

Patient data			
Name	Mrs. MEENAKSHI DEVI.	Patient ID	0852510120182.
Birthday	24-03-1994	Sample ID	B3529010.
Age at sample date	31.6	Sample Date	12-10-2025
Gestational age	11 + 4		
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	
PAPP-A	1.84 mIU/mL	0.65	Gestational age 11 + 3
fb-hCG	48.02 ng/mL	0.91	Method CRL Robinson
Risks at sampling date			Scan date 11-10-2025
Age risk	1:501		Crown rump length in mm 48.1
Biochemical T21 risk	1:1422		Nuchal translucency MoM 0.76
Combined trisomy 21 risk	1:7808		Nasal bone present
Trisomy 13/18 + NT	<1:10000		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 7808 women with the same data, there is one woman with a trisomy 21 pregnancy and 7807 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