

Study of discriminant factor M/H ratio in screening for β thalassemia trait

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Abstract

Introduction: Beta Thalassemia syndrome is a group of hereditary disorders characterized by genetic deficiency in the synthesis of beta globin chains located on chromosome 11. A major diagnostic challenge is to differentiate between mild microcytic hypochromic anaemia due to BTT from other causes such as iron deficiency (IDA), sideroblastic anaemia etc. The microcytic to hypochromic ratio (M/H ratio) is one of the simplest discriminant functions which is based on the fact that in Iron deficiency anaemia the RBCs are more hypochromic as compared to thalassemia trait. The present study was carried out in our diagnostic centre with the aim of evaluating the M/H ratio for screening patients for BTT.

Materials and Methods: This was a retrospective study carried out on a total of 200 patients between January to April 2018 coming to our diagnostic centre for complete blood count and Hb electrophoresis for evaluating the M/H ratio to screen for BTT. $HbA_2 > 3.5\%$ was considered to be diagnostic of BTT. M/H ratio as an indicator of differentiating between BTT and IDA was compared with other discriminant factors like England & Fraser, Mentzer Index, Shine & Lal Index and Shrivastava Index.

Results: There were 73 (36.5%) male and 127 (63.5%) females with a M:F ratio of 0.57:1. Out of 200 patients, 22 (11%) patients were diagnosed as BTT. Of these M/H ratio was <0.9 in 5 (22%) patients while >0.9 in 11 (77.27%) patients. England and Fraser index had highest specificity (94%) with a sensitivity of 23%. Shine and Lal had a specificity of 39% and sensitivity of 95%, Mentzer had a specificity of 83% and sensitivity of 45%, Shrivastava had sensitivity and specificity of 36% and 78% respectively while M/H ratio had specificity (70%) and sensitivity (77%).

Conclusion: M/H ratio provided by Advia 2120 hematology analyzer is as effective as preliminary screening tool for selection of samples for HbA_2 estimation. It is a rapid, automated formula provided without any additional cost to the patient.

Keywords: M/H ratio, BTT, IDA, Discriminant factor.

Introduction

Beta thalassemia syndrome is a group of hereditary disorders characterized by genetic deficiency in the synthesis of beta globin chains located on chromosome 11. The homozygous condition is called thalassemia major and is a severe transfusion dependent condition while heterozygotes have minor symptoms related to anemia and are called Beta thalassemia trait (BTT) or thalassemia minor.^{1,2} No specific treatment is needed for BTT patients. Thalassemias are more common in Mediterranean region, Africa, South East Asia with a prevalence rate of as high as 10% in these areas.

Thalassemia minor patients have a mild microcytic, hypochromic anaemia and are usually asymptomatic and do not have major morbid symptoms. However, a major diagnostic challenge is to differentiate between mild microcytic hypochromic anaemia due to BTT from other causes such as iron deficiency (IDA), sideroblastic anaemia etc.

A complete blood picture often gives an indication suggesting BTT due to presence of target cells and microcytic hypochromic RBCs on peripheral blood smear with an increased RBC count. An elevated HbA_2 level on Hb electrophoresis is confirmatory.

There are many discriminant formulas which have been published for distinguishing the cause of microcytosis as thalassemia based on RBC parameters.³⁻⁷ Some of these Discriminant functions are

simple to use while some require mathematical calculations and are complicated.³ The microcytic to hypochromic ratio (M/H ratio) is one of the simplest discriminant functions which is based on the fact that in Iron deficiency anaemia the RBCs are more hypochromic as compared to thalassemia trait.⁸⁻¹⁰

The Advia 2120 5-part hematology analyzer (Siemens) provides the percentage of microcytic to hypochromic cells. The M/H ratio is calculated by dividing the percentage of microcytic RBCs to the percentage of hypochromic RBCs and is automatically provided along with other CBC parameters by Advia 2120 hematology analyzer. A ratio of >0.9 is highly suggestive of BTT while ratio of <0.9 is suggestive of IDA.

The present study was carried out in our diagnostic centre with the aim of evaluating the M/H ratio for screening patients for BTT.

