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Date of Report	20-02-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Saumya**	Patient ID	021902180020
Birthday	01-01-88	Sample ID	10413734
Age at delivery	31.1	Sample Date	18/02/19
Gestational age by	11+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	86	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	1.26	mIU/ml	0.75
fb-hCG	33.26	ng/ml	0.74

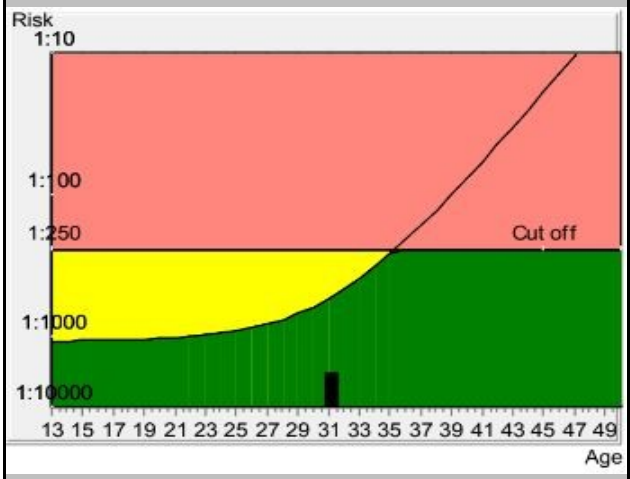
Risks at sampling date

Age Risk	1:534
Biochemical T21 Risk	1:3325
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18	<1:10000

Ultrasound Data

Gestational age	11+3
Method	CRL (<>Hadlock)
Scan Date	18-02-19
CRL measurments	45
Nuchal translucency MoM	0.97
Nasal bone	Present
Sonographer	DR.ANKUSH DHANADIA
Qualifications in measuring NT	MD

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