Date of Report 22-11-2023

PRISCA 5.2.0.13

Patient Data			
Name	**MRS PAYAL**	Patient ID	012311210104
Birthday	14-07-1997	Sample ID	11837757
Age at Sample date	26.4	Sample Date	21-11-2023
Gestational age	12+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	47	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.78 mIU/ml	0.49
fb-hCG	26.3 ng/ml	0.68

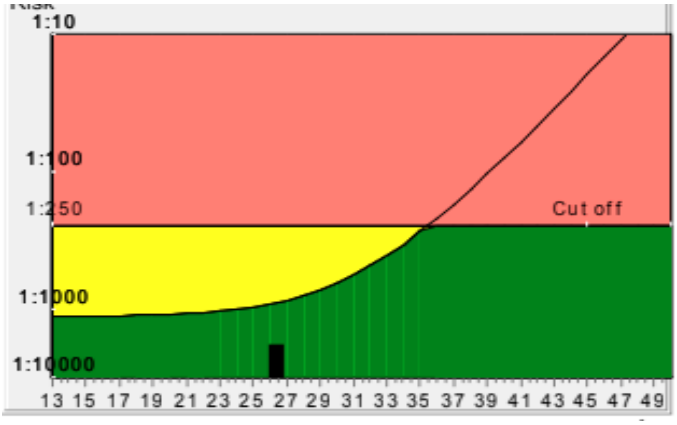
Risks at sampling date

Age Risk	1:902
Biochemical T21 risk	1:2262
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+5
Method	CRL (<>Robinson)
Scan date	21-11-2023
Crown rump length in mm	62.2
Nuchal translucency MoM	0.87
Nasal bone	PRESENT
Sonographer	DR.AMENDA
Qualifications in measuring NT	MD

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