Date of Report	19/04/19
PRISCA	5.0.2.37

Patient Data

Name	Mrs Sarita	Patient ID	011904160408
Birthday	12/5/1996	Sample ID	10455307
Age at delivery	23.4	Sample Date	16-04-19
Gestational age	17+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	48	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
AFP	61.72 ng/ml	1.47
uE3	1.12 mIU/ml	1.29
hCG	31082.5 ng/ml	1.43

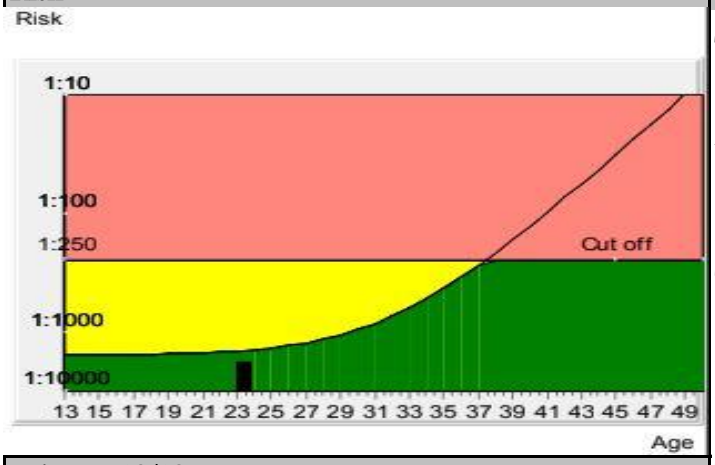
Risks at sampling date

Age Risk	1:1452
Biochemical T21 Risk	1:5659
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	14+1
Method	CRL (<>Robinson)
Scan Date	
NT mom	1/0/1900
Crown rump length (mm)	65.1
Nuchal translucency MoM	1
Nasal Bone	Present
Sonographer	DR.I.HANGLOO
Qualifications in measuring NT	PGD

Risk



Trisomy 13/18 + NT

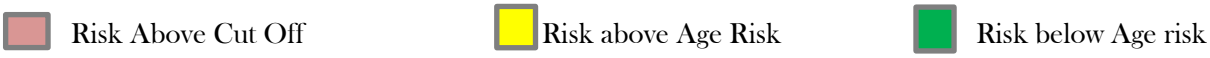
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

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 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