Date of Report	25/04/19
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Payal**	Patient ID	011904230333
Birthday	9/10/1990	Sample ID	10482895
Age at delivery	28.5	Sample Date	24/04/19
Gestational age	12+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	62	Diabetes	no	Pregnancies	unknown
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.82 mIU/ml	0.92
fb-hCG	64.9 ng/ml	1.39

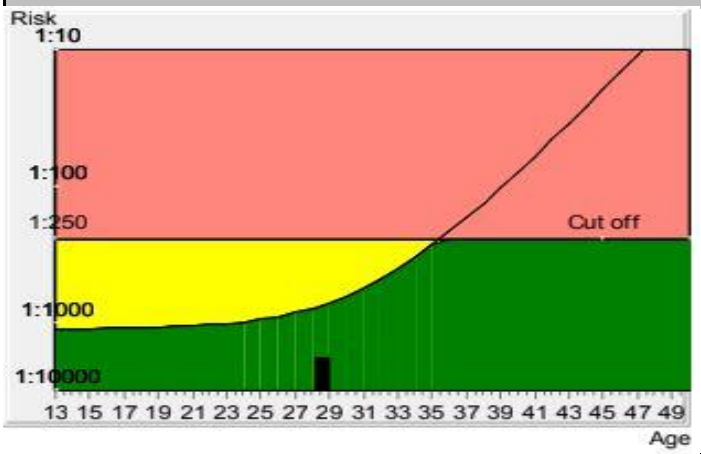
Ultrasound Data

Gestational age	12+2
Method	CRL (<>Robinson)
Scan Date	23-04-19
Crown Rump Length (mm)	55
Nuchal translucency MoM	0.69
Nasal Bone	present
Sonographer	DR.SHRUTI SANGWAN
Qualification in measuring NT	MD

Risks at sampling date

Age Risk	1:755
Biochemical Trisomy 21 Risk	1:1809
Combined Trisomy 21 Risk	1:9652
Trisomy 13/18 + NT	<1:10000

Risk



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 9652 women with the same data, there is one woman with a trisomy 21 pregnancy and 9651 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