Date of Report	17/10/19
PRISCA	5.0.2.37

Patient Data

Name	Mrs Sarita w/o Mahesh	Patient ID	011910150234
Birthday	8/8/1993	Sample ID	10553350
Age at delivery	26.2	Sample Date	15/10/19
Gestational age	12+1		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	54.3	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	2.18 mIU/ml	0.5
fb-hCG	24.9 ng/ml	0.49

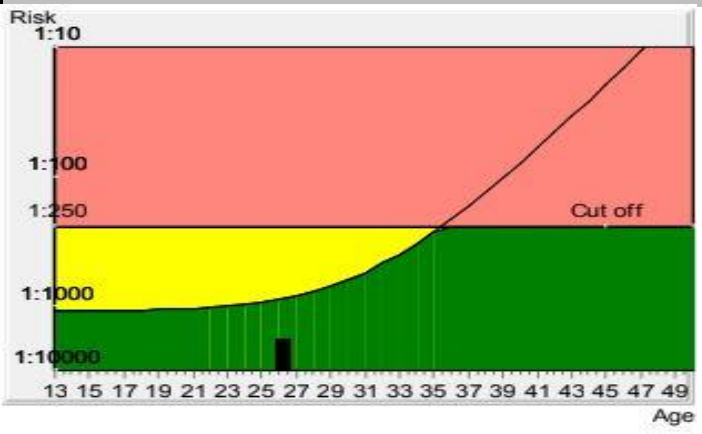
Ultrasound Data

Gestational age	12+0
Method	CRL (<>Robinson)
Scan Date	15/10/19
Crown Rump Length (mm)	52.7
Nuchal translucency MoM	1.14
Nasal Bone	present
Sonographer	DR. RAJESH YADAV
Qualification in measuring NT	CON. RADIOLOGIST

Risks at sampling date

Age Risk	1:893
Biochemical Trisomy 21 Risk	1:4487
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Risk



Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates 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.
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 nuchal translucency) is <1:10000, which represents a low risk.

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