

Prisca 5.1.0.17
Date of report: 03-08-2025

N A

Patient data			
Name		Mrs. PRANEETHA	
Patient ID		0212507310016	
Birthday		07-12-1997	
Sample ID		A2291466	
Age at sample date		27.6	
Sample Date		31-07-2025	
Gestational age		11 + 6	
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.86 mIU/mL	0.59	11 + 6
fb-hCG	44.74 ng/mL	0.91	Method
Risks at sampling date			CRL Robinson
Age risk			31-07-2025
Biochemical T21 risk			Crown rump length in mm
Combined trisomy 21 risk			55
Trisomy 13/18 + NT			Nuchal translucency MoM
			0.89
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 9293 women with the same data, there is one woman with a trisomy 21 pregnancy and 9292 women with not affected pregnancies.	
1:250		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!	
1:1000		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
1:10000		The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
Age			
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