JITM Diagnostics

Patient data			
Name	MRS. ANGURWATI	Patient ID	
Birthday	2/07/2000	Sample ID	2506017342/NOD
Age at sample date	24.9	Sample Date	11/06/2025
Gestational age	10 + 5		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	47	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	10 + 5
PAPP-A	2.9 mIU/ml	1.39	Method	CRL Robinson
fb-hCG	192 ng/ml	3.19	Scan date	11/06/2025

Risks at sampling date		Ultrasound data (cont.)	
Age risk	1:898	Crown rump length in mm	41.3
Biochemical T21 risk	1:602	Nuchal translucency MoM	0.77
Combined trisomy 21 risk	1:3227	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR. NEERU BHARDWAJ
		Qualifications in measuring NT	M.D

Risk

Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 3227 women with the same data, there is one woman with a trisomy 21 pregnancy and 3226 women with not affected pregnancies.
The free beta HCG level is high.
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!
The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

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