Date of Report	03-11-2020
PRISCA	5.0.2.37

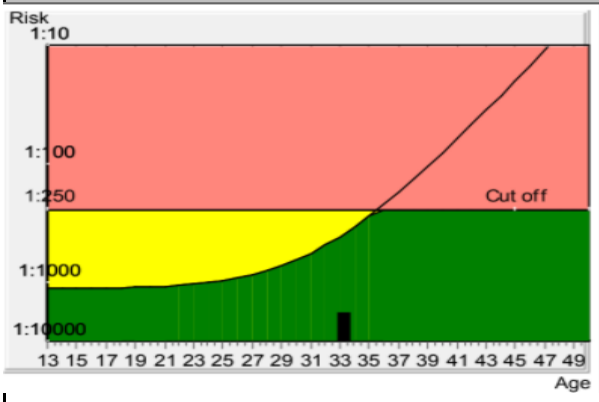
Patient Data			
Name	MRS YAMINI SAXENA	Patient ID	012011010152
Birthday	07-08-1987	Sample ID	DPLTA00294572
Age at delivery	33.2	Sample Date	1/11/2020
Gestational age	13+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	74	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	1.85 mIU/ml	0.45	Method	CRL(<>Robinson
fb-hCG	19.65 ng/ml	0.47	Scan date	30-10-2020
			Crown rump length in mm	60.2

Risks at sampling date			
Age Risk	1:396	Nuchal translucency MOM	0.7
Biochemical T21 risk	1:1594	Nasal bone	Present
Combined Trisomy 21 Risk	1:8780	Sonographer	DR.ARUN MAHAJAN
Trisomy 13/18 + NT	<1:10000	Qualification in measuring NT	HMC

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