Date of Report	20-04-2021
PRISCA	5.0.2.37

Patient Data

Name	**MRS. MAMTA**	Patient ID	012104160186
Birthday	16-07-1986	Sample ID	10918058
Age at delivery	34.8	Sample Date	16/04/2021
Gestational age	13+4		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	54	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.3 mIU/ml	0.73
fb-hCG	48.3 ng/ml	1.13

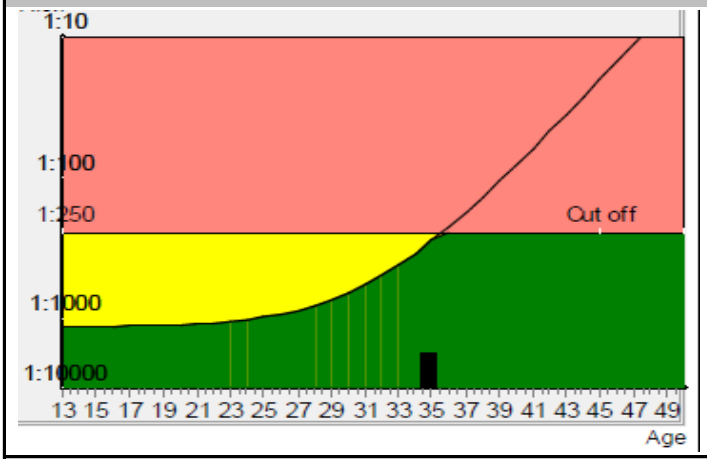
Ultrasound Data

Gestational age	13+0
Method	CRL(<>Robinson
Scan date	15-04-2021
Crown rump length in mm	66.2
Nuchal translucency MOM	0.61
Nasal bone	Present
Sonographer	Dr Namita Sharma
Qualification in measuring NT	C/R

Risks at sampling date

Age Risk	1:297
Biochemical T21 risk	1:691
Combined Trisomy 21 Risk	1:3796
Trisomy 13/18 + NT	<1:10000

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 3796 women with the same data, there is one woman with a trisomy 21 pregnancy and 3795 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 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

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