

Patient data			
Name	Mrs.ANANYA KUMAR	Patient ID	4053629
Birthday	30/12/00	Sample ID	4053629
Age at delivery	22.5	Sample Date	23/12/22
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	unknown
Weight	44.5	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	10 mIU/ml	2.01	Gestational age 11 + 6
fb-hCG	32.8 ng/ml	0.84	Method CRL Robinson
Risks at term			Scan date 18/12/22
Age risk		1:1481	Crown rump length in mm 53.9
Biochemical T21 risk		<1:10000	Nuchal translucency MoM 0.77
Combined trisomy 21 risk		<1:10000	Nasal bone unknown
Trisomy 13/18 + NT		<1:10000	Sonographer Dr.Balajeet Kaur
			Qualifications in measuring NT FMF
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

