Date of Report	17-05-2024
PRISCA	5.2.0.13

Patient Data

Name	**MRS. MAYA DEVI F3**	Patient ID	102405160016'
Birthday	15-08-1986	Sample ID	11904755
Age at Sample date	37.8	Sample Date	16-05-2024
Gestational age	13+6		

Correction factors

Fetuses	2	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	56.2	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	13.5 mIU/ml	0.78
fb-hCG	148.5 ng/ml	2.71

Risks at sampling date

Age Risk	1:149
Biochemical T21 risk	>1:50
Combined trisomy 21 risk	1:189
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+6
Method	CRL (<>Robinson)
Scan date	09-05-2024
Crown rump length in mm	65.8
Nuchal translucency MoM	1.01
Nasal bone	PRESENT
Sonographer	DR.
Qualifications in measuring NT	MBBS
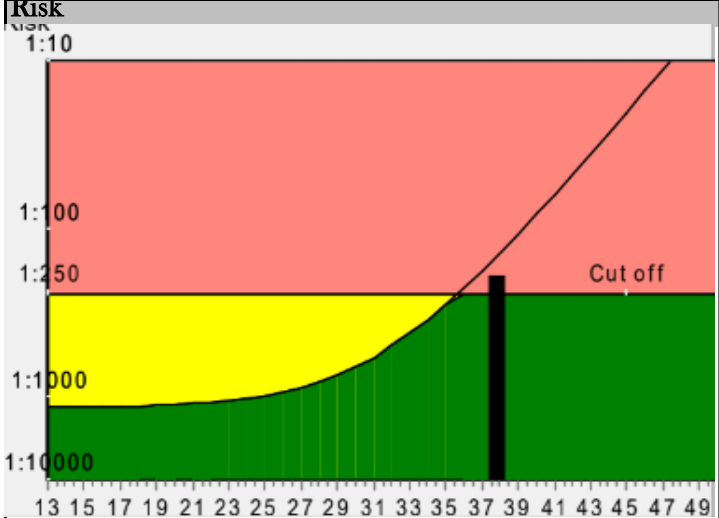
Risk

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 above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 189 women with the same data, there is one woman with a trisomy 21 pregnancy and 188 women with not affected pregnancies. The free beta HCG level is high. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.

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