

Prisca 5.1.0.17  
Date of report: 18-10-2025

NA

Patient data			
Name	Mrs. POOJA MANI	Patient ID	0012510170356
Birthday	10-12-1990	Sample ID	B3287472
Age at sample date	34.9	Sample Date	17-10-2025
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	67.7	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.53 mIU/mL	0.33	13 + 1
fb-hCG	34.9 ng/mL	1.05	Method
			CRL Robinson
			Scan date
			16-10-2025
Risks at sampling date			Trisomy 21
Age risk		1:289	Crown rump length in mm
Biochemical T21 risk		1:97	72
Combined trisomy 21 risk		1:649	Nuchal translucency MoM
Trisomy 13/18 + NT		1:9571	0.72
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			NA
			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 649 women with the same data, there is one woman with a trisomy 21 pregnancy and 648 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
Trisomy 13/18 + NT			
<p><b>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:9571, which represents a low risk.</b></p>			

Sign of Physician