Date of Report	14-04-2021
PRISCA	5.0.2.37

Patient Data			
Name	MRS RANIPAL	Patient ID	012104120241
Birthday	24-05-1987	Sample ID	10924811
Age at delivery	33.9	Sample Date	12/04/2021
Gestational age	13+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	61.7	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

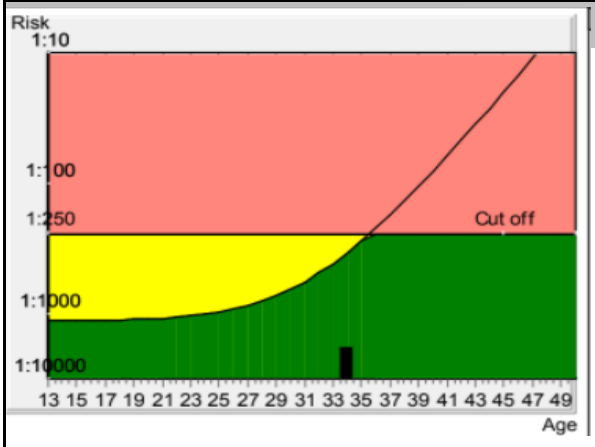
Parameter	Value	Corr Mom
PAPP-A	3.95 mIU/ml	0.77
fb-hCG	56.8 ng/ml	1.3

Ultrasound Data

Gestational age	12+6
Method	CRL(<>Robinson
Scan date	12-04-2021
Crown rump length in mm	64.4
Nuchal translucency MOM	0.61
Nasal bone	Present
Sonographer	DR.HARSHA SEHGAL
Qualification in measuring NT	FMF,UK

Risks at sampling date

Age Risk	1:349
Biochemical T21 risk	1:659
Combined Trisomy 21 Risk	1:3647
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 3647 women with the same data, there is one woman with a trisomy 21 pregnancy and 3646 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