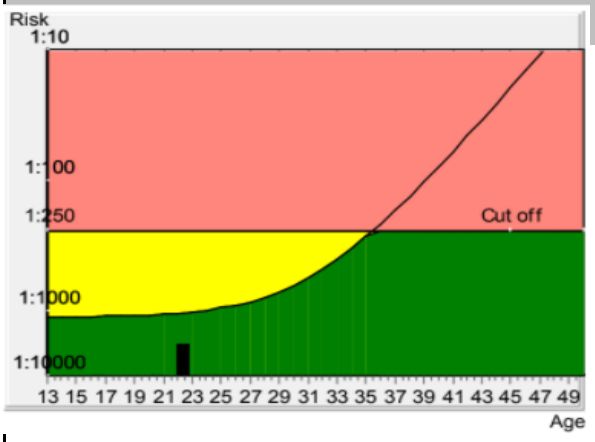
Date of Report	17-03-2021
PRISCA	5.0.2.37

Patient Data			
Name	MRS VANDANA	Patient ID	012103160009
Birthday	20-10-1998	Sample ID	10868522
Age at delivery	22.4	Sample Date	16/03/2021
Gestational age	12+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	60	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+3
PAPP-A	4.08 mIU/ml	0.89	Method	CRL(<>Robinson
fb-hCG	54.2 ng/ml	1.17	Scan date	15-03-2021
			Crown rump length in mm	60.3
Risks at sampling date			Nuchal translucency MOM	0.89
Age Risk		1:1047	Nasal bone	Present
Biochemical T21 risk		1:3565	Sonographer	DR.RANJAN KUMAR
Combined Trisomy 21 Risk		<1:10000	Qualification in measuring NT	MD
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 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

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