Date of Report	10-10-2019
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Jyoti**	Patient ID	011909290149
Birthday	25-12-1991	Sample ID	10550091
Age at delivery	27.8	Sample Date	29/09/19
Gestational age	12+5		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	61	Diabetes	no	Pregnancies	0
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	4.5 mIU/ml	0.95
fb-hCG	21.3 ng/ml	0.47

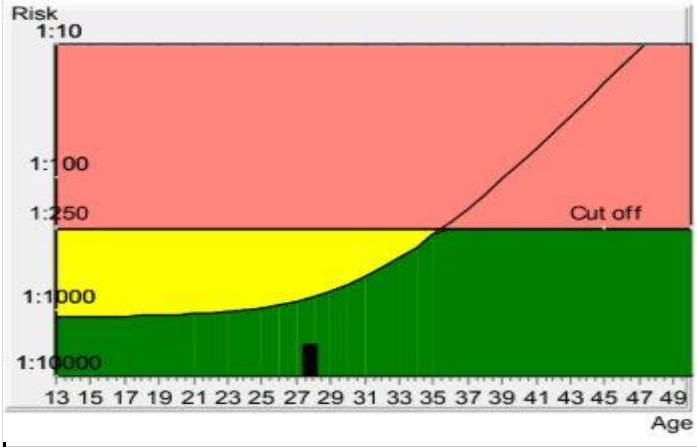
Risks at sampling date

Age Risk	1:817
Biochemical Trisomy 21 Risk	<1:10000
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	13+0
Method	CRL (<>Robinson)
Scan Date	02-10-2019
Crown Rump Length (mm)	65.3
Nuchal translucency MoM	0.84
Nasal Bone	present
Sonographer	DR. VARUN SHARMA
Qualification in measuring NT	

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)

TThe 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

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