

Prisca

5.2.0.13

Date of report: 14/01/2025

Patient data			
Name	MRS. PRIYANKA	Patient ID	
Birthday	21/05/1992	Sample ID	2501019072/NOD
Age at sample date	32.6	Sample Date	13/01/2025
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.31 mIU/ml	0.49	12 + 6
fb-hCG	36.5 ng/ml	0.84	Method
			CRL Robinson
			Scan date
			13/01/2025
Risks at sampling date			Crown rump length in mm
Age risk	1:439		67
Biochemical T21 risk	1:720		Nuchal translucency MoM
Combined trisomy 21 risk	1:4205		0.76
Trisomy 13/18 + NT	<1:10000		Nasal bone
			present
			Sonographer
			DR. (MRS.) NEERJA CHOPRA
			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 4205 women with the same data, there is one woman with a trisomy 21 pregnancy and 4204 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