

2015

Is Hemoglobin Variant Analysis Helpful in the Diagnostic Work-up of Patients Revealing Microcytic Erythrocytosis on Complete Blood Count?

Upasana Joneja, MD

Thomas Jefferson University Hospital, Philadelphia, PA

Alina Dulau-Florea

NIH, Bethesda MD

Gene Gulati, Ph.D

Department of Pathology, Anatomy, and Cell Biology, Jefferson Medical College of Thomas Jefferson

University, Philadelphia, PA works at: <https://jdc.jefferson.edu/pacbresidentposters>



is part of the Medical Anatomy Commons, Medical Cell Biology Commons, and the Medical Pathology

Commons at Thomas Jefferson University

© 2015 Joneja, MD, Dulau-Florea, Alina; Gulati, Ph.D, Gene; and Gong, MD, Jerald Z.

Thomas Jefferson University

[Let us know how access to this document benefits you](https://jdc.jefferson.edu/pacbresidentposters/16)

Recommended Citation

Joneja, MD, Upasana; Dulau-Florea, Alina; Gulati, Ph.D, Gene; and Gong, MD, Jerald Z., "Is Hemoglobin Variant Analysis Helpful in the Diagnostic Work-up of Patients Revealing Microcytic Erythrocytosis on Complete Blood Count?" (2015). *Department of Pathology, Anatomy, and Cell Biology Resident's Posters*. Paper 16.

<https://jdc.jefferson.edu/pacbresidentposters/16>

This Article is brought to you for free and open access by the Jefferson Digital Commons. The Jefferson Digital Commons is a service of Thomas Jefferson University's [Center for Teaching and Learning \(CTL\)](#). The Commons is a showcase for Jefferson books and journals, peer-reviewed scholarly publications, unique historical collections from the University archives, and teaching tools. The Jefferson Digital Commons allows researchers and interested readers anywhere in the world to learn about and keep up to date with Jefferson scholarship. This article has been accepted for inclusion in Department of Pathology, Anatomy, and Cell Biology Resident's Posters by an authorized administrator of the Jefferson Digital Commons. For more information, please contact: JeffersonDigitalCommons@jefferson.edu.

Is Hemoglobin Variant Analysis Helpful in the Diagnostic Work-up of Patients Revealing Microcytic Erythrocytosis on Complete Blood Count?

Upasana Joneja¹, Alina Dulau-Florea², Gene L. Gulati¹, Jerald Z. Gong¹

¹Department of Pathology, Cell Biology and Anatomy, Thomas Jefferson University Hospital, Philadelphia, PA

²Department of Laboratory Medicine, Clinical Center, National Institutes of Health, Bethesda, MD

INTRODUCTION

Microcytic erythrocytosis is an abnormal CBC (complete blood count) finding that is under-recognized, poorly understood, and consequently under-utilized in patient care. It is characterized by decreased MCV and increased RBC count. Its etiology is likely multifactorial and includes thalassemias and hemoglobinopathies. The focus of our study was to determine the relative prevalence of hemoglobin-associated disorders in patients revealing microcytic erythrocytosis on CBC and to demonstrate whether or not hemoglobin variant analysis should be included in the diagnostic work-up of such cases.

METHODS

Data from all CBCs within a 2 year period at Jefferson were reviewed, and patients meeting the inclusion criteria of microcytosis (MCV <80 fL) and erythrocytosis (RBC count $\geq 5.2 \times 10^6$ T/L (Female) and $> 6 \times 10^6$ T/L (Male)) were selected. The sample size was 137 patients with 87 females and 50 males; ages ranged from: 1-91 years with median age of 46 years. The racial distribution comprised of 49 Caucasians, 44 African Americans, 24 Asians and 20 unspecified. Hemoglobin values for the patients ranged from 7.2– 16.1 g/dL, MCVs ranged from 54.5-79 fL and RBC count ranged from $5.2-7.5 \times 10^6$ T/L. 74 patients had mild anemia while 63 patients had hemoglobin levels within the gender-specified reference range. The results of hemoglobin variant analysis performed by high-performance liquid chromatography on these patients were reviewed.

