

Patient data			
Name	Mrs. B PALLAVI	Patient ID	0012506250240
Birthday	22-10-2001	Sample ID	A2299349
Age at sample date	23.7	Sample Date	25-06-2025
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	56	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.9 mIU/mL	0.41	12 + 3
fb-hCG	38.41 ng/mL	0.94	Method
Risks at sampling date			CRL Robinson
Age risk	1:1017		Scan date
Biochemical T21 risk	1:794		23-06-2025
Combined trisomy 21 risk	1:4949		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		60.9
			Nuchal translucency MoM
			0.63
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 4949 women with the same data, there is one woman with a trisomy 21 pregnancy and 4948 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