

Patient data			
Name	MRS. SALONI JINDAL	Patient ID	2503063964/NOD
Birthday	30/05/1994	Sample ID	2503063964/NOD
Age at sample date	30.8	Sample Date	31/03/2025
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.6 mIU/ml	0.64	Gestational age 12 + 4
fb-hCG	53.7 ng/ml	1.20	Method CRL Robinson
Risks at sampling date			Scan date 31/03/2025
Age risk		1:579	Crown rump length in mm 64.1
Biochemical T21 risk		1:833	Nuchal translucency MoM 0.73
Combined trisomy 21 risk		1:4784	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. (MRS.) NEERJA CHOPRA
			Qualifications in measuring NT MD
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 4784 women with the same data, there is one woman with a trisomy 21 pregnancy and 4783 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