Date of Report	26/01/2020
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Farhana**	Patient ID	012001250071
Birthday	15/12/1988	Sample ID	10565915
Age at delivery	31.5	Sample Date	25/01/2020

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	68	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr MoM
AFP	93.5 ng/ml	1.85
uE3	1.55 ng/ml	1.21
hCG	9591.5 mIU/ml	0.5

Risks at sampling date

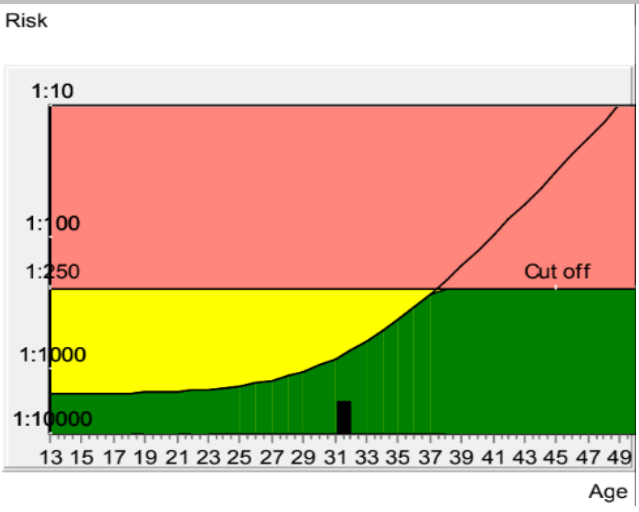
Age Risk	1:794
Trisomy 21 Risk	<1:10000
Combined Trisomy 21 Risk	<1:10000
Trisomy 18 Risk	<1:10000

Ultrasound Data

Gestational age	18+6
Method	CRL (<>Robinson)

CRL: 67.8 mm, NT 1.8 mm

Risk



Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 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 patient combined risk presumes the NT measurement was done according to accepted guidelines.

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!

Trisomy 18

The calculated risk for Trisomy 18 is <1:10000, which indicates a low risk

Neural Tube Defect (NTD) Screening

The corrected MoM for AFP (1.85) is located in the low risk area for neural tube defects.

The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!

Risk Above Cut Off | Risk above Age Risk | Risk below Age risk