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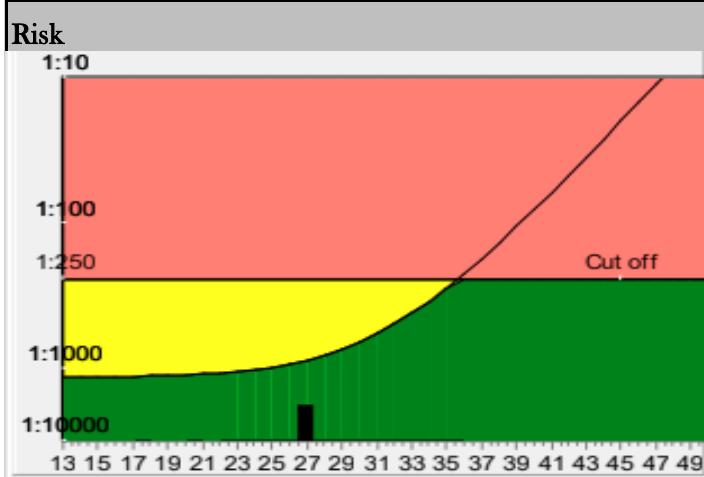
Date of Report 6/2/2024
PRISCA 5.2.0.13

Patient Data	
Name	PRIYANKA
Birthdate	26/03/1997
Age at sample	26.9
Gestational age	13+5

Correction factors	
Fetuses	1 IVF
Weight in kg	55.9
Smoker	NO
Diabetes	NO
Origin	Asian
Previous trisomy 21	unknown
Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+4
PAPP-A	5.90 mIU/ml	0.66	Method	CRL (<>Robinson)
fb-hCG	56.5 ng/ml	2.12	Scan date	4/2/2024

Risks at sampling date		Down's Syndrome Risk (Trisomy 21 Screening)	
Age Risk	1:902	Crown rump length in mm	62.2
Biochemical T21 risk	1:358	Nuchal translucency MoM	0.69
Combined trisomy 21 risk	1:2207	Nasal bone	Present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR.ABHISHEKH
		Qualifications in measuring NT	MBBS



Trisomy 13/18 + NT
The calculated risk for Trisomy 13/18 (with NT) is < 1:10000, which indicates a low risk

The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 2207 women with the same data, there is one woman with a trisomy 21 pregnancy and 2206 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 the risk calculations are statistical approaches and have no diagnostic value! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off
 Risk above Age Risk
 Risk below Age risk