Date of Report	14/02/2020
PRISCA	5.0.2.37

Patient Data

Name	Mrs Neha Garg	Patient ID	062002120031
Birthday	07/01/1986	Sample ID	10658529
Age at delivery	34.1	Sample Date	12/02/2020
Gestational age	12+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	73	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.6 mIU/ml	1.05
fb-hCG	61.4 ng/ml	1.38

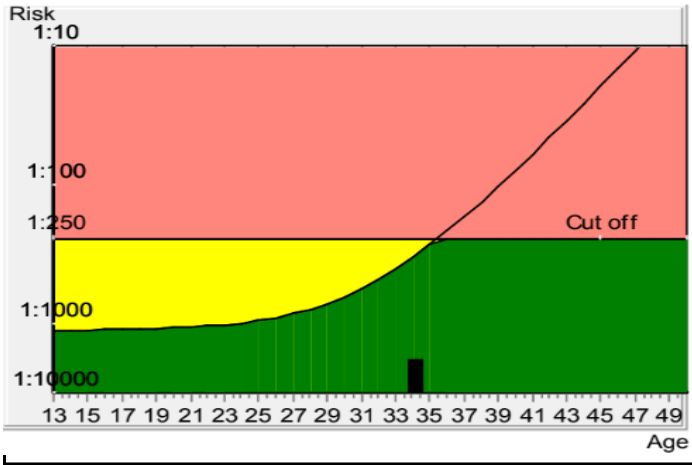
Risks at sampling date

Age Risk	1:328
Biochemical Trisomy 21 Risk	1:1071
Combined Trisomy 21 Risk	1:5513
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+1
Method	CRL (<>Robinson)
Scan Date	10/2/2020
Crown Rump Length (mm)	54
Nuchal translucency MoM	0.55
Nasal Bone	present
Sonographer	DR. SAPNA SHARMA
Qualification in measuring NT	DNB, HMC

Risk



Trisomy 13/18 + NT

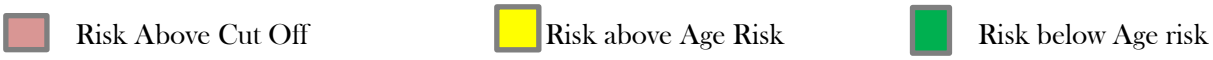
The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.

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 5513 women with the same data, there is one woman with a trisomy 21 pregnancy and 5512 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