

LABORATORY TEST REPORT

Name	: Mrs. AYSHA JULEKHA	
Sample ID	: B3585048	
Age/Gender	: 29 Years/Female	Reg. No : 0692508260016
Referred by	: Dr. SELF	SPP Code : SPL-AS-006
Referring Customer	: HEALTH ZONE	Collected On : 26-Aug-2025 11:14 AM
Primary Sample	: Whole Blood	Received On : 27-Aug-2025 03:49 PM
Sample Tested In	: Serum	Reported On : 28-Aug-2025 11:50 AM
Client Address	:	Report Status : Final Report

CLINICAL BIOCHEMISTRY

Test Name	Results	Units	Biological Reference Interval
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[PDF Attached](#)

Double Marker

Free -Beta -HCG (Method: CLIA)	37.84	ng/mL	< 2 :Non-Pregnant 5.4 - 393.4 : Pregnant
PAPP-A (Method: CLIA)	3.08	mIU/mL	< 0.1 : Non-Pregnant 0.1-19.5 : Pregnant

Interpretation:

DISORDER	SCREEN POSITIVE/HIGH RISK CUT OFF
Trisomy 21 (Down)	< 1:250
Trisomy 18/13	< 1:100
DISORDER	SCREEN NEGATIVE/LOW RISK CUT OFF
Trisomy 21 (Down)	> 1:250
Trisomy 18/13	> 1:100

Note:Statistical evaluation has been done using CE marked PRISCA 5 software. · Screening tests are based on statistical analysis of patient demographic and biochemical data. They simply indicate a high or low risk category. Confirmation of screen positives is recommended by Chorionic Villus Sampling (CVS). · The interpretive unit is MoM (Multiples of Median) which takes into account variables such as gestational age (ultrasound), maternal weight, race, insulin dependent Diabetes, multiple gestation, IVF (Date of Birth of Donor, if applicable), smoking & previous history of Down syndrome. Accurate availability of this data for Risk Calculation is critical. · Ideally all pregnant women should be screened for Prenatal disorders irrespective of maternal age. The test is valid between 9-13.6 weeks of gestation, but ideal sampling time is between 10-13 weeks gestation. · First trimester detection rate of Down syndrome is 60% with a false positive rate of 5%. A combination of Nuchal translucency, Nasal bone visualization and biochemical tests (Combined test) increases the detection rate of Down syndrome to 85% at the same false positive rate.

Comments:First trimester screening for Prenatal disorders (Trisomy 21, 18 & 13) is essential to identify those women at sufficient risk for a congenital anomaly in the fetus to warrant further evaluation and followup. For Open neural tube defects, second trimester screening before 20 weeks is recommended. These are screening procedures which cannot discriminate all affected pregnancies from all unaffected pregnancies. Screening cutoffs are established by using MoM values that maximize the detection rate and minimize false positives.

*** End Of Report ***



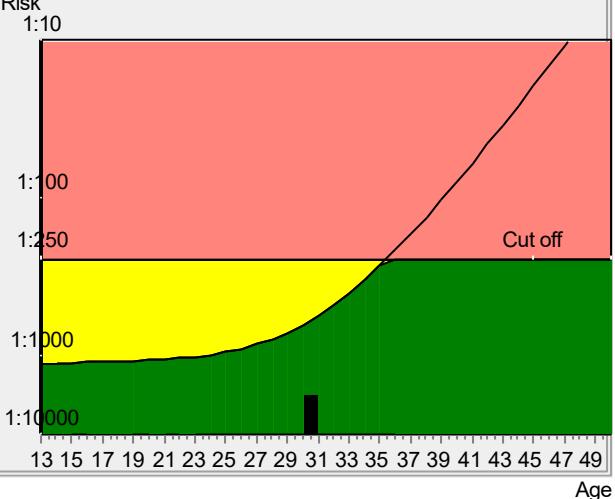
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DR. LAVANYA LAGISSETTY
MD BIOCHEMISTRY

Prisca 5.1.0.17

Date of report: 28/08/25

NA

Patient data			
Name	Mrs. AYSHA JULEKHA	Patient ID	0692508260016
Birthday	12/02/95	Sample ID	B3585048
Age at sample date	30.5	Sample Date	26/08/25
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	66	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	3.08 mIU/mL	0.82	Gestational age 12 + 2
fb-hCG	37.84 ng/ml	0.97	Method CRL Robinson
Risks at sampling date			
Age risk	1:601		Scan date 25/08/25
Biochemical T21 risk	1:2567		Crown rump length in mm 60
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.83
Trisomy 13/18 + NT	<1:10000		Nasal bone unknown
Sonographer NA			
Qualifications in measuring NT MD			
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.			
<p>After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
			
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

 below cut off Below Cut Off, but above Age Risk above cut off