Date of Report	19-01-2024
PRISCA	5.2.0.13

Patient Data

Name	**MRS MANISHA**	Patient ID	012401180136
Birthday	25-05-1997	Sample ID	11813715
Age at Sample date	26.7	Sample Date	18-01-2024
Gestational age	13+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	55	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	6.3 mIU/ml	0.76
fb-hCG	18.5 ng/ml	0.63

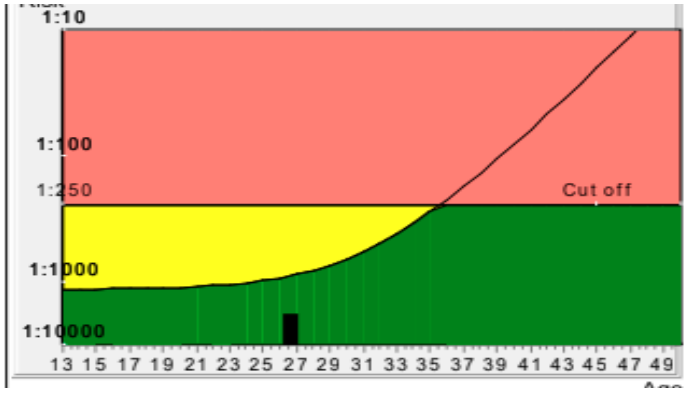
Risks at sampling date

Age Risk	1:907
Biochemical T21 risk	1:8195
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	13+0
Method	CRL (<>Robinson)
Scan date	15-01-2024
Crown rump length in mm	67.4
Nuchal translucency MoM	1.11
Nasal bone	PRESENT
Sonographer	DR MAMTA
Qualifications in measuring NT	MBBS

Risk



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 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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

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