low-dose rasburicase is a highly effective agent for the management of hyperuricemia associated with TLS. We also find that the cost of rasburicase reduces by one-sixth when using a fixed low dose for TLS. All patients at potential risk and majority of high-risk patients responded to a reduced dose, indicating that in appropriately monitored patients, single dose followed by dosing as needed can be cost saving. Our results are similar to the results of Hummel et al.[4] wherein fifty patients were studied to evaluate the efficacy of low-dose rasburicase in TLS [Table 3].[4-11]

Conclusion

As per our knowledge, this is the largest study conducted for evaluating the efficacy of low-dose rasburicase. A reduced dose of rasburicase at 1.5 mg single dose (repeated only

Table 2:	Risk str	atification	of TLS	patients
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Table 2. Misk stratification of TLB patients					
Low risk	Intermediate risk	High risk			
Multiple myeloma	Neuroblastoma, GCT, SCLC	AML with TLC >100,000/mm ³			
CML	CLL with TLC >50,000/mm ³	ALL with TLC $>$ 100,000/mm ³ or LDH $>$ 2 \times ULN			
CLL with TLC <50,000/mm ³	AML with either TLC >25,000-1,00,000/ mm3 or with LDH >2 × ULN	Stage III/IV Burkitt's lymphoma, any stage Burkitt's lymphoma with LDH >2 × ULN			
Hodgkin's lymphoma	Intermediate-grade NHL with LDH >2 × ULN	Stage III/IV lymphoblastic lymphoma or any stage lymphoblastic lymphoma with LDH >2 × ULN			
AML with TLC <25,000/mm³ and LDH <2*ULN	ALL with TLC <100,000/mm ³ and LDH <2 × ULN	Intermediate-risk disease with renal dysfunction			
Adult ALCL	Burkitt's lymphoma with LDH <2 × ULN	Intermediate-risk disease with elevated serum uric acid or potassium levels			

LDH – Lactate dehydrogenase; ULN – Upper limit of normal; NHL – Non-Hodgkin's lymphoma; CML – Chronic myeloid leukemia; CLL – Chronic lymphocytic leukemia; AML – Acute myeloid leukemia; ALCL – Anaplastic large cell lymphoma; GCT – Germ cell tumor; SCLC – Small cell lung cancer; TLC – Total leukocyte count, ALL – Acute lymphoblastic leukemia

Table 3: Other studies on efficacy of low dose rasburicase					
Study	n	Malignancy	Dose of rasburicase	Number of doses	
Lee et al.[5]	3	ALL	0.08-0.26 mg/kg	1	
McDonnell et al.[6]	11	3 NHL B-cell, 1 Burkitt's lymphoma, 3	0.0232-0.1361 mg/kg	1	
		AML, 1 CMML, 1 MDS			
Liu et al. ^[7]	8	3 AML, 2 ALL, 2 NHL, 1 CML	0.141-0.178 mg/kg	1	
Trifilio et al.[8]	43	20 plasma cell dyscrasias, 10 NHL, 7	3 mg	1, except for 6 additional	
		AML, 3 CLL, 1 MDS		doses (2 doses, 1.5 mg; 4 doses, 3 mg)	
Hutcherson et al.[9]	11		0.045-0.1 mg/kg	1, except for 1 additional dose (0.1 mg/kg)	
Hummel et al.[4]	50	14 NHL, 9 AML, 7 CLL, 6 CMP/MDS	0.031-0.11 mg/kg	Given as 1	
		5 ALL, 5 multiple, 4 solid tumor		(25 of 50 patients) to 3 doses	
Reeves and Bestul.[10]	17	14 NHL, 3 AML	7.5 mg	1	
Campara et al.[11]	21	9 AML, 3 NHL, 3 multiple myeloma, 3	0.11-0.24 mg/kg	1	
		myelofibrosis, 2 chronic leukemia,			
		1 plasma cell leukemia			
Our study	90	7 multiple myeloma, 10 CLL, 6 CML, 22 ALL, 17 GCT, 09 Burkitt's lymphoma, 08 AML, 11 DLBCL	1.5 mg	1 or 2 doses depending on the response to single dose	

NHL – Non-Hodgkin's lymphoma; AML – Acute myeloid leukemia; CML – Chronic myeloid leukemia; CLL – Chronic lymphocytic leukemia; GCT – Germ cell tumor; ALL – Acute lymphocytic leukemia; CMML – Chronic myelomonocytic leukemia; MDS – Myelodysplastic syndrome; CMP – Common myeloid progenitors; DLBCL – Diffuse large B-cell lymphoma; ALL – Acute lymphoblastic leukemia

if necessary clinically) is very efficacious in TLS. We conclude that a low dose of rasburicase (1.5 mg or 3 mg) is cost saving and effective in reducing serum UA (especially for low-risk and intermediate-risk TLS) and the higher dose as recommended by the US FDA probably is required only for patients with high risk of TLS.

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Conflicts of interest

There are no conflicts of interest.

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