

Date of Report 19-10-2024
PRISCA 5.2.0.13

Patient Data			
Name	MRS. GUNJAN W/O MANJEET	Patient ID	12410180159
Birthday	28-02-2000	Sample ID	11966742
Age at Sample date	24.6	Sample Date	18-10-2024
Gestational age	13+3		

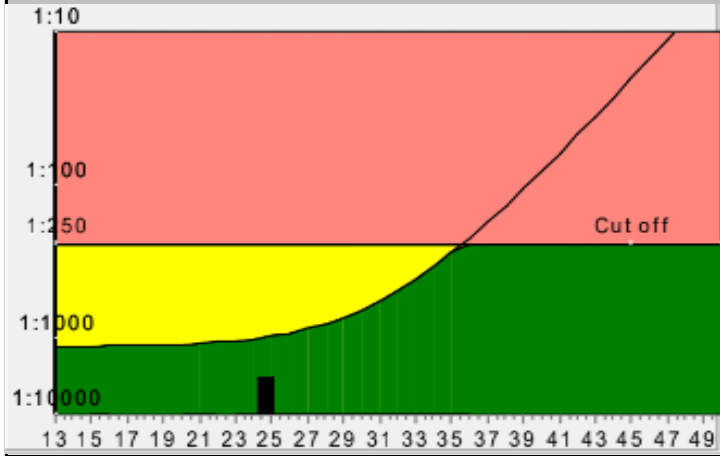
Correction factors			
Fetuses	1 IVF	unknown	Previous trisomy 21 unknown
Weight in kg	56.8	Diabetes NO	Pregnancies unknown
Smoker	NO	Origin Asian	

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom	
PAPP-A	6.9 mIU/ml	0.86	Method CRL (<>Robinson)
fb-hCG	78.5 ng/ml	2.69	Scan date 18-10-2024

Risks at sampling date		
Age Risk	1:1006	Crown rump length in mm 72.1
Biochemical T21 risk	1:403	Nuchal translucency MoM 0.84
Combined trisomy 21 risk	1:2296	Nasal bone PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer DR. AMENDA
		Qualifications in measuring NT MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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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 2296 women with the same data, there is one woman with a trisomy 21 pregnancy and 2295 women with not affected pregnancies. The free beta HCG level is high.

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).

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

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