Date of Report	1/1/2020
PRISCA	5.0.2.37

Patient Data

Name	Mrs Neha Singh	Patient ID	011912310088
Birthday	29/12/1987	Sample ID	10496968
Age at delivery	32	Sample Date	31/12/19
Gestational age	12+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	68.4	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	2.85 mIU/ml	0.77
fb-hCG	51.4 ng/ml	1.14

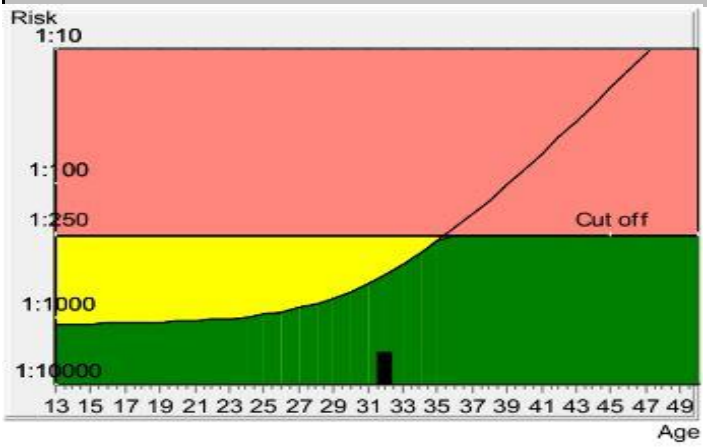
Ultrasound Data

Gestational age	12+0
Method	CRL (<>Robinson)
Scan Date	28/12/2019
Crown Rump Length (mm)	54.1
Nuchal translucency MoM	0.63
Nasal Bone	present
Sonographer	DR. HARMEET KAUR
Qualification in measuring NT	MD

Risks at sampling date

Age Risk	1:482
Biochemical Trisomy 21 Risk	1:1247
Combined Trisomy 21 Risk	1:6764
Trisomy 13/18 + NT	<1:10000

Risk



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low 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 6764 women with the same data, there is one woman with a trisomy 21 pregnancy and 6763 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