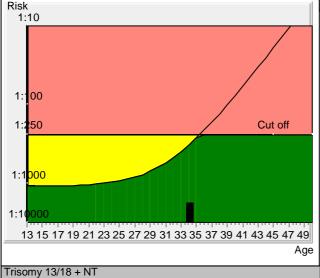
D-87, SEC-2 NOIDA

Prisca 5.2.0.13

Date of report: 8/12/2024

Patient data						
Name	MRS. RAMEEZA RAFI			Patient ID		
Birthday	27/09/1990			Sample ID		2412013268/NOD
Age at sample date	34.2			Sample Date		7/12/2024
Gestational age	12 + 2					
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	55.3	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound data		
Parameter	Value Corr. MoN		Gestational age		12 + 2	
PAPP-A	1.2 mIU/ml		0.30	Method		CRL Robinson
fb-hCG	20.3 ng/ml		0.43	Scan date		7/12/2024
Risks at sampling date				Crown rump length in mm		60.4
Age risk	1:319			Nuchal translucency MoM		1.08
Biochemical T21 risk	risk 1:494			Nasal bone		present
Combined trisomy 21 risk 1:1875			Sonographer		DR. SHUBHAM SAHAI	
Trisomy 13/18 + NT 1:636			Qualifications in measuring NT ME			
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 1875 women with the same data, there is one woman with a trisomy 21 pregnancy and 1874 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!

Sign of Physician

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:636, which represents a low risk.