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Date of Report	23-03-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs RANJANA**	Patient ID	051903220029
Birthday	09-05-91	Sample ID	10450782
Age at delivery	27.9	Sample Date	22/03/2019
Gestational age by	12+6		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	65.3	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

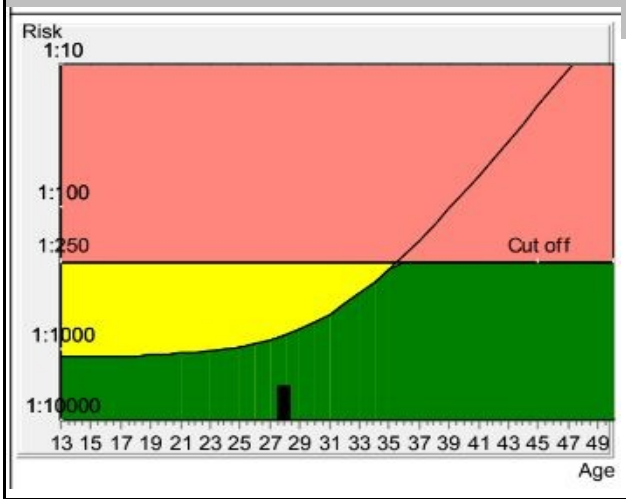
Parameter	Value		Corr Mom
PAPP-A	1.84	mIU/ml	0.47
fb-hCG	17.18	ng/ml	0.41

Ultrasound Data

Gestational age	12+6
Method	CRL (<>Hadlock)
Scan Date	22-03-19
CRL measurments	64.1
Nuchal translucency MoM	0.67
Nasal bone	Present
Sonographer	DR. A. DHANADIA
Qualifications in measuring NT	MD

Risks at sampling date

Age Risk	1:814
Biochemical T21 Risk	1:4960
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18	<1:10000



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 4960 women with the same data, there is one woman with a trisomy 21 pregnancy and 4959 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 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