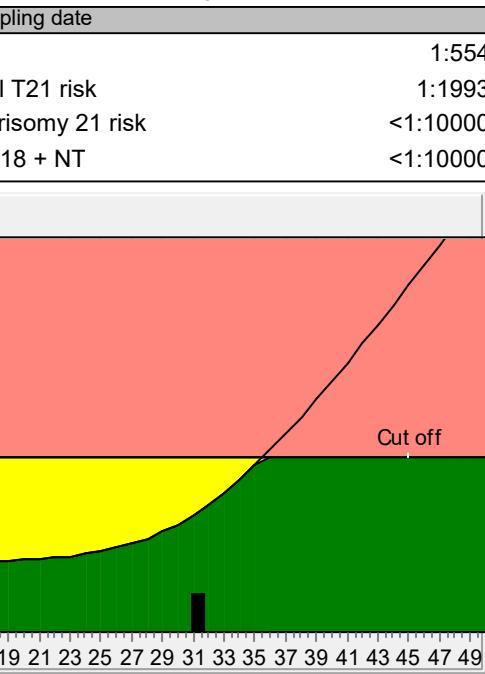


Prisca 5.1.0.17
Date of report: 11-03-2025

DR

S MANJULA MBBS

Patient data		Test results		Report date				
Name	Mrs. POOJA MORE	Patient ID		0662503110131				
Birthday	15-11-1993	Sample ID		B2566338				
Age at sample date	31.3	Sample Date		11-03-2025				
Gestational age	13 + 2							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	71	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data			Ultrasound data					
Parameter	Value	Corr. MoM	Gestational age	13 + 1				
PAPP-A	4.3 mIU/mL	0.99	Method		CRL Robinson			
fb-hCG	41.31 ng/mL	1.26	Scan date	10-03-2025	71			
Risks at sampling date			Crown rump length in mm					
Age risk		1:554	Nuchal translucency MoM		0.68			
Biochemical T21 risk		1:1993	Nasal bone	present				
Combined trisomy 21 risk		<1:10000	Sonographer	DR S MANJULA MBBS				
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	DMCH				
Risk			Trisomy 21					
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.					
1:100			After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.					
1:250			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.					
1:1000			Please note that risk calculations are statistical approaches and have no diagnostic value!					
1:10000			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 held 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.								

Sign of Physician

below cut off

 Below Cut Off, but above Age Risk

above cut off