To: Vatsalya Maternity and Nursing Home-Virar(E)

Ground Floor Mohak Chambers 2.

Near Rane Talav, Manvelpada,

Talav, Manvelpada, Virar East, Palghar

Maharashtra

Virar - 401305

Contact: 8830521446

Report Of: Mrs. RUPALI AMOL PAWAR

Pt. Contact: 8108926402



Sample ID	2300118044
Patient ID	1002356348
Collected on	14/07/2023
Received on	20/07/2023 10:19
Registered on	19/07/2023 18:53
Reported on	20/07/2023 13:34
Referred by	DR.SUNITA R.MORE

Hemoglobinopathy Screening				
Patient Name: Mrs. RUPALI AMOL PAWAR	Sample Type: Whole Blood EDTA	4		
Date of Birth/Age: 12/06/1993	Gender: FEMALE	City: MUMBAI		
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	0.4%	<2%	
P2*	3.8%	<4.6%	
HbA0	86.7%	85 - 95%	
HbA2/HbE	3.0%	1.8 - 3.5%	
HbD	ABSENT	Absent	
HbS	ABSENT	Absent	

^{*}The mentioned P2 value from BioRad Variant-II HPLC system is equivalent of HbA1c value in BioRad D10 system

Interpretation

Chromatogram shows normal hemoglobin pattern.

Hemoglobin, PCV and RBC count are reduced. However, red cell indices are normocytic normochromic.

Suggestions

Please correlate clinically.

Verified by Mr. Pradip Kadam Incharge Biochemistry

Bede

Dr. A. Dasgupta MD, PhD, Consultant Hematopathologist

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HPLC Findings

Patient Data 2300118044 Sample ID: Patient ID: Name: Physician: Sex: DOB:

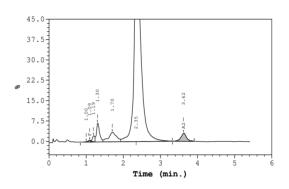
Comments:

Analysis Data Analysis Performed: Injection Number: 07/15/2023 13:03:26 Run Number: 315 Rack ID: Tube Number:

Report Generated: Operator ID: 07/15/2023 14:03:44

	Calibrated	Calibrated		Peak	
Peak Name	Area %	Area %	Time (min)	Area	
Unknown		0.1	1.00	1242	
F	0.4		1.09	7972	
Unknown		1.0	1.19	19745	
P2		3.8	1.30	71805	
P3		5.0	1.70	94604	
Ao		86.7	2.35	1633536	
A2	3.0		3.62	55422	

Total Area: 1,884,325



Analysis comments:

F Concentration = 0.4 A2 Concentration = 3.0

Important Blood Indices (from CBC Analysis)

Parameters	Result	Reference Range	Units
Hemoglobin (Hb)	10.74 🗥	12 - 15	g/dL
RBC Count	3.54 ▲	3.8 - 4.8	x 10 ⁶ /μL
Hematocrit	29.60 🗥	36 - 46	%
Mean Corpuscular Volume (MCV)	83.60	83 - 101	fL
Mean Corpuscular Hb (MCH)	30.30	27 - 32	pg
Mean Corpuscular Hb Conc. (MCHC)	36.30 ▲	31.5 - 34.5	g/dL
RBC Distribution Width (RDW) (CV)	15.40 	11.6 - 14	%
RBC Distribution Width (RDW) (SD)	40.80	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.
- Mild to moderately elevated fetal hemoglobin (HbF) values are observed during pregnancy, hypoxia, chronic kidney disease, use of certain 3. drugs, myelodysplastic syndromes (MDS), aplastic anemia and conditions of stress hemopoiesis.
- Cases with borderline HbA2 levels (3.1-3.9%) could represent Silent Beta-thalassemia trait, or co-existent iron deficiency or Alpha-thalassemia 4. in a case of Beta-thalassemia trait. They need to be investigated further by appropriate tests.
- Confirmatory molecular tests for Beta-thalassemia traits and abnormal hemoglobin disorders (e.g. HbS, HbE, and HbD), followed by subsequent 5. prenatal diagnosis (If required) are available at our centre.
- The mentioned P2 value from BioRad Variant-II HPLC system is equivalent of HbA1c value in BioRad D10 system

Disclaimers:

- 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.
- Each sample received at Lilac Insights' processing centre is handled with the utmost sensitivity and care. All samples received on Sundays and 3. National holidays are stored as per specific guidelines for the respective specimens and processed on the next day.
- 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. A. Dasgupta MD, PhD, Consultant Hematopathologist Page 2

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