Date of Report	25/8/2021
PRISCA	5.1.0.17

Patient Data			
Name	MRS. PAYAL	Patient ID	012108240019
Birthday	5/6/1989	Sample ID	10960487
Age at delivery	32.2	Sample Date	24/8/2021
Gestational age	11+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	58	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	4.35 mIU/ml	1.28
fb-hCG	96.2 ng/ml	1.86

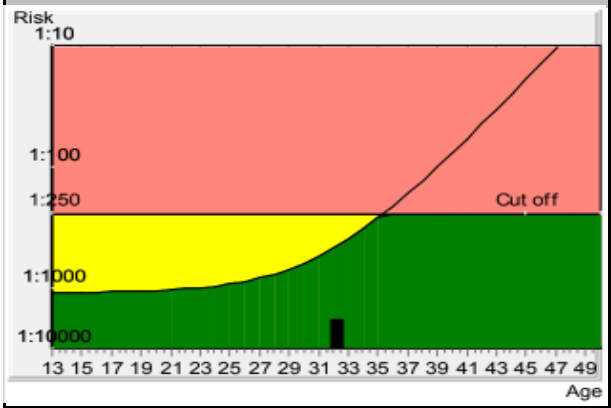
Risks at sampling date

Age Risk	1:452
Biochemical T21 risk	1:1065
Combined Trisomy 21 Risk	1:5256
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	11+4
Method	CRL(<>Robinson
Scan date	23/08/2021
Crown rump length in mm	47.3
Nuchal translucency MOM	0.85
Nasal bone	Present
Sonographer	DR. JYOTIKA
Qualification in measuring NT	MBBS, MD

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 5256 women with the same data, there is one woman with a trisomy 21 pregnancy and 5255 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