

Patient data			
Name	MRS. VARSHA		Patient ID
Birthday	26/07/92	Sample ID	2504033820/NOD
Age at sample date	32.7	Sample Date	16/04/25
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.75 mIU/ml	0.68	Gestational age 13 + 1
fb-hCG	57.3 ng/ml	1.35	Method CRL Robinson
Risks at sampling date			Scan date 16/04/25
Age risk	1:438		Crown rump length in mm 72.9
Biochemical T21 risk	1:578		Nuchal translucency MoM 0.77
Combined trisomy 21 risk	1:3310		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer DR. NEERU BHARDWAJ
			Qualifications in measuring NT M.D
Risk			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 3310 women with the same data, there is one woman with a trisomy 21 pregnancy and 3309 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