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Date of Report	06-03-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Bhawana Tiwari**	Patient ID	011903040255
Birthday	27-01-87	Sample ID	10433027
Age at delivery	32.1	Sample Date	04/03/19
Gestational age by	11+1		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	76	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	1.54	mIU/ml	0.81
fb-hCG	36.59	ng/ml	0.72

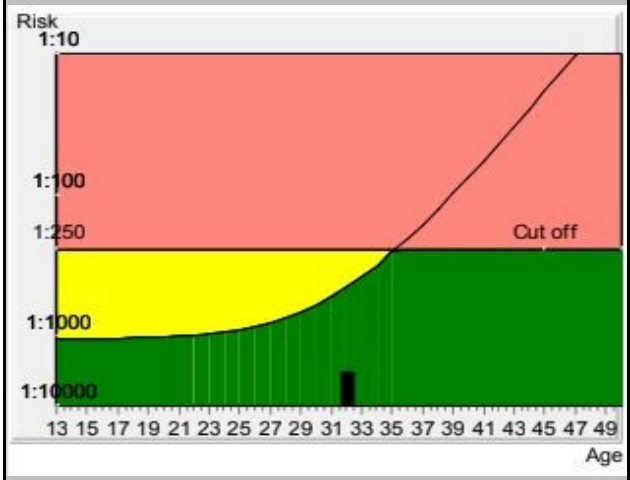
Risks at sampling date

Age Risk	1:450
Biochemical T21 Risk	1:3567
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18	<1:10000

Ultrasound Data

Gestational age	11+0
Method	CRL (<>Hadlock)
Scan Date	04-03-19
CRL measurments	41.2
Nuchal translucency MoM	1.03
Nasal bone	Present
Sonographer	DR.SANJEEV KUMAR SINGHAL
Qualifications in measuring NT	MBBS,PGDUS,DMRD

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