Date of Report	24/07/2019
PRISCA	5.0.2.37

Patient Data		
Name	Mrs Pooja	Patient ID
Birthday	8/7/1994	Sample ID
Age at delivery	25	Sample Date
Gestational age	12+6	

Patient Data	
Patient ID	021907220017
Sample ID	10571833
Sample Date	22/07/19

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	50	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data		
Parameter	Value	Corr Mom
PAPP-A	2.81 mIU/ml	0.45
fb-hCG	39.55 ng/ml	0.83

Risks at sampling date	
Age Risk	1:971
Biochemical Trisomy 21 Risk	1:1281
Combined Trisomy 21 Risk	1:2036
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	12+5
Method	CRL (<>Robinson)
Scan Date	22/07/19
Crown Rump Length (mm)	61.4
Nuchal translucency MoM	1.32
Nasal Bone	present
Sonographer	DR.SHWETA AGARWAL
Qualification in measuring NT	DNB

Risk

Down's Syndrome Risk (Trisomy 21 Screening)
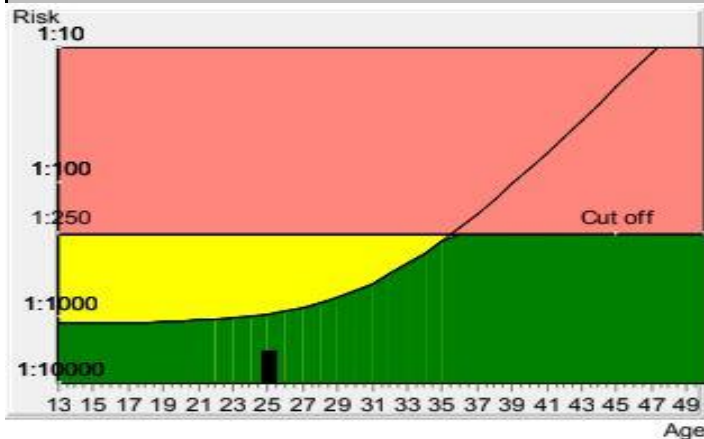
TThe calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 2036 women with the same data, there is one woman with a trisomy 21 pregnancy and 2035 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!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

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