

Patient data			
Name	MRS. SANDHYA GHARANA		Patient ID
Birthday	14/01/1989		Sample ID
Age at sample date	36.0		Sample Date
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	67	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			no
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.61 mIU/ml	0.78	12 + 3
fb-hCG	41.6 ng/ml	0.95	Method
			CRL Robinson
			Scan date
			22/01/2025
Risks at sampling date			Crown rump length in mm
			61.5
Age risk	1:215		Nuchal translucency MoM
Biochemical T21 risk	1:850		1.13
Combined trisomy 21 risk	1:2441		Nasal bone
Trisomy 13/18 + NT	<1:10000		present
			Sonographer
			DR. SHAILEN SINGHA
			Qualifications in measuring NT
			M.D
			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 2441 women with the same data, there is one woman with a trisomy 21 pregnancy and 2440 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