ejpmr, 2021,8(11), 449-451



EUROPEAN JOURNAL OF PHARMACEUTICAL AND MEDICAL RESEARCH

www.ejpmr.com

Research Article ISSN 2394-3211 EJPMR

STUDY OF ACUTE BLOOD TRANSFUSION REACTIONS IN PATIENTS OF BETA THALASSEMIA

Dr. Narahari Venkata Susmitha^{1*}, Dr. Priti Kamble², Dr. Chilukuru J. Satya Mani Deepika³, Dr. Suhas Kulkarni⁴ and Dr. Anil B. Kurane⁵

¹Junior Resident, Department of Paediatrics, Dr. DY Patil Medical College Hospital and Research Institute, Kolhapur, Maharashtra, India.

²Assistant Professor, Department of Paediatrics, Dr. DY Patil Medical College Hospital and Research Institute, Kolhapur, Maharashtra, India.

³Junior Resident, Department of Paediatrics, Dr. DY Patil Medical College Hospital and Research Institute, Kolhapur,

Maharashtra, India.

⁴Associate Professor, Department of Paediatrics, Dr. DY Patil Medical College Hospital and Research Institute, Kolhapur, Maharashtra, India.

⁵Professor and HOD, Department of Paediatrics, Dr. DY Patil Medical College Hospital and Research Institute, Kolhapur, Maharashtra, India.

*Corresponding Author: Dr. Narahari Venkata Susmitha

Junior Resident, Department of Paediatrics, Dr. DY Patil Medical College Hospital and Research Institute, Kolhapur, Maharashtra, India.

Article Received on 01/09/2021

Article Revised on 22/09/2021

Article Accepted on 12/10/2021

ABSTRACT

Background: The thalassemias are the most common genetic disorder on a worldwide basis.^[2] Early and regular blood transfusion decreases the complications of severe anemia and prolongs survival. Adverse events which occur in association with the transfusion of blood products are commonly called transfusion reactions.^[11] **Material and Methods:** This cross-sectional study included 54 beta(β) thalassemia patients whose age ranged between 1yr to 20 years. Pre transfusion hemoglobin assessment was done for all patients. Acute blood transfusion reactions were noted in details. **Results:** Most common transfusion reaction was fever. By fisher exact test, significant association was observed between pre transfusion Hemoglobin level and reaction. It is observed that % of reaction was more(22%) in patients with HB level <7.5, than in patients with HB level >7.5 (6%), which was statistically significant(P value< 0.0001). **Conclusion:** Acute Febrile non haemolytic reactions were common among patients with low pre- transfusion hemoglobin and Fever was the most common transfusion reaction noted.

KEYWORDS: β-thalassemia, blood transfusion, transfusion reactions, pre- transfusion haemoglobin.

INTRODUCTION

The thalassemias are the most common genetic disorder on a worldwide basis.^[1] Early and regular blood transfusion decreases the complications of severe anemia and prolongs survival. The recommended treatment for beta thalassaemia major involves lifelong regular blood transfusions, usually administered every two to five weeks, to maintain the pre transfusion hemoglobin level above 9-10.5 g/dl. Yet, transfusion carries the risk of complications. Therefore knowing different adverse effects of blood transfusion represents a great issue in managing thalassemia patients. Adverse events which occur in association with the transfusion of blood products are commonly called transfusion reactions.^[3] The aim of the present work is to study the prevalence of acute blood transfusion reactions among β-thalassemia major patients in the Thalassemia day care centre of Dr D Y Patil Hospital.

ACUTE BLOOD TRANSFUSION REACTIONS

| Febrile | Afebrile |
|------------------------|-------------------|
| Acute haemolytic | Allergic reaction |
| Febrile non-haemolytic | Anaphylaxis |
| Bacterial sepsis | |
| TRALI | |

AIM

To study acute blood transfusion reactions in patients of beta thalassemia.

OBJECTIVES

- \circ To know which type of transfusion reaction is common.
- To find out correlation between pre transfusion hemoglobin and acute blood transfusion reactions in beta thalassemia patients.

MATERIAL AND METHODS

This cross-sectional study included 54 beta(β) thalassemia patients whose age ranged between 1yr to 20 years coming for regular follow-up and blood transfusion at Thalassemia day care centre of Dr D Y Patil hospital in the last six months. After explaining the procedure to the patients and acquiring informed consent from the patients or guardians, 1ml of venous blood was withdrawn under complete aseptic conditions into EDTA vial to perform ABO and Rhesus(Rh) blood grouping and haemoglobin level before transfusion. All patients were subjected to thorough history taking with special emphasis on blood transfusions regarding rate of blood transfusion, type of received blood, and history of previous transfusion reactions. Packed red cells were transfused. Transfusion was begun within 30 minutes of removing from storage temperature (2-6 deg c) and completed within 4 hours, if ambient temperature was 22-25 deg c. In case of high ambient temperatures, shorter out of refrigerator time was used. Clinical assessment was done for each patient during the transfusion session in order to detect any transfusion reaction. Acute blood transfusion reactions were noted in detail.

