Date of Report	12/11/2019
PRISCA	5.0.2.37

Patient Data

Name	Mrs Sapna Khowal F2	Patient ID	071911090009
Birthday	4/8/1986	Sample ID	10506190
Age at delivery	33.3	Sample Date	09/11/19
Gestational age	13+1		

Correction factors

Fetuses	2	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	57	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	6.56 mIU/ml	0.59
fb-hCG	133.05 ng/ml	1.39

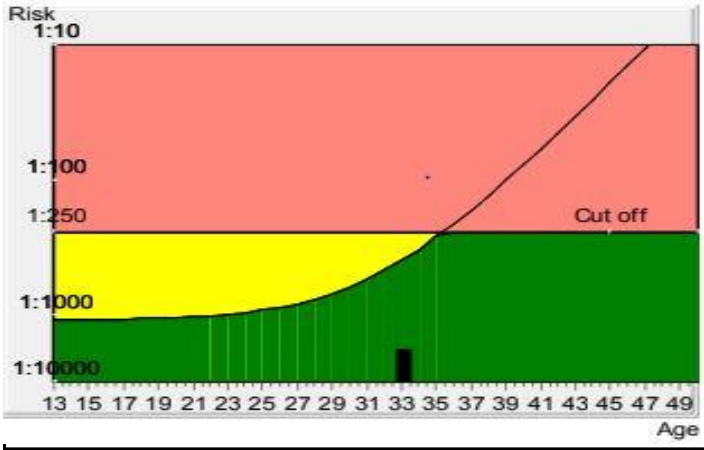
Ultrasound Data

Gestational age	12+5
Method	CRL (<>Robinson)
Scan Date	8/11/2019
Crown Rump Length (mm)	67.2
Nuchal translucency MoM	0.88
Nasal Bone	present
Sonographer	DR. S.P. JAYANT
Qualification in measuring NT	MD

Risks at sampling date

Age Risk	1:396
Biochemical Trisomy 21 Risk	1:342
Combined Trisomy 21 Risk	1:1884
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)

The 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 1884 women with the same data, there is one woman with a trisomy 21 pregnancy and 1883 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