

Patient data			
Name	MRS. SANDHYA	Patient ID	
Birthday	4/08/2000	Sample ID	2407010873/NOD
Age at sample date	23.9	Sample Date	6/07/2024
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.36 mIU/ml	1.03	11 + 4
fb-hCG	24.6 ng/ml	0.55	Method
			CRL Robinson
Risks at sampling date			Scan date
Age risk	1:1014		27/06/2024
Biochemical T21 risk	<1:10000		Crown rump length in mm
Combined trisomy 21 risk	<1:10000		51
Trisomy 13/18 + NT	<1:10000		Nuchal translucency MoM
			1.09
			Nasal bone
			present
			Sonographer
			DR. PREETHAM
			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 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><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

below cut off	Below Cut Off, but above Age Risk	above cut off
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