Date of Report	17-12-2020
PRISCA	5.0.2.37

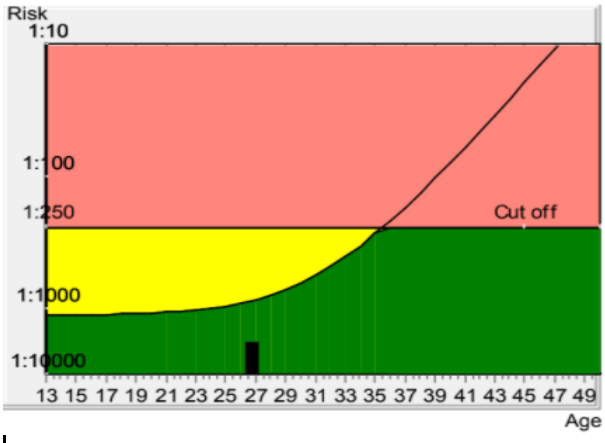
Patient Data			
Name	MRS PRIYA KUMARI	Patient ID	052012160008
Birthday	19-02-1994	Sample ID	10792537
Age at delivery	26.8	Sample Date	16/12/2020
Gestational age	12+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	53.4	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	5.3 mIU/ml	0.96
fb-hCG	116.5 ng/ml	2.45

Ultrasound Data	
Gestational age	12+4
Method	CRL(<>Robinson
Scan date	15-12-2020
Crown rump length in mm	62.1
Nuchal translucency MOM	1.06
Nasal bone	Present
Sonographer	DR.SHYAM
Qualification in measuring NT	

Risks at sampling date	
Age Risk	1:876
Biochemical T21 risk	1:565
Combined Trisomy 21 Risk	1:1940
Trisomy 13/18 + NT	<1:10000



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 1940 women with the same data, there is one woman with a trisomy 21 pregnancy and 1939 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

Trisomy 13/18 + NT

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

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