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Date of Report	03-04-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Ankita**	Patient ID	011904030053
Birthday	06-04-94	Sample ID	10470944
Age at delivery	25.0	Sample Date	03/04/2019
Gestational age by	12+3		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	63	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	4.15	mIU/ml	1.19
fb-hCG	60.45	ng/ml	1.35

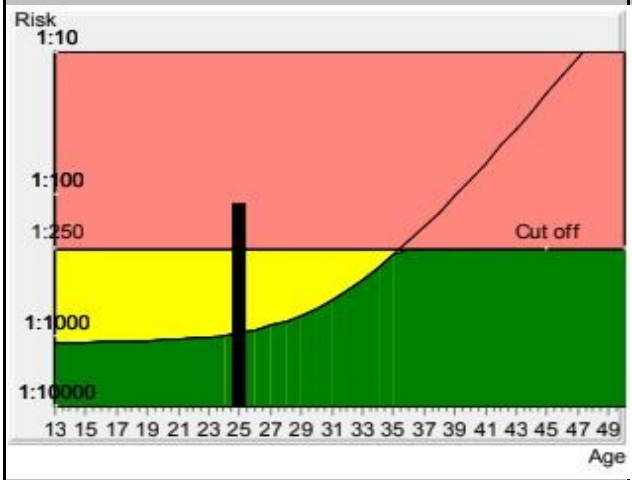
Risks at sampling date

Age Risk	1:958
Biochemical T21 Risk	1:4215
Combined Trisomy 21 Risk	1:115
Trisomy 13/18	1:7961

Ultrasound Data

Gestational age	12+1
Method	CRL (<>Hadlock)
Scan Date	01-04-19
CRL measurments	56
Nuchal translucency MoM	2.23
Nasal bone	Present
Sonographer	DR.AMIT PANT
Qualifications in measuring NT	HMC

Risk



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:7961, which represents a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 115 women with the same data, there is one woman with a trisomy 21 pregnancy and 114 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

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