

JITM Diagnostics

Patient data			
Name	MRS. ETI SHARMA		Patient ID
Birthday	4/07/1993		Sample ID
Age at sample date	31.7		Sample Date
Gestational age	11 + 5		25/03/2025
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.1 mIU/ml	0.69	11 + 5
fb-hCG	68.2 ng/ml	1.35	Method
Risks at sampling date			CRL Robinson
Age risk	1:491		Scan date
Biochemical T21 risk	1:668		25/03/2025
Combined trisomy 21 risk	1:1095		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		52
			Nuchal translucency MoM
			1.29
			Nasal bone
			present
			Sonographer
			DR. ANKIT KHANDELWAL MBBS DNB
			Qualifications in measuring NT
			MD
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 1095 women with the same data, there is one woman with a trisomy 21 pregnancy and 1094 women with not affected pregnancies.</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