Materials and Methods

This was a retrospective study carried out on a total of 200 patients between January to April 2018 coming to our diagnostic centre for complete blood count and Hb electrophoresis for evaluating the M/H ratio to screen for BTT.

The patients of both sexes and all age groups were included in the study. The patients were divided into 0-10, 11-20, 21-30, 31-40, 41-50, 51-60 and more than 60 years of age in both the sexes. Venous blood samples

were collected into K2EDTA tubes and were subjected to complete blood count on Advia 2120 within one hour of sample collection.

The Advia analyzer was calibrated and 3 levels of controls were run before analyzing the specimens. HbA₂ was assayed on BIO-RAD D10 HPLC electrophoresis analyzer (Bio-Rad Laboratories Hercules, CA, USA). All patients with Hb <12 gm% (WHO criteria for anaemia) were included in the study. HbA₂ > 3.5% was considered to be diagnostic of BTT. Differential values for different discriminant indices were applied as defined in their original reports like Mentzer Index, Shine & Lal Index, England & Fraser Index and Shrivastava index.¹¹⁻¹⁴

Results

A total of 200 patients of both sexes and all ages were included in the study. There were 73 (36.5%) male and 127 (63.5%) females with a M:F ratio of 0.57:1.

Maximum patients 76 (38%) were in the 21 – 30 years of age group followed by 42 (21%) in 0-10 and 35 (17.5%) in 31 – 40 years of age and 35 (17.5%) in 11-20 years of age group. (Table 1)

The Hb% concentration between 9-11 gm% was found in 44 (22%) patients, followed by 43 (21.5%) between 7.1 – 9.0 gm%. Maximum patients had Hb >11

gm% i.e. 66(30.5%). Only 20 patients (10%) had Hb below 5 gm% (Table 2) RBCs count <5.0 million/cumm was found in 157 patients (78.5%) while RBC count >5.0 million/cumm was found in 43 (21.5%) patients. An interesting observation was that 13 BTT cases had RBC count below 5 million/ cumm as compared to 9 who had RBC count greater than 5.0 million/cumm (Table 3). Out of 200 patients, 22 (11%) patients were diagnosed as BTT. Of these M/H ratio was <0.9 in 5 (22%) patients while >0.9 in 11 (77.27%) patients (Table 4).

When M/H ratio was divided into different categories it was observed that 50% patients of BTT had M/H ratio above 1.80. 13.6% had M/H ratio between 1.21 – 1.50 and 4.54% had M/H ratio between 1.51 – 1.80.

The mean age was 22.9 ± 12.9 years. The mean Hb was 9.3 ± 3.2 gm%. Mean RBC count was $4.2 \pm 1.0 \times 10^3$ /ul, mean PCV was $30.4 \pm 9.5\%$, Mean MCV was 73.6 ± 20.0 fl, Mean M/H ratio was 0.89 ± 0.9 and Mean HbA₂ was $2.8 \pm 1.0\%$.

M/H ratio as an indicator of differentiating between BTT and IDA was compared with other discriminant factors like England & Fraser, Mentzer Index, Shine & Lal Index and Shrivastava Index (Table 5).

Table 1: Showing demographic data of patients

Age in years	Male	Female	Total	%
0 - 10	29	13	42	21
11.0 - 20	12	23	35	17.5
21 - 30	21	55	76	38
31 - 40	6	29	35	17.5
41 - 50	2	4	6	3
51 - 60	2	3	5	2.5
> 60	1	0	1	0.5
Total	73 (36.5%)	127 (63.5%)	200	

Table 2: Showing distribution of patients according to hematological parameters

Hb (gm%)	Male	Female	Total	%
<5.0	8	12	20	10
5.1 - 7.0	16	16	32	16
7.1 -9.0	10	33	43	21.5
9.1 - 11.0	11	33	44	22
>11.0	28	33	61	30.5
Total	73	127		
RBC Count	Male	Female	Total	%
<5.0	44	113	157	78.5
>5.0	29	14	43	21.5
Total	73	127		

Table 3: Showing distribution of patients according to RBC count

RBC Count	Normal	Thalassemia Minor
<5.0	144	13
>5.0	34	9
Total	178 (89%)	22 (11%)

