Date of Report		2/8/2021
PRISCA		5.1.0.17

Patient Data

Name	**MRS.PREETI KUMARI**	Patient ID	012108010013
Birthday	21/05/1987	Sample ID	10960741
Age at delivery	34.2	Sample Date	01/08/2021
Gestational age	13+1		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	63.9	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.18 mIU/ml	0.61
fb-hCG	52.5 ng/ml	1.23

Risks at sampling date

Age Risk	1:330
Biochemical T21 risk	1:408
Combined Trisomy 21 Risk	1:2376
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	13+0
Method	CRL(<>Robinson
Scan date	31/07/2021
Crown rump length in mm	68.2
Nuchal translucency MOM	0.7
Nasal bone	Present
Sonographer	DR. RANJAN KUMAR
Qualification in measuring NT	MD
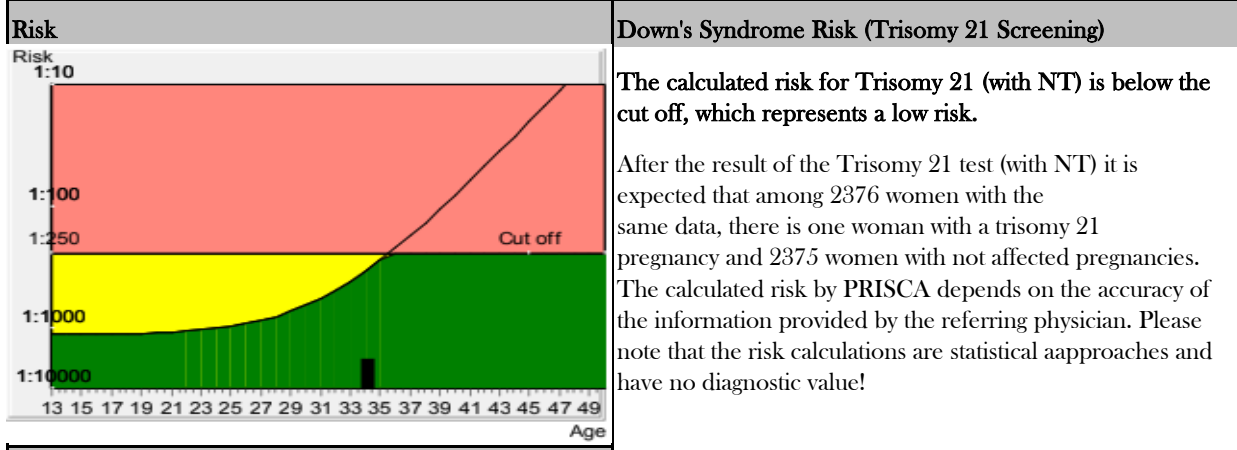
Risk

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 2376 women with the same data, there is one woman with a trisomy 21 pregnancy and 2375 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