Date of Report	24/09/19
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Mamta**	Patient ID	011909220170
Birthday	23/11/1983	Sample ID	10543473
Age at delivery	35.8	Sample Date	22/09/19
Gestational age	12+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	58	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	4.3 mIU/ml	0.95
fb-hCG	19.2 ng/ml	0.40

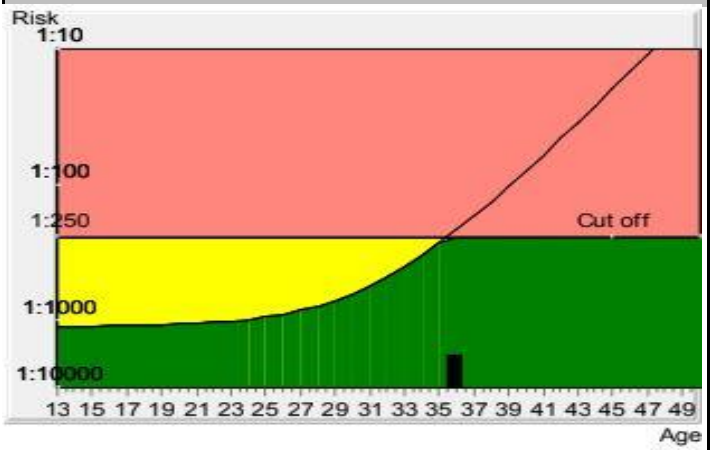
Risks at sampling date

Age Risk	1:225
Biochemical Trisomy 21 Risk	1:8220
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+2
Method	CRL (<>Robinson)
Scan Date	22/09/19
Crown Rump Length (mm)	53
Nuchal translucency MoM	0.71
Nasal Bone	present
Sonographer	DR. SHRUTI SANGWAN
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 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