Table 4: Showing distribution of patients according to micro/hypo ratio

Micro/Hypo Ratio	Normal		Thalassemia Minor	
	Total Number	%	Total Number	%
<0.9	124	69.6	5	22.7
>0.9	54	30.3	17	77.27
Total	178		22	
Micro/Hypo	Thalassemia Minor		Normal	
	Total Patients	Percent	Total Patients	Percent
</= 0.2	2	9.09	43	24.1
0.3 - 0.60	0	0	45	25.2
0.61 - 0.90	3	13.6	36	20.4
0.91 - 1.20	2	9.09	27	15.1
1.21 - 1.50	3	13.6	8	4.49
1.51 - 1.80	1	4.54	5	2.8
> 1.80	11	50	14	7.86
Total	22		178	

Table 5: Comparison of discriminant functions for identifying thalassemia in patients

Discriminant function	Formula	Optimal Cut off	T.Minor		Normal HbA2		Sensitivity	Specificity
			<0	>0	<0	>0		
England & Fraser	MCV-RBC-(5xHb)-3.4	0	5	17	11	167	23%	94%
			<13	>13	<13	>13		
Mentzer	MCV/RBC	13	10	12	31	147	45%	83%
			<1530	>1530	<1530	>1530		
Shine & Lal	MCV ² xMCH/100	1530	21	1	108	70	95%	39%
			<3.8	>3.8	<3.8	>3.8		
Srivastava	MCH/RBC	3.8	9	13	40	138	36%	78%
			>0.9	<0.9	>0.9	<0.9		
M/H Ratio	MIC/HPO	0.9	17	5	54	124	77%	70%

Table 6: Sensitivity, specificity, positive predictive value (PPV), negative predictive value (NPV), and Youden's index to discriminate between BTT and normal HbA2

Indices	Sensitivity	Specificity	PPV	NPV	Youden Index (sensitivity+specificity)- 100
Mentzer					
BTT	45%	83%	24.4%	92.5%	28%
Normal HbA2	83%	45%	92.5%	24.4%	
Shine & Lal					
BTT	95%	39%	16.3%	98.6%	34%
Normal HbA2	39%	95%	98.6%	16.3%	
Srivastava					
BTT	36%	78%	18.4%	91.4%	14%
Normal HbA2	78%	36%	91.4%	18.4%	
England & Fraser					
BTT	23%	94%	31.3%	90.8%	17%
Normal HbA2	94%	23%	90.8%	31.3%	
M/H Ratio					
BTT	77%	70%	23.9%	3.9%	7%
Normal HbA2	70%	77%	3.9%	23.9%	

Table 6: Shows the sensitivity, specificity, PPV, NPV and Youden Index of all discriminant factors.

Discussion

Anaemia is the most common presenting symptom of a variety of diseases. Iron deficiency anaemia and

Beta thalassemia trait being common causes of anaemia. However, differentiating between IDA and BTT is a challenging problem and has important

clinical implications because both of these have different causes, prognosis and treatment. A missed case of BTT may have potential homozygous offspring. Hence, it is very crucial to have a high index of suspicion for BTT based on CBC parameters so that it can be diagnosed and proper counseling of family members can be done.¹⁵ Many discriminant indices have been proposed to differentiate and screen for IDA and BTT but none has 100% specificity or sensitivity.

An ideal discriminant index should have high sensitivity and specificity. A combination of two or more CBC parameters in discriminant indices can lead to a great improvement in screening of anaemic patients for BTT.^{16,17}

In recent years several studies have been carried out to assess the usefulness in the differential diagnosis of BTT by using M/H ratio.¹⁸⁻²⁰

In IDA, RBCs are produced which have decreased Hb concentration and high hypochromic cell population while in BTT, microcytes are generally smaller with more preserved Hb concentration and is characterized by increase in RBC% due to chronic increase in erythropoiesis.

In our study, we found that England and Fraser index had highest specificity (94%) with a sensitivity of 23%. Shine and Lal had a specificity of 39% and sensitivity of 95%, Mentzer had a specificity of 83% and sensitivity of 45%, Srivastava had sensitivity and specificity of 36% and 78% respectively while M/H ratio had specificity (70%) and sensitivity (77%).

