

Prisca 5.1.0.17
Date of report: 26-11-2025

Patient data			
Name Mrs. PALLAVI ROHIT RAMTEKE		Patient ID 0372511250100	
Birthday 26-01-1996		Sample ID A1559982	
Age at sample date 29.8		Sample Date 25-11-2025	
Gestational age 13 + 3			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 48	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 0
PAPP-A	4.26 mIU/mL	0.59	Method CRL Robinson
fb-hCG	33.52 ng/mL	0.94	Scan date 22-11-2025
Risks at sampling date			Crown rump length in mm 69.7
Age risk 1:680			Nuchal translucency MoM 0.63
Biochemical T21 risk 1:1425			Nasal bone present
Combined trisomy 21 risk 1:8058			Sonographer NA
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT Sonographer
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 8058 women with the same data, there is one woman with a trisomy 21 pregnancy and 8057 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.			

Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off