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Date of Report	04-02-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs ANAMIKA**	Patient ID	011902030072
Birthday	20-05-90	Sample ID	10396653
Age at delivery	28.7	Sample Date	03/02/19
Gestational age	12+3		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	45.9	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	3.18	mIU/ml	0.63
F-BhCG	63.08	ng/ml	1.26

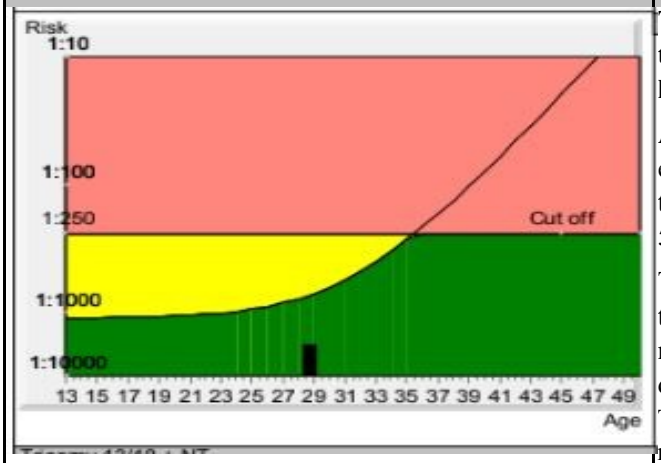
Risks at sampling date

Age Risk	1:742
Biochemical T21 Risk	1:927
Combined T21 Risk	1:5376
Trisomy 18	<1:10000

Ultrasound Data

Gestational age	12+3
Method	CRL (<>Hadlock)
Scan Date	03-02-19
CRL measurments	59 mm
NT Mom	0.65
Nasal Bone	Present
Sonographer	DR.PRAKASH LALCHANDANI
Qualifications in measuring NT	MD

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 5376 women with the same data, there is one woman with a trisomy 21 pregnancy and 5375 women with not affected pregnancies.

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 18 is <1:10000, which represents a low risk.

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