Date of Report	1/9/2019
PRISCA	5.0.2.37

Patient Data

Name	Mrs Chanchal	Patient ID	011909030150
Birthday	21/08/1989	Sample ID	10586287
Age at delivery	30	Sample Date	03/09/19
Gestational age	12+0		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	52	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	4.12 mIU/ml	0.95
fb-hCG	37.6 ng/ml	0.72

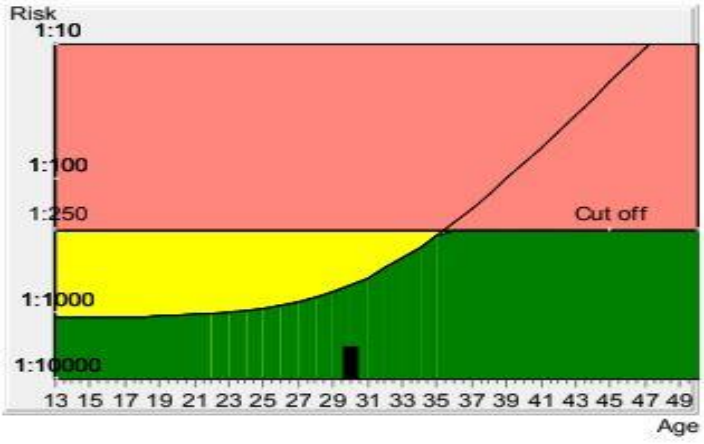
Risks at sampling date

Age Risk	1:630
Biochemical Trisomy 21 Risk	1:7119
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+0
Method	CRL (<>Robinson)
Scan Date	3/9/2019
Crown Rump Length (mm)	53.1
Nuchal translucency MoM	1.2
Nasal Bone	present
Sonographer	Dr. Shweta SM
Qualification in measuring NT	HMC 7111

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)

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 more than 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