



Date of Report 09-01-2024
PRISCA 5.2.0.13

Patient Data	
Name	MRS SHREYA
Birthdate	14-05-1998
Age at Sample date	25.6
Gestational age	12+2

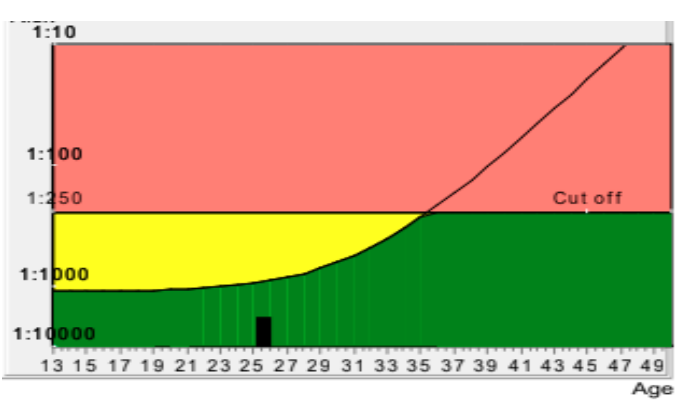
Correction factors	
Fetuses	1 IVF
Weight in kg	60.8
Smoker	NO
Diabetes	NO
Origin	Asian
Previous trisomy 21	unknown
Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom
PAPP-A	3.17 mIU/ml	0.65
fb-hCG	24.7 ng/ml	0.62

Risks at sampling date	Ultrasound Data
Age Risk	1:924
Biochemical T21 risk	1:5836
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000
Gestational age	11+6
Method	CRL (<>Robinson)
Scan date	03-01-2024
Crown rump length in mm	51.8
Nuchal translucency MoM	0.84
Nasal bone	PRESENT
Sonographer	DR NIDHI
Qualifications in measuring NT	RADIOLOGIST

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 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 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