

LABORATORY REPORT

Patient Details		Specimen Details		Prescription Details	
Name	: Mrs SHWETA M	Specimen	: Serum	Test	: Double Marker 1st Trimester Screening
Husband Name	:	Collected on	: 17-Sep-2025, 19:36	Clinician	: Dr. Buragadda Srinadh
Gender	: Female	Registration Date	: 17-Sep-2025, 19:35	Hospital	: DR B SRINADH
DOB	: 04-Dec-1990	Report Date	: 20-Sep-2025, 13:25	City	: Hyderabad
Weight	: 71 kg	Patient ID	:		
Ethnicity	: Asian	Lab ID	: 50906301056		

Ongoing Pregnancy		Sonography Details		Prior Risk Factors	
Last Menstrual Period	: 25-Jun-2025	Crown Rump Length (mm)	: 55mm (12 weeks 1 days)	Smoking	: None
Conception Method	: Spontaneous	Biparietal Length (mm)	: N/A	Diabetes	:
Number of Fetus	: 1	Nasal Bone	: Present	Previous Trisomy 21 pregnancies	:
EDD (By CRL)	: 31-Mar-2026	Scan Date	: 17-09-2025		
Maternal Age at Term	: 35 Years	GA at Collection	: 12 weeks		

Assistance Details	Method	Transfer Date	Egg Extraction Date	Age At Extraction
	N/A	N/A	N/A	N/A

Parameters Assessed

Out Of Range

Within Range

PAPP-A (By Auto Delfia)

Result : 8120.00 mU/L

Multiple of Median : 3.82



FBHCG (By Auto Delfia)

Result : 62.39 ng/mL

Multiple of Median : 1.5



NT (By Scan)

Result : 1.5 mm

Multiple of Median : 1.2



Risk Assessment – Aneuploidy

(These results were analyzed with LifeCycle software from PerkinElmer Life and Analytical Sciences))

Low Risk

Intermediate Risk

Increased Risk

Down Syndrome (T21) Biochemical Risk (BR) – 1:4545 Final Risk (FR) – <1:10000 Age Risk (AR) – 1:400 Risk Result – Low Risk	Edward Syndrome (T18) Final Risk (FR) – <1:10000 Age Risk (AR) – 1:3601 Risk Result – Low Risk	Patau Syndrome (T13) Final Risk (FR) – <1:10000 Age Risk (AR) – <1:10000 Risk Result – Low Risk
AR BR FR	AR FR	AR FR

Legend: Low Risk (Green), Intermediate Risk (Yellow), Increased Risk (Red)

Interpretation

Screening test shows low risk for T13/T18/T21.

