Date of Report	04-12-2020
PRISCA	5.0.2.37

Patient Data

Name	MRS JYOTSANA	Patient ID	012011300135
Birthday	15-11-1996	Sample ID	10787779
Age at delivery	24	Sample Date	30/11/2020
Gestational age	13+4		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	57.3	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	5.3	mIU/ml	0.78
fb-hCG	15.3	ng/ml	0.36

Risks at sampling date

Age Risk	1:1033
Biochemical T21 risk	<1:10000
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	13+4
Method	CRL(<>Robinson
Scan date	30-11-2020
Crown rump length in mm	74.6
Nuchal translucency MOM	0.88
Nasal bone	Present
Sonographer	DR.DEEPAK BHATIA
Qualification in measuring NT	MBBS,MD
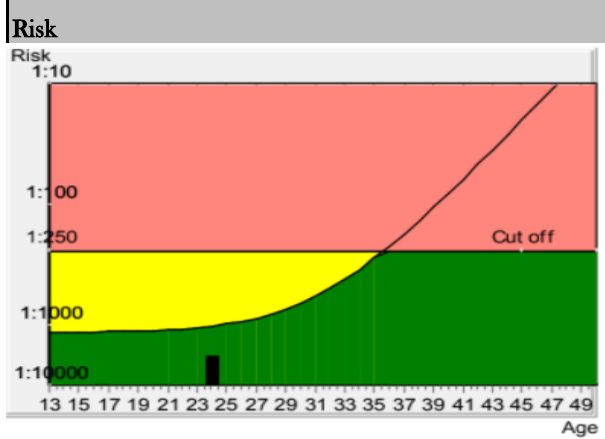
Risk

Down's Syndrome Risk (Trisomy 21 Screening)

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 aapproaches and have no diagnostic value!

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Trisomy 13/18 + NT

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

🟥 Risk Above Cut Off	🟨 Risk above Age Risk	🟩 Risk below Age risk