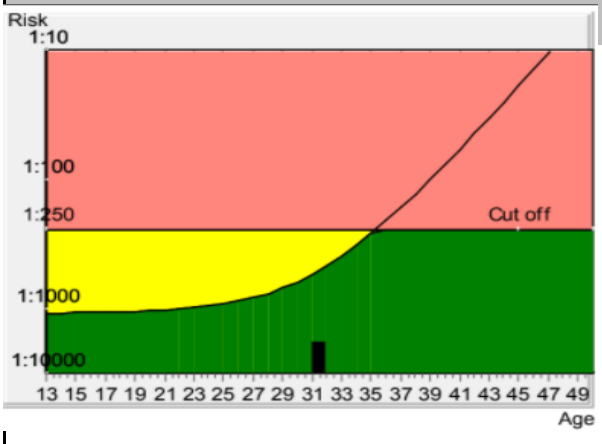
Date of Report	15-03-2021
PRISCA	5.0.2.37

Patient Data			
Name	MRS MANISHA	Patient ID	012103140006
Birthday	07-10-1989	Sample ID	10868219
Age at delivery	31.4	Sample Date	14/03/2021
Gestational age	11+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	62	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+3
PAPP-A	2.35 mIU/ml	0.8	Method	CRL(<>Robinson
fb-hCG	88.7 ng/ml	1.72	Scan date	13-03-2021
			Crown rump length in mm	45

Risks at sampling date			
Age Risk	1:510	Nuchal translucency MOM	0.64
Biochemical T21 risk	1:531	Nasal bone	Present
Combined Trisomy 21 Risk	1:3027	Sonographer	DR.VIDYA SAGAR
Trisomy 13/18 + NT	<1:10000	Qualification in measuring NT	MD



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 3027 women with the same data, there is one woman with a trisomy 21 pregnancy and 3026 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