Date of Report	13-03-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Pallavi**	Patient ID	011902280152
Birthday	31-05-91	Sample ID	10381020
Age at delivery	28.3	Sample Date	01/03/2019
Gestational age by	15+3		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	65	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
AFP	35.6	ng/ml	1.08
uE3	0.65	ng/ml	1.38
hCG	32622.50	mIU/ml	1.00

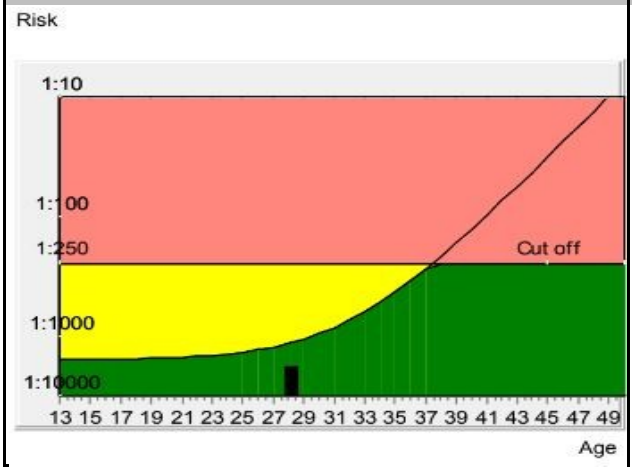
Risks at sampling date

Age Risk	1:1142
Biochemical T21 Risk	1:5052
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18	<1:10000

Ultrasound Data

Gestational age	12+3
Method	CRL
Scan Date	20-02-19
CRL measurments	58.8
Nuchal translucency MoM	0.72
Nasal bone	
Sonographer	Dr. Sanjeev Kumar Singhal
Qualifications in measuring NT	MBBS, PGDUS, DMRD

Risk



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 is < 1:10000, which represents a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (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 risk calculations are statistical approaches and have no diagnostic value!

The patient combined risk presumes the NT measurement was done according to accepted guidelines

The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

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