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LABORATORY REPORT

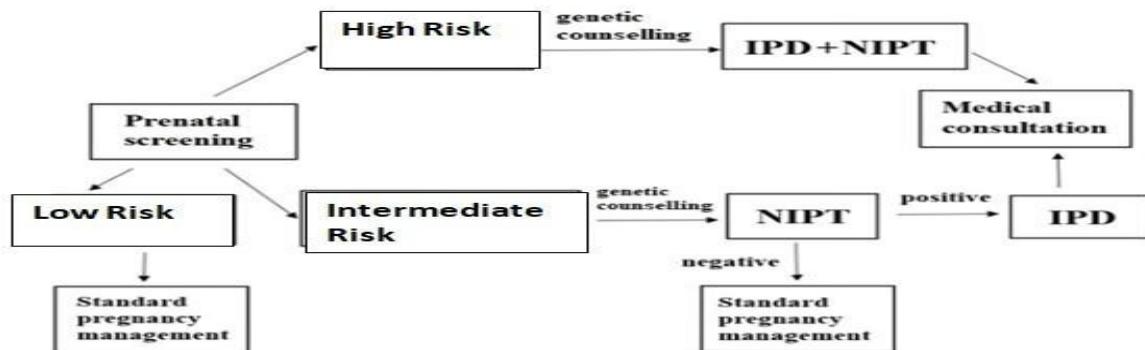
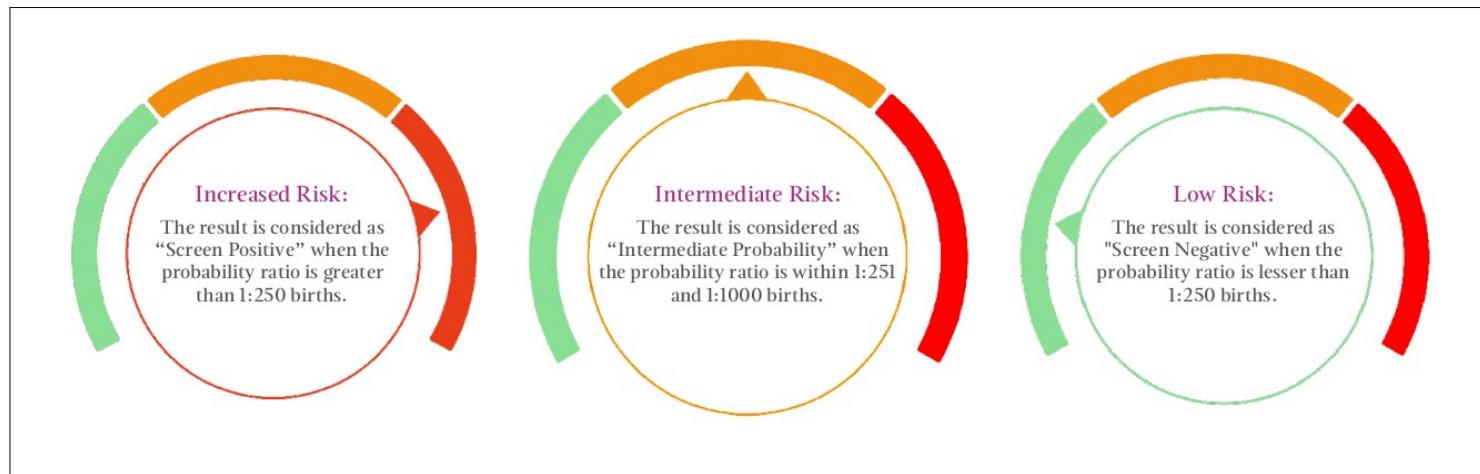
Caution

It must be clearly understood that this is a screening test, and therefore cannot be used to reach a definitive diagnosis. A low-risk result doesn't guarantee that your baby won't have one of these conditions. Likewise, an increased-risk result doesn't guarantee that your baby will be born with one of these conditions, and that further confirmatory tests must be performed in consultation with your healthcare provider.

FBHCG:Free beta human chorionic gonadotrophin | PAPPA:Pregnancy-associated plasma protein A | NT:Nuchal translucency | MoM:Multiple of median | NM:Not mentioned

Understanding Reported Final Risk

Prenatal screening gives a risk estimate after analyzing. The risk estimate is in the form of a ratio. For example, if the reported final risk is 1:1280, it means that of 1280 pregnancies with similar values, one baby is likely to be affected with the screened condition



NIPT, non-invasive prenatal testing; IPD, invasive prenatal diagnostics (Amniocentesis, CVS)

Disclaimer

1. This interpretation assumes that patient and specimen details are accurate and correct.
2. Ultrasound observations / measurements if not performed as per imaging guidelines may lead to erroneous risk assessments, and NSRL does not bear responsibility for results arising due to such errors



Dr. Harveen Bhusari

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MD Pathology

End of Report



Dr. Akash Shah

MD Pathology
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THE BEST START FOR YOUR BABY WITH NEWBORN SCREENING!

Dear Parents,

Congratulations on your journey to parenthood! As you prepare to welcome your baby, we want to introduce you to Newborn Screening (NBS)—a simple, essential test that can make a lifetime of difference.

What is Newborn Screening?

Newborn Screening is a painless, quick test performed after **48 hours of birth**. A few drops of blood from your baby's heel are screened for **over 50 metabolic, genetic, and hormonal disorders** that may not be visible at birth but can cause serious health complications if left undetected.

Why is it Important?

Even if your baby looks healthy, certain hidden conditions can affect their growth and development. Early detection through NBS helps in:

- Preventing severe complications like developmental delays, organ damage, or life-threatening conditions.
- Starting early treatment to manage or even cure certain disorders.
- Ensuring a healthier future with timely medical care.

Why Choose Neuberg Diagnostics for Newborn Screening?

NABL-Accredited Lab – Ensuring accurate, reliable, and high-quality testing.

- Expert Genetic Counselors
If a screening result is positive, our specialists provide personalized guidance and next steps.
- Globally Recommended Screening – Trusted by leading health organizations.
- One Test, Lifetime Benefits – Early detection leads to early intervention and better health outcomes.

**Give your baby the best start in life.
Ask your doctor about Newborn Screening today!**

For more details

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