Date of Report	19/05/2019
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Cheena**	Patient ID	011905170250
Birthday	4/8/1989	Sample ID	10609611
Age at sample date	29.8	Sample Date	17/05/2019
Gestational age by CRL	13+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	76	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.69 mIU/ml	1.18
fb-hCG	35.1 ng/ml	0.91

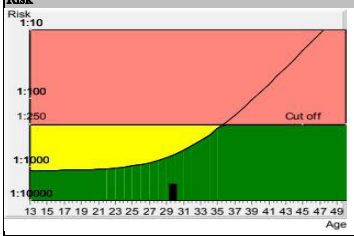
Risks at sampling date

Age Risk	1:687
Biochemical T21 Risk	1:7411
Combined Trisomy 21 Risk	1:9461
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age by CRL	13+0
Method	CRL (<>Robinson)
Scan Date	14/05/2019
Crown rump length (mm)	63
Nuchal translucency MoM	1.36
Nasal Bone	Present
Sonographer	DR. SHRUTI SANGWAN
Qualifications in measuring NT	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 more than 9641 women with the same data, there is one woman with a trisomy 21 pregnancy.
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