RESULTS

Hemoglobin Variant Analysis Pattern Consistent With	Number of Patients	Percentage of Patients
Beta-thalassemia trait	70	51.1
Delta/beta-thalassemia trait	1	0.7
Hemoglobin E disease	4	2.9
Hereditary persistence of fetal hemoglobin (HPFH)	1	0.7
Possible HPFH	4	2.9
Beta-thalassemia with HPFH	2	1.5
Delta/beta-thalassemia with iron deficiency anemia	1	0.7
Hemoglobin C trait with beta-thalassemia	1	0.7
Hemoglobin C trait with possible alpha-thalassemia	2	1.5
Hemoglobin C with HPFH	1	0.7
Unidentified hemoglobinopathy with possible delta/beta-thalassemia	2	1.5
Hemoglobin S trait combined with beta and possible alpha-thalassemia	3	2.2
Hemoglobin S trait with beta-thalassemia	1	0.7
Normal pattern	44*	32.1
Total # of patients	137	100

Table 1: Relative prevalence of hemoglobin-associated conditions in patients revealing microcytic erythrocytosis on CBC.

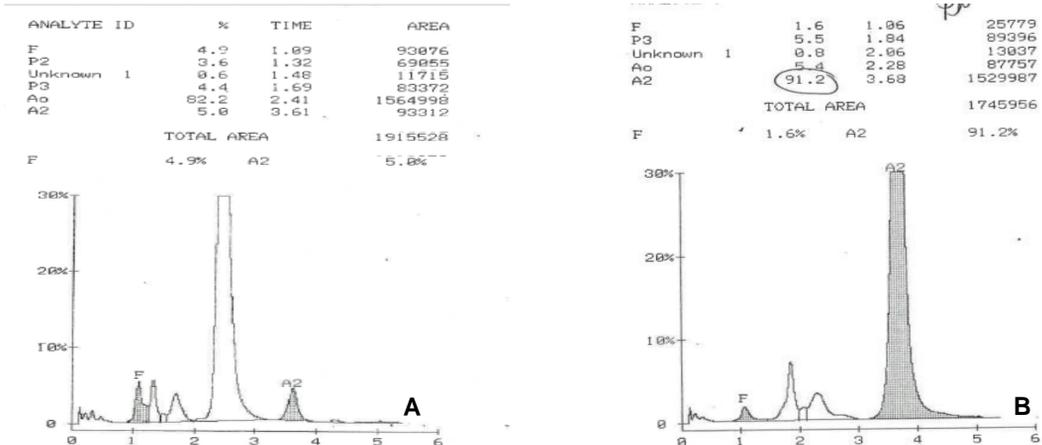


Figure 1A) HPLC results of a patient with beta thalassemia trait B) HPLC results of a patient with Hemoglobin E disease

DISCUSSION

- 93 of the 137 (67.9%) had a thalassemia trait and/or a hemoglobinopathy. Amongst these, beta-thalassemia trait topped the list, followed by hemoglobin E disease, possible HPFH, HPFH and delta-beta-thalassemia.
- The diagnosis of possible HPFH was entertained based on mild elevation of hemoglobin F (2.3% to 3.1%) by HPLC. Since HPFH is typically associated with normal MCV or RBC count, the microcytic erythrocytosis noted in these patients likely represent an associated undiscovered alpha-thalassemia trait.
- Majority, if not all, of the 44 cases with normal hemoglobin variance analysis pattern also represent patients with possible α -thalassemia trait, Microcytic erythrocytosis is uncommon in normal patients and alpha thalassemia is a diagnosis of exclusion (requires DNA analysis studies not performed here)
- Compound heterozygous conditions for either hemoglobinopathies and/or thalassemia(s) comprised 12 cases. Unspecified abnormal hemoglobins detected by the HPLC method remain unidentified as no additional analyses had been performed in both cases.
- The diagnosis of delta/beta-thalassemia and concomitant iron deficiency anemia made in 1 patient was supported by the laboratory findings of increased hemoglobin F, decreased hemoglobin A2 and decreased serum ferritin.
- With all these potential considerations, over 96% of the studied cases of microcytic erythrocytosis may be categorized as having a possible thalassemia trait and/or a hemoglobinopathy.

CONCLUSION

The answer to the question our study title proposed i.e. “Is Hemoglobin Variant Analysis Helpful in the Diagnostic Work-up of Patients Revealing Microcytic Erythrocytosis on Complete Blood Count?” is unequivocally “yes”. Based on the observed extremely high positive yield of hemoglobin variant analysis in determining the etiology of erythrocytosis, we recommend that the abnormality of microcytic erythrocytosis should automatically be flagged and such specimens should be reflexively processed for hemoglobin variant analysis for each previously undiagnosed patient.