

Patient data			
Name	MRS. SWETA	Patient ID	
Birthdate	12/11/1995	Sample ID	2409041483/NOD
Age at sample date	28.9	Sample Date	19/09/2024
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.25 mIU/ml	1.14	Gestational age 12 + 4
fb-hCG	20.6 ng/ml	0.47	Method CRL Robinson
Risks at sampling date			Scan date 19/09/2024
Age risk		1:736	Crown rump length in mm 64.5
Biochemical T21 risk		<1:10000	Nuchal translucency MoM 0.73
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	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 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

