

Patient data				
Name		Mrs. SHAMALA	Patient ID	0842508200019
Birthday		21-08-1998	Sample ID	B2998721
Age at sample date		27.0	Sample Date	20-08-2025
Gestational age		11 + 4		
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	38	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	
PAPP-A	2.22 mIU/mL	0.51	10 + 6	
fb-hCG	48.54 ng/mL	0.79	Method	
Risks at sampling date			CRL Robinson	
Age risk			15-08-2025	
Biochemical T21 risk			Crown rump length in mm	
Combined trisomy 21 risk			42.5	
Trisomy 13/18 + NT			Nuchal translucency MoM	
			1.01	
			Nasal bone	
			unknown	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			NA	
Risk			Trisomy 21	
			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 7101 women with the same data, there is one woman with a trisomy 21 pregnancy and 7100 women with not affected pregnancies. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Trisomy 13/18 + NT				
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.				

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