RESULTS

Fever was the most common transfusion reaction noted.

| Type of reaction | No | % |
|--------------------|----|-------|
| Fever | 6 | 40 |
| Fever with chills | 5 | 33.33 |
| Chills | 1 | 6.66 |
| Rash | 1 | 6.66 |
| Red coloured urine | 2 | 13.33 |
| Headache | 2 | 13.33 |



| Association Of With Reaction | Hb Level | | |
|---------------------------------|----------|----|-------|
| | Reaction | | |
| | YES | NO | TOTAL |
| HB LEVEL | | | |
| <7.5 | 12 | 4 | 16 |
| >7.5 | 3 | 35 | 38 |
| TOTAL | 15 | 39 | 54 |

By fisher exact test, there was significant association observed between pre- transfusion hemoglobin level and reaction.

It was observed that % of reaction was more(22%) in patients with HB level <7.5, than in patients with HB level >7.5 (6%), which was statistically significant with P value (P< 0.0001).

DISCUSSION

The treatment of thalassemia major is blood transfusions to maintain the hemoglobin level.^[4] The present study was designed to explore the prevalence of acute blood transfusion reactions among 54 B- thalassemia patients who receive regular blood transfusion in the thalassemia day care centre of Dr D Y Patil hospital during a period of 6 months. Acute transfusion reactions were developed in 15 patients (28%), of which 12 patients with pretransfusion HB > 7.5 (22%) and 3 patients with per transfusion HB < 7.5 (6%). It shows that the blood transfusion reactions were common in the patients with low pre- transfusion hemoglobin. The mean pretransfusion hemoglobin level was 8.5±1.16 g/dl which is lower than similar studies in which median baseline hemoglobin concentration of 10.0 g/dl was observed.^[4] The low hemoglobin concentration level among the patients compared to similar studies may be related to the low availability of blood due to low rate of blood donation as this blood was given to the patients free of charge.

Allergic reactions developed in 15 patients during the period of study in which fever (40%) and fever with chills(33.33%) were more common. Other acute transfusion reactions noted were chills(6.66%), rash(6.66%), red coloured urine(13.33%) and headache(13.33%).

CONCLUSION

Fever was the most common transfusion reaction noted and acute febrile non haemolytic reactions were common among the patients with low pre- transfusion hemoglobin.

As transfusion reactions are more common in patients with pre transfusion hemoglobin less then 7.5gm/dl and hemoglobin drops by 1gm/dl per week after transfusion, we can counsel patients accordingly and can prevent reactions. As acute febrile reactions are more common, we can minimise them by giving anti pyretics and anti histaminics prior to transfusion.

REFERENCES

- Borgna-Pignatti C, Cappellini MD, De Stefano P, Del Vecchio GC, Forni GL, Gamberini MR, et al. Survival and complications in thalassemia. Ann N Y Acad Sci, 2005; 1054: 40–7. [PubMed] [Google Scholar]
- El-Beshlawy A, Kaddah N, Moustafa A, Mouktar G, Youssry I. Screening for β-thalassemia carriers in

Egypt: Significance of the osmotic fragility test. East Mediterr Health J, 2007; 13: 780–6. [PubMed] [Google Scholar]

- Katz EA. Blood transfusion: Friend or foe. AACN Adv Crit Care, 2009; 20: 155–63. [PubMed] [Google Scholar]
- Cario H, Stahnke K, Kohne E. Beta-thalassemia in Germany. Results of cooperative beta-thalassemia study. Klin Padiatr, 1999; 211: 431–7. [PubMed] [Google Scholar]
- Rebulla P. Transfusion reactions in thalassemia. A survey from the Cooley care. The Cooley care Cooperative Group. Haematologica, 1990; 75: 122–7. [PubMed] [Google Scholar]
- Climent-Peris C, Vélez-Rosario R. Immediate transfusion reactions. P R Health Sci J, 2001; 20: 229–35. [PubMed] [Google Scholar]
- Martínez-Díaz H, Frye-Maldonado AC, Climent-Perís C, Vélez-Rosario R. Evaluation of serologic markers for transfusion transmitted infectious diseases for allogeneic blood donors in Puerto Rico. P R Health Sci J, 1997; 16: 255–8. [PubMed] [Google Scholar]