Date of Report	25-02-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Preeti w/o Nitish**	Patient ID	011902240102
Birthday	26-01-89	Sample ID	10426847
Age at delivery	30.1	Sample Date	24/02/19
Gestational age by	12+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	58.5	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	1.45	mIU/ml	0.44
fb-hCG	22.28	ng/ml	0.46

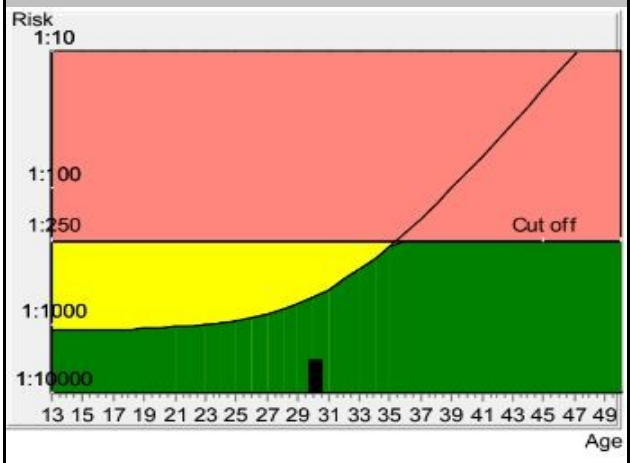
Ultrasound Data

Gestational age	11+6
Method	CRL (<>Hadlock)
Scan Date	23-02-19
CRL measurments	52.1
Nuchal translucency MoM	0.72
Nasal bone	Present
Sonographer	SHAILENDER AGGARWAL
Qualifications in measuring NT	MBBS, MDS

Risks at sampling date

Age Risk	1:627
Biochemical T21 Risk	1:2590
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18	<1:10000

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

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

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