

JITM Diagnostics

Patient data			
Name	MRS. SHRUTI	Patient ID	
Birthday	24/12/1998	Sample ID	2504016826/NOD
Age at sample date	26.3	Sample Date	9/04/2025
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	69	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.67 mIU/ml	2.36	12 + 2
fb-hCG	20.1 ng/ml	0.46	Method
			CRL Robinson
Risks at sampling date			Scan date
Age risk		1:897	8/04/2025
Biochemical T21 risk		<1:10000	Crown rump length in mm
Combined trisomy 21 risk		<1:10000	59.5
Trisomy 13/18 + NT		<1:10000	Nuchal translucency MoM
			0.84
			Nasal bone
			present
			Sonographer
			DR. NEERJA CHOPRA
			Qualifications in measuring NT
			M.D
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician

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