Date of Report	31-10-18
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Pooja W/o Sandeep Featus 1**	Patient ID	011810290346
Birthday	24-07-93	Sample ID	10337532
Age at delivery	25.3	Sample Date	29/10/18
Gestational age	13+4		

Correction factors					
Fetuses	2	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	65.3	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	3.87	mIU/ml	0.42
fb-hCG	36.86	ng/ml	0.44

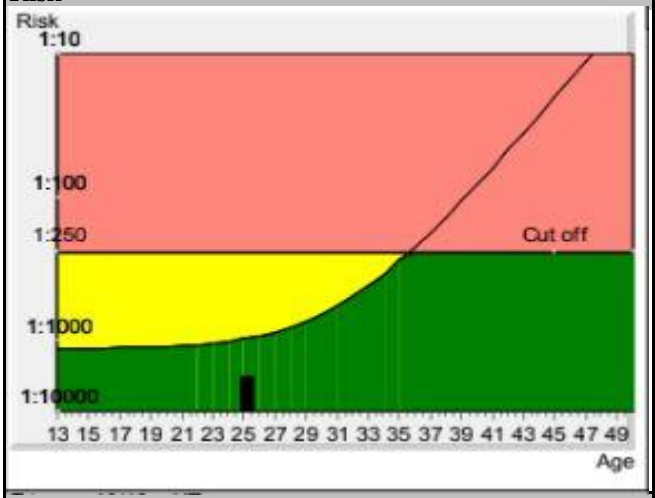
Risks at sampling date

Age Risk	1:983
Biochemical T21 Risk	1:3790
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	13+3
Method	CRL (<>Robinson)
Scan Date	29-10-18
Crown rump length (mm)	73
Nuchal translucency MoM	0.5
Nasal Bone	Present
Sonographer	DR.SURENDER SINGH
Qualifications in measuring NT	MBBS,MD

Risk



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 3790 with the same data, there is one women with a trisomy 21 pregnancy and 3789 women with not affected pregnacies.

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 patient combined risk presumes the NT measurement was done according to accepted guidelines.

The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!

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