Date of Report	28/02/2020
PRISCA	5.0.2.37

Patient Data

Name	Mrs Preeti Taneja	Patient ID	012002270103
Birthday	1/1/1988	Sample ID	10642390
Age at delivery	32.2	Sample Date	27/02/2020
Gestational age	13+0		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	73.3	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.18 mIU/ml	0.76
fb-hCG	63.5 ng/ml	1.53

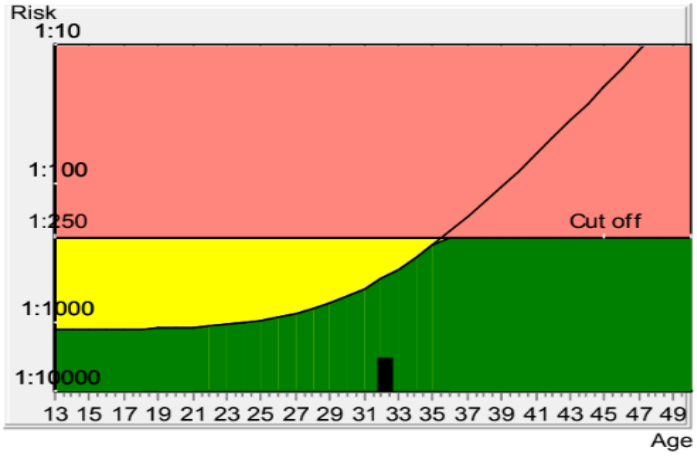
Risks at sampling date

Age Risk	1:480
Biochemical Trisomy 21 Risk	1:603
Combined Trisomy 21 Risk	1:3415
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+6
Method	CRL (<>Robinson)
Scan Date	26/02/2020
Crown Rump Length (mm)	65.7
Nuchal translucency MoM	0.79
Nasal Bone	present
Sonographer	DR. ANKIT BHARGAVA
Qualification in measuring NT	MD

Risk



Trisomy 13/18 + NT

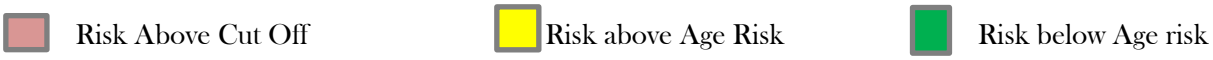
The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.

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 3415 women with the same data, there is one woman with a trisomy 21 pregnancy and 3414 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