Date of Report	21/01/2020
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Mandeep Kaur**	Patient ID	012001300025
Birthday	6/4/1994	Sample ID	10520499
Age at delivery	25.8	Sample Date	30/01/2020
Gestational age	13+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	60	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.16 mIU/ml	0.84
fb-hCG	52.6 ng/ml	1.25

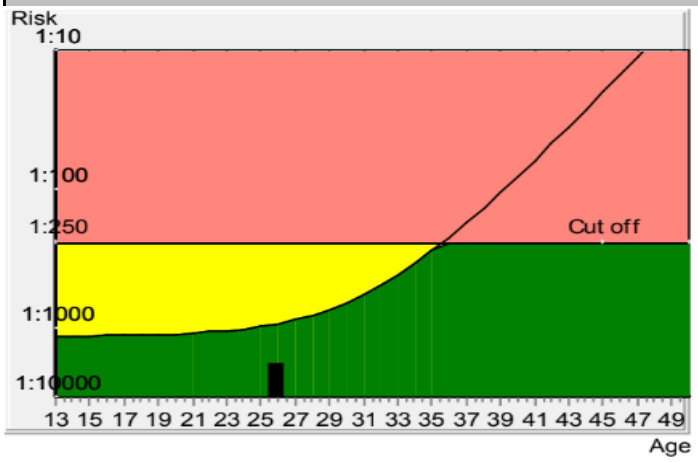
Risks at sampling date

Age Risk	1:952
Biochemical T21 Risk	1:2418
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	13+3
Method	CRL (<>Robinson)
Scan Date	28/01/20
Crown rump length (mm)	69.9
Nuchal translucency MoM	0.8
Nasal Bone	Present
Sonographer	Dr. Vikas Deswal
Qualifications in measuring NT	C/R

Risk



Trisomy 13/18 + NT

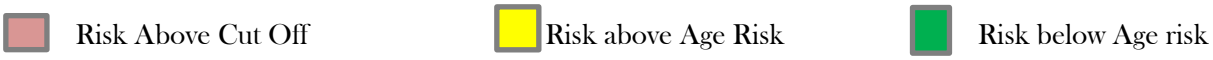
The calculated risk for Trisomy 13/18 (with NT) is 1:10000, which indicates a low risk

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 10000 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