Date of Report	20-03-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Gayatri**	Patient ID	011903180146
Birthday	09-01-89	Sample ID	10444394
Age at delivery	30.2	Sample Date	18/03/19
Gestational age by	13+1		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	75	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	1.63	mIU/ml	0.45
fb-hCG	18.55	ng/ml	0.47

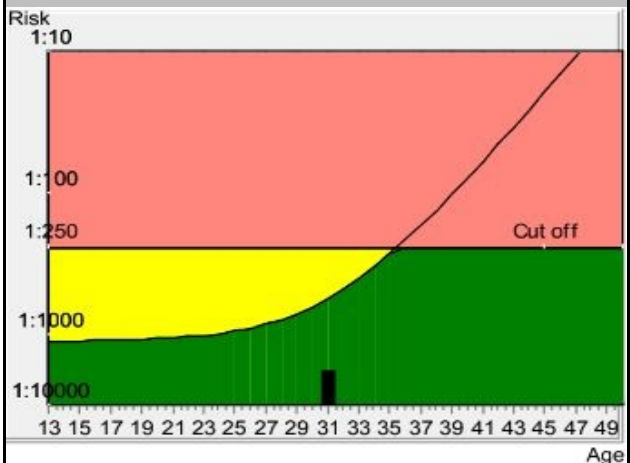
Ultrasound Data

Gestational age	13+0
Method	CRL (<>Hadlock)
Scan Date	17-03-19
CRL measurments	68.3
Nuchal translucency MoM	1.04
Nasal bone	Present
Sonographer	DR.RAKHI BANSAL
Qualifications in measuring NT	MBBS,DMRD

Risks at sampling date

Age Risk	1:645
Biochemical T21 Risk	1:2576
Combined Trisomy 21 Risk	<1:10000
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 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 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