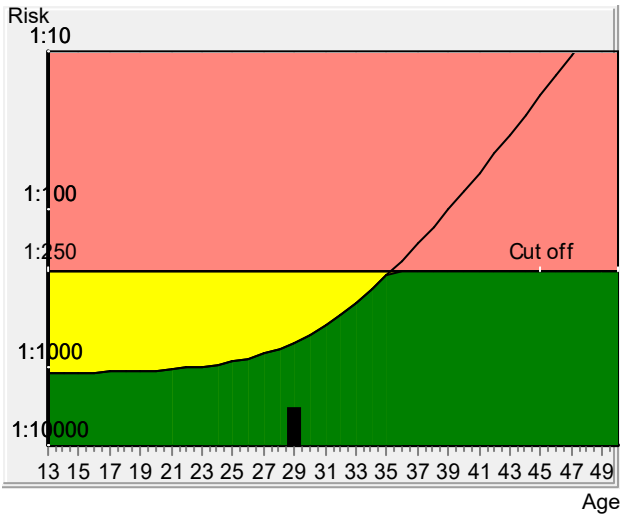


Prisca 5.1.0.17
Date of report: 20-11-2025

N A

Patient data						
Name		Mrs. PRIYANKA YADAV		Patient ID	038251118011C	
Birthday		14-12-1996		Sample ID	B3963642	
Age at sample date		28.9		Sample Date	18-11-2025	
Gestational age		11 + 5				
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown	
Weight	40	diabetes	no			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age			11 + 5
PAPP-A	2.06 mIU/mL	0.46	Method			CRL Robinson
fb-hCG	46.27 ng/mL	0.79	Scan date			18-11-2025
Risks at sampling date			Crown rump length in mm			52
Age risk		1:707	Nuchal translucency MoM			0.79
Biochemical T21 risk		1:1117	Nasal bone			present
Combined trisomy 21 risk		1:6574	Sonographer			N A
Trisomy 13/18 + NT		<1:10000	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 6574 women with the same data, there is one woman with a trisomy 21 pregnancy and 6573 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