

Patient data			
Name		Mrs. HARINI REDDY	Patient ID
Birthday		12-02-1998	0212506120010
Age at sample date		27.3	Sample ID
Gestational age		12 + 6	B3000763
			Sample Date
			12-06-2025
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.84 mIU/mL	0.39	12 + 6
fb-hCG	37.07 ng/mL	0.95	Method
Risks at sampling date			CRL Robinson
Age risk		1:849	Scan date
Biochemical T21 risk		1:566	12-06-2025
Combined trisomy 21 risk		1:3540	Crown rump length in mm
Trisomy 13/18 + NT		<1:10000	68
			Nuchal translucency MoM
			0.81
			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 3540 women with the same data, there is one woman with a trisomy 21 pregnancy and 3539 women with not affected pregnancies. The PAPP-A level is low. 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