

Patient data			
Name	Mrs. MAHESHWARI	Patient ID	0012502010358
Birthday	09-03-2000	Sample ID	A111011C
Age at sample date	24.9	Sample Date	01-02-2025
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	47	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.93 mIU/mL	0.70	Gestational age 11 + 6
fb-hCG	63.49 ng/mL	1.24	Method CRL Robinson
Risks at sampling date			Scan date 31-01-2025
Age risk	1:947		Crown rump length in mm 53.6
Biochemical T21 risk	1:1591		Nuchal translucency MoM 0.77
Combined trisomy 21 risk	1:8966		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer NA
			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 8966 women with the same data, there is one woman with a trisomy 21 pregnancy and 8965 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