In the study conducted by Eloisa et al, Shine and Lal provided the best sensitivity (100%) but low specificity (22.8%). Srivastava index had a good specificity (91.4%) but a sensitivity of 86.8%.²¹ In their study the M/H ratio shared a sensitivity of 99.2% and specificity of 77.1%.

In the study by A. Vehapoglu et al. they found Mentzer index to be most reliable index with the highest sensitivity of 98.7% and specificity of 82.3%.²²

Al Fadhli et al in their study concluded that England and Fraser index had the highest Youden index value (98.2%) for differentiating between IDA and BTT.²³

In a study conducted by Ehsani et al in 2009, Mentzer Index (90%) was found to be the best index for differentiating IDA and BTT.²⁴ Similar results indicating Mentzer Index to be highly sensitive (90.9%) was observed by Ghafouri et al.²⁵

A study by Kabir AL et al, however showed 96% sensitivity for M/H ratio with a predictive value of 90.4% in screening for BTT.²⁶ We observed that with a cut off value of 1.8 for M/H ratio, we could diagnose BTT with maximal diagnostic sensitivity but at expense of specificity.

Our study. However has some limitations as the inclusion criteria may be different from other studies as our aim was to investigate patients with microcytosis for BTT rather than for IDA. Moreover, we included

subjects with normal Hb also who were registered for Hb electrophoresis as a part of family studies. Moreover, our sample size is also small which may have lead to difference in sensitivity and specificity of M/H ratio as well as Youden index in our study as compared to other studies.

Conclusion

Several discriminant indices have been in use to differentiate between IDA & BTT and M/H ratio provided by Advia 2120 hematology analyzer is as effective as preliminary screening tool for selection of samples for HbA₂ estimation. It is a rapid, automated formula provided without any additional cost to the patients. Large prospective population studies are needed to assess the efficiency of M/H ratio for screening patients with BTT.

Conflict of Interest: None

References

1. [Rachmilewitz EA, Giardina PJ. How I treat thalassemia. *Blood*. 2011 Sep 29. 118(13):3479-88.
2. Galanello R, Sanna S, Perseu L, Sollaino MC, Satta S, Lai ME, et al. Amelioration of Sardinian beta0 thalassemia by genetic modifiers. *Blood*. 2009 Oct 29. 114(18):3935-7.
3. Eldibany MM, Totonchi KF, Joseph NJ, Rhone D. Usefulness of certain red blood cell indices in diagnosing and differentiating thalassemia trait from iron-deficiency anemia. *Amer J Clin Pathol* 1999;111:676-82.
4. Ntaios G, Chatzinikolaou A, Saouli Z, Girtovitis F, Tsapanidou M, Kaiapa G, Kontoninas Z, Nikolaidou A, Savopoulos C, Pidonia I, Alexiou-Daniel S. Discrimination indices as screening tests for beta-thalassaemic trait. *Ann Hematol* 2007;86:487-91.
5. Urrechaga E. Discriminant value of % microcytic/% hypochromic ratio in the differential diagnosis of microcytic anemia. *Clin Chem Lab Med* 2008;46:1752-8.
6. Urrechaga E, Borque L, Escanero JF. The role of automated measurement of red cell subpopulations on the Sysmex XE 5000 analyzer in the differential diagnosis of microcytic anemia. *Int J Lab Hematol* 2011;33:30-6.
7. Beyan C, Kaptan K, Ifran A. Predictive value of discrimination indices in differential diagnosis of iron deficiency anemia and beta-thalassemia trait. *Eur J Haematol* 2007;78:524-6.
8. Urrechaga E. Red blood cell microcytosis and hypochromia in the differential diagnosis of iron deficiency and β -thalassaemia trait. *Int J Lab Hematol* 2009;31:528-34.
9. d'Onofrio G, Zini G, Ricerca BM, Mancini S, Mango G. Automated measurement of red blood cell microcytosis and hypochromia in iron deficiency and beta-thalassemia trait. *Arch Pathol Lab Med* 1992;116:84-9.
10. Lafferty JD, Crowther MA, Ali MA, Levine M. The evaluation of various mathematical RBC indices and their efficacy in discriminating between

