

Patient data			
Name	MRS. MAN CHANDA	Patient ID	
Birth day	26/09/1993	Sample ID	2409030460/NOD
Age at sample date	31.0	Sample Date	14/09/2024
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	60.5	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.35 mIU/ml	0.76	Gestational age 13 + 3
fb-hCG	56.3 ng/ml	1.40	Method CRL Robinson
Risks at sampling date			Scan date 14/09/2024
Age risk		1:586	Crown rump length in mm 75.4
Biochemical T21 risk		1:927	Nuchal translucency MoM 0.54
Combined trisomy 21 risk		1:5181	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. NEERU BHARDWAJ
			Qualifications in measuring NT M.D
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 5181 women with the same data, there is one woman with a trisomy 21 pregnancy and 5180 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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician

