Date of Report	08-03-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Jyoti**	Patient ID	021903070015
Birthday	05-10-83	Sample ID	10413249
Age at delivery	35.4	Sample Date	07/03/19
Gestational age by	13+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	66	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	4.36 mIU/ml	1.07
fb-hCG	59.1 ng/ml	1.43

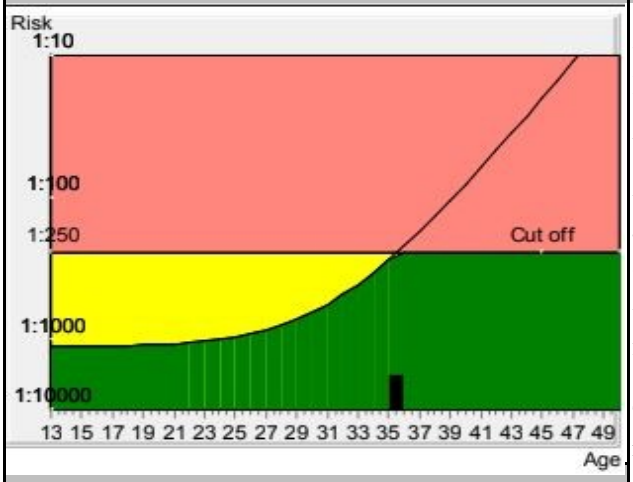
Ultrasound Data

Gestational age	12+6
Method	CRL (<>Hadlock)
Scan Date	06-03-19
CRL measurments	64.9
Nuchal translucency MoM	0.72
Nasal bone	Present
Sonographer	DR.RITU JAIN
Qualifications in measuring NT	

Risks at sampling date

Age Risk	1:252
Biochemical T21 Risk	1:789
Combined Trisomy 21 Risk	1:4059
Trisomy 13/18	<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 4059 women with the same data, there is one woman with a trisomy 21 pregnancy and 4058 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 risk calculations are statistical approach and have no diagnostic values!

The patient combined risk presumes the NT measurement was done according to accepted guidelines

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