

Date of Report 29-10-2023  
PRISCA 5.2.0.13

Patient Data	
Name	MRS SNEHA
Birthdate	01-07-1988
Age at Sample date	35.3
Gestational age	13+0

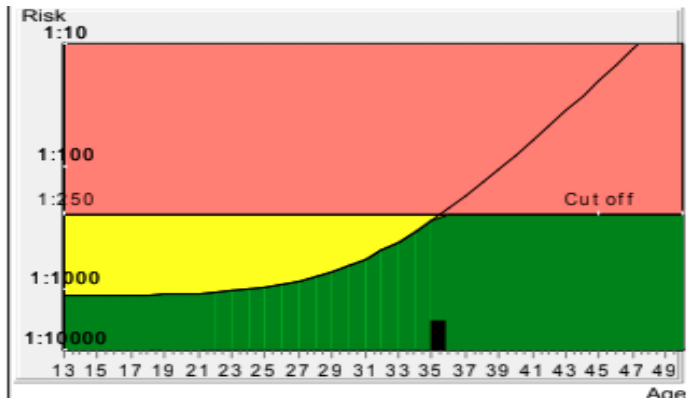
Correction factors	
Fetuses	1 IVF
Weight in kg	58.5
Smoker	NO
Diabetes	NO
Origin	Asian
Previous trisomy 21	unknown
Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom
PAPP-A	3.52 mIU/ml	0.53
fb-hCG	16.2 ng/ml	0.49

Risks at sampling date	Ultrasound Data
Age Risk	1:258
Biochemical T21 risk	1:1519
Combined trisomy 21 risk	1:8051
Trisomy 13/18 + NT	<1:10000
	Gestational age 13+0
	Method CRL (<>Robinson)
	Scan date 28-10-2023
	Crown rump length in mm 67.5
	Nuchal translucency MoM 0.70
	Nasal bone PRESENT
	Sonographer DR.PRAVEEN
	Qualifications in measuring NT MBBS

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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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 8051 women with the same data, there is one woman with a trisomy 21 pregnancy and 8050 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 approaches 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 calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk. 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