Date of Report	03-04-19
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Sneha Kumari**	Patient ID	011904020185
Birthday	25-12-86	Sample ID	10396242
Age at delivery	32.3	Sample Date	02/04/2019
Gestational age by	13+1		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	55	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	5.12	mIU/ml	0.97
fb-hCG	28.54	ng/ml	0.66

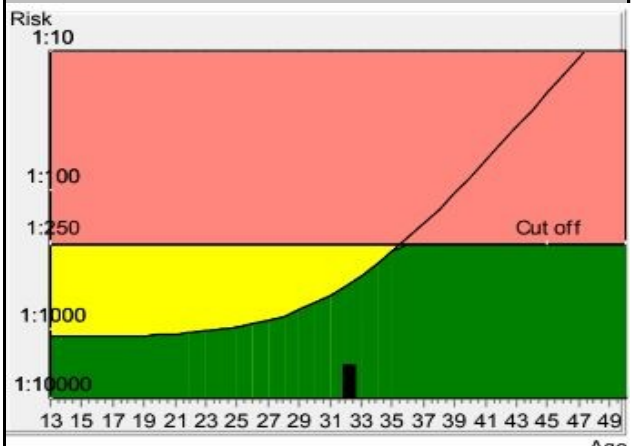
Ultrasound Data

Gestational age	13+1
Method	CRL (<>Hadlock)
Scan Date	02-04-19
CRL measurments	68
Nuchal translucency MoM	0.64
Nasal bone	Present
Sonographer	DR.PRAKASH LALCHANDANI
Qualifications in measuring NT	MD

Risks at sampling date

Age Risk	1:474
Biochemical T21 Risk	1:6674
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