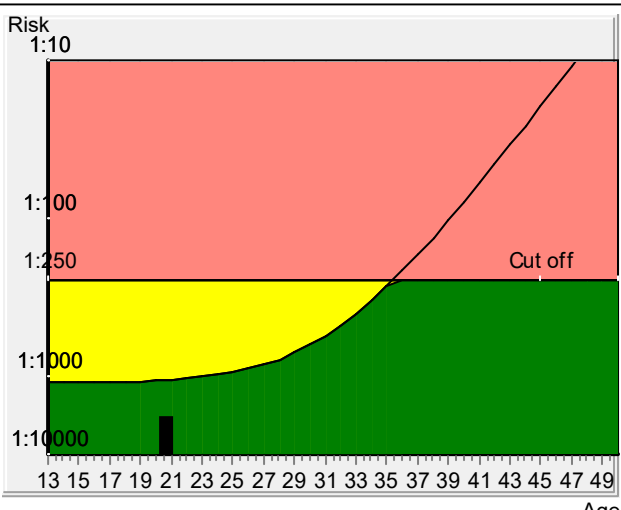


Prisca 5.1.0.17
Date of report: 03-02-2025

N A

Patient data			
Name		Mrs. MAHESHWARI W/O SHASHI	
Patient ID		0312502010024	
Birthday		21-06-2004	
Sample ID		A0936995	
Age at sample date		20.6	
Sample Date		01-02-2025	
Gestational age		12 + 2	
Correction factors			
Fetuses	1	IVF	no
Weight	41	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.91 mIU/mL	0.52	12 + 2
fb-hCG	42.36 ng/mL	0.83	Method
Risks at sampling date			CRL Robinson
Age risk	1:1071		Scan date
Biochemical T21 risk	1:2121		01-02-2025
Combined trisomy 21 risk	1:2808		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		60
			Nuchal translucency MoM
			1.36
			Nasal bone
			unknown
			Sonographer
			N A
			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 2808 women with the same data, there is one woman with a trisomy 21 pregnancy and 2807 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