JITM Diagnostics

Patient data			
Name	MRS. NIRMALA (B)	Patient ID	
Birthday	30/10/1994	Sample ID	2505043401/NOD (B)
Age at sample date	30.6	Sample Date	21/05/2025
Gestational age	13 + 3		

Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21	no
Weight	50	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 3
PAPP-A	10.1 mIU/ml	0.76	Method	CRL Robinson
fb-hCG	79.1 ng/ml	0.85	Scan date	21/05/2025

Risks at sampling date		Ultrasound data	
		Crown rump length in mm	76.8
Age risk	1:620	Nuchal translucency MoM	0.75
Biochemical T21 risk	1:3039	Nasal bone	present
Combined trisomy 21 risk	<1:10000	Sonographer	DR. JYOTI GUPTA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MD

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.
The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.
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.

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Sign of Physician

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