

To: Mauli Childrens Hospital And Maternity Home-Aurangabad

Opp Gajanan Mandir, Beside Malkapur Bank,

Pudling Nagar Road, Grkheda Parisar

Maharashtra

Aurangabad - 431001

Contact:

Report Of: Mrs. DR PRATIKSHA JAIN

Pt. Contact: 7276473733



Sample ID	2400207149
Patient ID	10024100031
Collected on	13/11/2024
Received on	14/11/2024 16:37
Registered on	14/11/2024 18:22
Reported on	15/11/2024 13:42
Referred by	Dr. Archana Patil

Hemoglob	inopathy Screening	
Patient Name: Mrs. DR PRATIKSHA JAIN	Sample Type: Whole Blood EDT/	Α
Date of Birth/Age: 05/11/1993	Gender: FEMALE	City: Aurangabad
Method: High Performance Liquid Chromatography (HPLC)	Blood Transfusion History: No	
Referral Reason or Clinical History:		

About the test

Hemoglobinopathy screening by high performance liquid chromatography is a blood test that is used for detecting quantitative and qualitative abnormalities of hemoglobin (Hb), namely, Thalassemia and Structural Hb variants (e.g. HbS) respectively. The test helps identify individuals with these disorders so that they can receive timely and appropriate treatment and care. Antenatal diagnosis of these disorders allows measures to reduce the chances of the birth of an affected baby. It is also possible to screen the newborns for hemoglobinopathies using this approach, thereby decreasing the mortality & morbidity associated with conditions like Sickle cell disorder.

	Test findings	
Hb Fraction	Observed Value (%)	Expected Value (%)
HbF	0.3%	<2%
P2*	3.5%	<4.6%
HbA0	87.1%	85 - 95%
HbA2/HbE	2.9%	1.8 - 3.5%
HbD	Absent	Absent
HbS	Absent	Absent

	Interpretation			
Chromatogram shows normal hemoglobin pattern. Hemoglobin and RBC parameters are normal.				

Verified by

Mr. Pradip Kadam
Incharge Biochemistry

Dr.Suresh Bhanushali MD (Path), Consultant Pathologist Page 1 of 2





Patient Name: Mrs. DR PRATIKSHA JAIN Sample ID: 2400207149

HPLC Findings

A2 Concentration = 2.9 %

Analysis comments:

Patient Data
Sample ID: 2400207149
Patient ID:
Name:
Physician:
Sex:
DOB:

Unknown

Unknown

P2

Α2

Comments:

 Analysis Performed:
 11/14/2024 22:38:16

 Injection Number:
 9124

 Run Number:
 596

 Rack ID:
 0004

 Tube Number:
 10

 Report Generated:
 11/14/2024 22:46:04

Operator ID:

Area

0.1

1.0

3.5

87.1

용

Retention Time (min)

0.90

1.01

1.09

1.21

1.33

1.72

2.38

3.67

Calibrated

0.3

2.9

	45.0		
	37.5		
	30.0		
96	22.5		
	15.0	1,000 1,000	
	7.5	2 3 8 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	

Time (min.)

Total Area: 2,646,521

660

2258

27768

92456

86314

2305484

F Concentration = 0.3
A2 Concentration = 2.9

Important Blood Indices (from CBC Analysis)

Parameters	Result	Reference Range	Units
Hemoglobin (Hb)	12.88	12 - 15	g/dL
RBC Count	3.96	3.8 - 4.8	x 10 ⁶ /μL
Hematocrit	39.70	36 - 46	%
Mean Corpuscular Volume (MCV)	100.20	83 - 101	fL
Mean Corpuscular Hb (MCH)	32.50 🔥	27 - 32	pg
Mean Corpuscular Hb Conc. (MCHC)	32.40	31.5 - 34.5	g/dL
RBC Distribution Width (RDW) (CV)	13.30	11.6 - 14	%
RBC Distribution Width (RDW) (SD)	45.50	39 - 46	fL

Notes:

- 1. Recent blood transfusions and iron deficiency can interfere with the results, repeat testing is recommended three months after the last blood transfusion. In case of iron deficiency, it is recommended to evaluate the result post-correction of iron deficiency.
- 2. Megaloblastic anemia can cause elevated HbA2 levels. A repeat assay is recommended after correction of VitB12 deficiency.
- 3. Mild to moderately elevated fetal hemoglobin (HbF) values are observed during pregnancy, hypoxia, chronic kidney disease, use of certain drugs,myelodysplastic syndromes (MDS), aplastic anemia and conditions of stress hemopoiesis.
- 4. Cases with borderline HbA2 levels (3.1-3.9%) could represent Silent Beta-thalassemia trait, or co-existent iron deficiency or Alpha-thalassemia in a case of Beta-thalassemia trait. They need to be investigated further by appropriate tests.
- 5. Confirmatory molecular tests for Beta-thalassemia traits and abnormal hemoglobin disorders (e.g. HbS, HbE, and HbD), followed by subsequent prenatal diagnosis (If required) are available at our centre.
- $6. \quad \text{The mentioned P2 value from BioRad Variant-II HPLC system is equivalent of HbA1c value in BioRad D10 system} \\$

Disclaimers:

- 1. The Hb-HPLC is a screening test that detects Beta-thalassemia and other hemoglobin variants. It does not identify Alpha-thalassemia and Silent Beta-thal-assemia carriers. DNA analysis is recommended to rule out Alpha-thalassemia and Silent Beta-thalassemia carriers.
- 2. The result must be interpreted in conjunction with the complete blood counts (CBC), VitB12 and iron profile of the individual.
- 3. Each sample received at Lilac Insights' processing centre is handled with the utmost sensitivity and care. All samples received on Sundays and National holidays are stored as per specific guidelines for the respective specimens and processed on the next day.
- 4. P2 peak in Bio Rad's Variant II HPLC platform represents glycated hemoglobin. It is elevated in uncontrolled diabetes.

Verified by

Mr. Pradip Kadam

Incharge Biochemistry

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Dr.Suresh Bhanushali MD (Path), Consultant Pathologist Page 2 of 2

