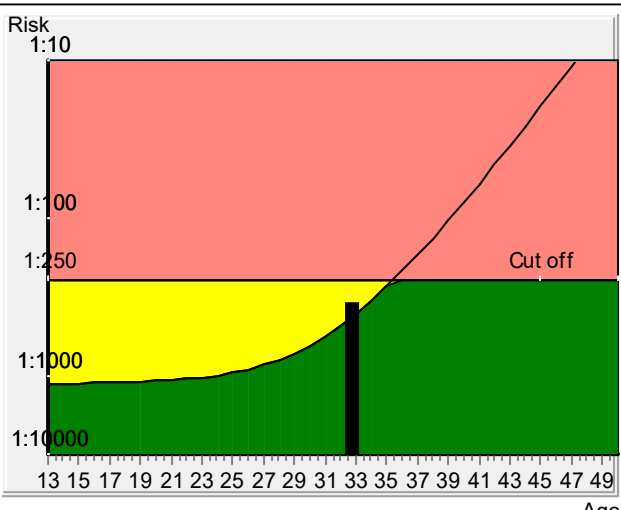


Prisca 5.1.0.17
Date of report: 21-07-2025

NA

Patient data				
Name		Mrs. MAMATHA	Patient ID 0012507210213	
Birthday		10-10-1992	Sample ID B3108629	
Age at sample date		32.8	Sample Date 21-07-2025	
Gestational age		12 + 3		
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 + 2	
PAPP-A	1.08 mIU/mL	0.20	Method CRL Robinson	
fb-hCG	40.15 ng/mL	0.84	Scan date 20-07-2025	
Risks at sampling date			Crown rump length in mm 59.5	
Age risk		1:422	Nuchal translucency MoM 0.71	
Biochemical T21 risk		>1:50	Nasal bone unknown	
Combined trisomy 21 risk		1:340	Sonographer NA	
Trisomy 13/18 + NT		1:740	Qualifications in measuring NT NA	
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 340 women with the same data, there is one woman with a trisomy 21 pregnancy and 339 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 1:740, which represents a low risk.				

Sign of Physician