Date of Report	05-05-2021
PRISCA	5.0.2.37

Patient Data			
Name	**MS. RAJNISH**	Patient ID	012104280045
Birthday	01-01-1989	Sample ID	10913519
Age at delivery	32.3	Sample Date	28/04/2021
Gestational age	12+6		

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

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	2.74 mIU/ml	0.49
fb-hCG	31.2 ng/ml	0.61

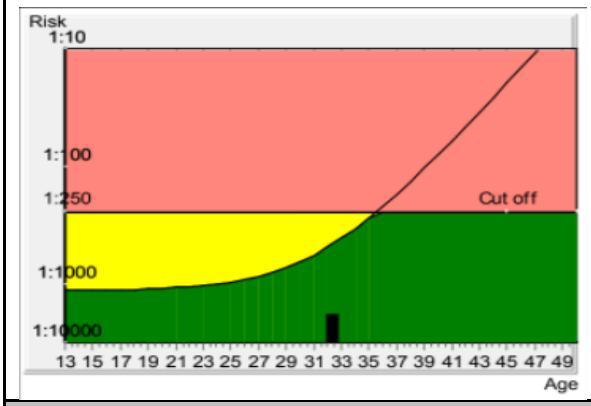
Ultrasound Data

Gestational age	12+5
Method	CRL(<>Robinson
Scan date	27-04-2021
Crown rump length in mm	64.3
Nuchal translucency MOM	0.97
Nasal bone	Present
Sonographer	Dr Ranjan Kumar
Qualification in measuring NT	MD

Risks at sampling date

Age Risk	1:465
Biochemical T21 risk	1:1507
Combined Trisomy 21 Risk	1:6952
Trisomy 13/18 + NT	<1:10000

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 6952 women with the same data, there is one woman with a trisomy 21 pregnancy and 6951 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