Date of Report	28-12-18
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Sweta Kumari**	Patient ID	011812260150
Birthday	20-10-90	Sample ID	10415591
Age at delivery	28.2	Sample Date	26/12/2018
Gestational age	12+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	72	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	3.45	mIU/ml	1.22
fb-hCG	76.78	ng/ml	1.76

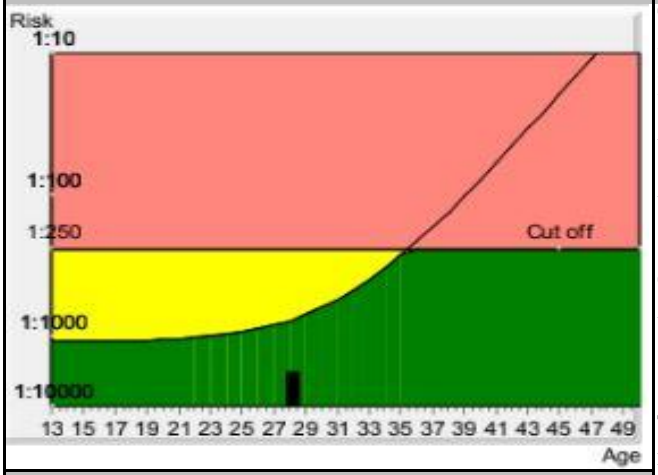
Risks at sampling date

Age Risk	1:776
Biochemical T21 Risk	1:1912
Combined Trisomy 21 Risk	1:8923
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	11+5
Method	CRL (<>Hadlock)
Scan Date	23-12-18
CRL	44
Nuchal translucency MoM	0.90
Nasal Bone	Present
Sonographer	DR.RAJESH ARORA
Qualifications in measuring NT	HMC

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)

The 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 8923 women with the same data, there is one woman with a trisomy 21 pregnancy and 8922 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approach and have no diagnostic values!

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

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