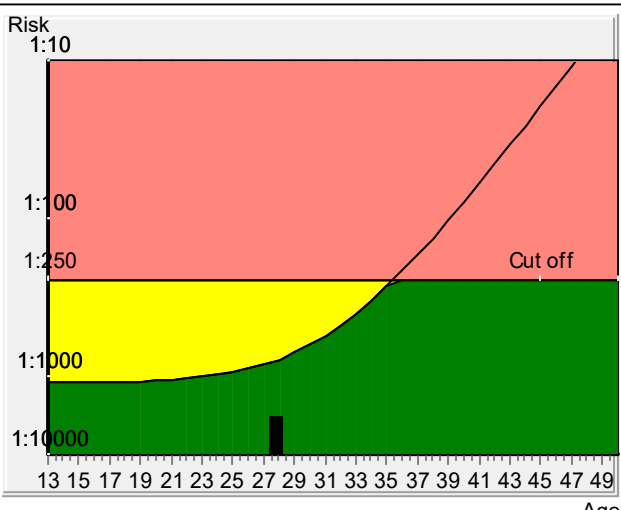


Prisca 5.1.0.17
Date of report: 25-10-2025

N A

Patient data						
Name		Mrs. T PRIYA		Patient ID	0012510240114	
Birthday		16-01-1998		Sample ID	B3101157	
Age at sample date		27.8		Sample Date	24-10-2025	
Gestational age		12 + 2				
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown	
Weight	44	diabetes	no			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age			12 + 1
PAPP-A	1.94 mIU/mL	0.38	Method			CRL Robinson
fb-hCG	40.09 ng/mL	0.81	Scan date			23-10-2025
Risks at sampling date			Crown rump length in mm			58
Age risk		1:804	Nuchal translucency MoM			0.46
Biochemical T21 risk		1:682	Nasal bone			present
Combined trisomy 21 risk		1:4258	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 4258 women with the same data, there is one woman with a trisomy 21 pregnancy and 4257 women with not affected pregnancies. The PAPP-A level is low. 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