Date of Report	11-12-2020
PRISCA	5.0.2.37

Patient Data

Name	MRS MANISHA	Patient ID	012012100009
Birthday	25-11-1986	Sample ID	10807700
Age at delivery	34	Sample Date	10/12/2020
Gestational age	13+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	55	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.2 mIU/ml	0.76
fb-hCG	41.6 ng/ml	0.96

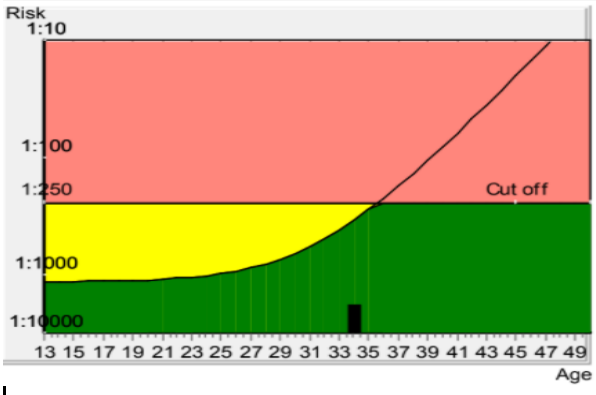
Ultrasound Data

Gestational age	13+2
Method	CRL(<>Robinson
Scan date	09-12-2020
Crown rump length in mm	56.8
Nuchal translucency MOM	0.33
Nasal bone	Present
Sonographer	DR.APARNA
Qualification in measuring NT	MD

Risks at sampling date

Age Risk	1:344
Biochemical T21 risk	1:1280
Combined Trisomy 21 Risk	1:6788
Trisomy 13/18 + NT	<1:10000

Risk



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low 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 6788 women with the same data, there is one woman with a trisomy 21 pregnancy and 6787 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

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