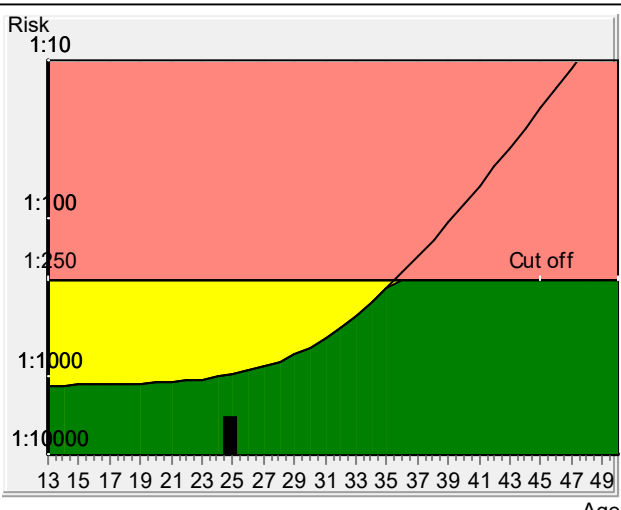


Prisca 5.1.0.17
Date of report: 14-10-2024

N A

Patient data			
Name		Mrs. MASARATH BEGUM	
Patient ID		0012410140229	
Birthday		01-01-2000	
Sample ID		A1302737	
Age at sample date		24.8	
Sample Date		14-10-2024	
Gestational age		13 + 2	
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	8.8 mIU/mL	1.50	12 + 6
fb-hCG	43.1 ng/mL	1.22	Method
Risks at sampling date			CRL Robinson
Age risk			1:995
Biochemical T21 risk			1:8879
Combined trisomy 21 risk			1:2227
Trisomy 13/18 + NT			<1:10000
Scan date			11-10-2024
Crown rump length in mm			68
Nuchal translucency MoM			1.74
Nasal bone			present
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 2227 women with the same data, there is one woman with a trisomy 21 pregnancy and 2226 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