Date of Report 14-12-19
PRISCA 5.0.2.37

Patient Data			
Name	MRS SANDHYA	Patient ID	011912130051
Birthday	25-04-94	Sample ID	10528880
Age at delivery	25.6	Sample Date	13/12/2019
Gestational age	12+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	52	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	4.61 mIU/ml	0.95
fb-hCG	50.1 ng/ml	1

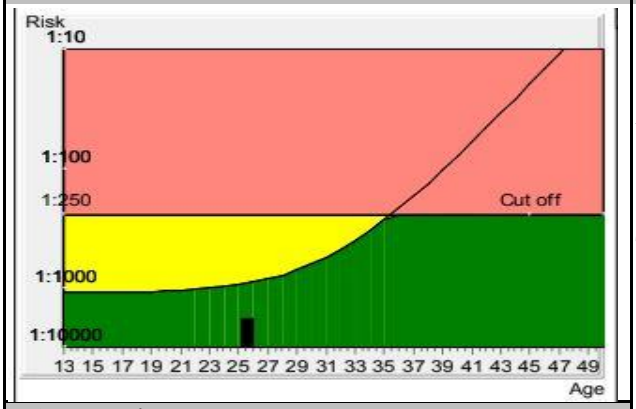
Ultrasound Data

Gestational age	12+1
Method	CRL (<>Robinson)
Scan Date	12-12-19
Crown rump length in mm	55.7
Nuchal translucency MoM	1.16
Nasal Bone	Present
sonographer	DR.RUBY RAHUL
Qualification in measuring NT	CON. RADIOLOGIT

Risks at sampling date

Age Risk	1:925
Biochemical T21 risk	1:5205
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

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 and 9999 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 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