

Patient data			
Name	MRS. SHOIBA ANSARI	Patient ID	
Birthday	1/01/1994	Sample ID	2412021881/NOD
Age at sample date	30.9	Sample Date	12/12/2024
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	47	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies			no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	10.3 mIU/ml	1.79	Gestational age 12 + 3
fb-hCG	105 ng/ml	2.19	Method CRL Robinson
Risks at sampling date			Scan date 10/12/2024
Age risk		1:573	Crown rump length in mm 62.4
Biochemical T21 risk		1:1615	Nuchal translucency MoM 0.88
Combined trisomy 21 risk		1:7284	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. SANDEEP GOEL
			Qualifications in measuring NT M.D
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 7284 women with the same data, there is one woman with a trisomy 21 pregnancy and 7283 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

