

JITM Diagnostics

Patient data			
Name	MRS. SWETA	Patient ID	
Birthday	22/05/1988	Sample ID	2507002552/NOD
Age at sample date	37.1	Sample Date	2/07/2025
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	10.1 mIU/ml	1.69	
fb-hCG	56.1 ng/ml	1.37	
Risks at sampling date			
Age risk	1:172		Gestational age
Biochemical T21 risk	1:1421		13 + 2
Combined trisomy 21 risk	1:6495		Method
Trisomy 13/18 + NT	<1:10000		CRL Robinson
			Scan date
			1/07/2025
			Crown rump length in mm
			74.7
			Nuchal translucency MoM
			0.71
			Nasal bone
			present
			Sonographer
			DR. NEERJA CHOPRA
			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 6495 women with the same data, there is one woman with a trisomy 21 pregnancy and 6494 women with not affected pregnancies. 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