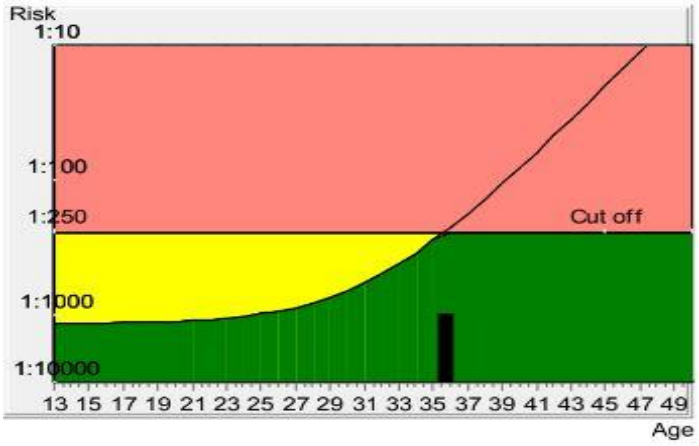
Date of Report	10/5/2019
PRISCA	5.0.2.37

Patient Data			
Name	Mrs Sangeeta	Patient ID	011905090118
Birthday	1/8/1983	Sample ID	10461087
Age at delivery	35.8	Sample Date	09/05/19
Gestational age	13+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	48	Diabetes	no	Pregnancies	unknown
Smoker	no	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+3
PAPP-A	4.55 mIU/ml	0.55	Method	CRL (<>Robinson)
fb-hCG	58.26 ng/ml	1.30	Scan Date	8/5/2019
			Crown Rump Length (mm)	73.5
Risks at sampling date			Nuchal translucency MoM	0.93
Age Risk		1:238	Nasal Bone	present
Biochemical Trisomy 21 Risk		1:194	Sonographer	DR.ANANT ANUPAM
Combined Trisomy 21 Risk		1:986	Qualification in measuring NT	MBBS,DMRD
Trisomy 13/18 + NT		<1:10000		

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)

TThe 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 986 women with the same data, there is one woman with a trisomy 21 pregnancy and 985 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 the risk calculations are statistical aapproaches and have no diagnostic value!

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

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