- thalassemic and non-thal- assemic microcytosis. *Amer J Clin Pathol* 1996;106:201–5.
11. W. C. Mentzer Jr., “Differentiation of iron deficiency from thalassaemia trait,” *The Lancet*, vol. 1, no. 7808, p. 882, 1973.
 12. I. Shine and S. Lal, “A strategy to detect β thalassaemia minor,” *The Lancet*, vol. 1, no. 8013, pp. 692–694, 1977.
 13. J. M. England and P. M. Fraser, “Differentiation of iron deficiency from thalassaemia trait by routine blood-count,” *The Lancet*, vol. 1, no. 7801, pp. 449–452, 1973.
 14. P. C. Srivastava, “Differentiation of thalassemia minor from iron deficiency,” *The Lancet*, vol. 2, pp. 154–155, 1973.
 15. Giordano PC, Bouva MJ, Harteveld CL. A confidential inquiry estimating the number of patients affected with sickle cell disease and thalassemia major confirms the need for a prevention strategy in the Netherlands. *Hemoglobin* 2004;28:287–96.
 16. Eldibany MM, Totonchi KF, Joseph NJ, Rhone D. Usefulness of certain red blood cell indices in diagnosing and differentiating thalassemia trait from iron-deficiency anemia. *Am J Clin Pathol* 1999;111:676–82.
 17. Demir A, Yarali N, Fisgin T, Duru F, Kara A. Most reliable indices in differentiation between thalassemia trait and iron deficiency anemia. *Pediatr Int* 2002;44:612–6.
 18. d’Onofrio G, Zini G, Ricerca BM, Mancini S, Mango G. Automated measurement of red blood cell microcytosis and hypochromia in iron deficiency and beta-thal- semia trait. *Arch Pathol Lab Med* 1992;116:84–9.
 19. Robertson EP, Pollock A, Yau KS, Chan CL. Use of Tech- nicon H*1 technology in routine thalassemia screening. *Med Lab Sci* 1992;49:259–64.
 20. Jimenez CV, Minchinela J, Ros J. New indices from the H*2 analyser improve differentiation between heterozygous beta or delta beta thalassaemia and iron deficiency anaemia. *Clin Lab Haem* 1995;17:151–5.
 21. Elo’s Urrechaga Discriminant value of % microcytic/% hypochromic ratio in the differential diagnosis of microcytic anemia *Clin Chem Lab Med* 2008;46(12):1752–1758 .
 22. Aysel Vehapoglu, Gamze Ozgurhan, Aysegul Dogan Demir, Selcuk Uzuner, Mustafa Atilla Nursoy, Serdar Turkmen, and Arzu Kacan Hematological Indices for Differential Diagnosis of Beta Thalassemia Trait and Iron Deficiency Anemia: Anaemia volume 2014 (2014).
 23. S. M. AlFadhli, A. M. Al-Awadhi, and D. AlKhalidi, “Validity assessment of nine discriminant functions used for the differentiation between Iron deficiency anemia and thalassemia minor,” *Journal of Tropical Pediatrics*, vol. 53, no. 2, pp. 93–97, 2007.
 24. A. Ehsani, E. Shahgholi, M. S. Rahiminejad, F. Seighali, and A. Rashidi, “A new index for discrimination between iron deficiency anemia and beta-thalassemia minor: results in 284 patients,” *Pakistan Journal of Biological Sciences*, vol. 12, no. 5, pp. 473–475, 2009.
 25. M. Ghafouri, L. Mostaan Sefat, and L. Sharifi, “Comparison of cell counter indices in differentiation of beta thalassemia trait and iron deficiency anemia,” *The Scientific Journal of Iranian Blood Transfusion Organization*, vol. 2, no. 7, pp. 385–389, 2006.
 26. Kabir AL, Dipta TF, Rahman MH, Mahfuz H, Ahmed M, Rahman M, Nasreen T Auto-analyzer based screening of microcytic hypochromic ratio to differentiate thalassaemia and non *Bangladesh Med Res Counc Bull* 2013;39:146-147.