



Date of Report 13-12-2023  
PRISCA 5.2.0.13

Patient Data			
Name	MRS SHEETAL 7168	Patient ID	012312110017
Birthday	25-01-2000	Sample ID	011839511
Age at Sample date	23.9	Sample Date	11-12-2023
Gestational age	13+2		

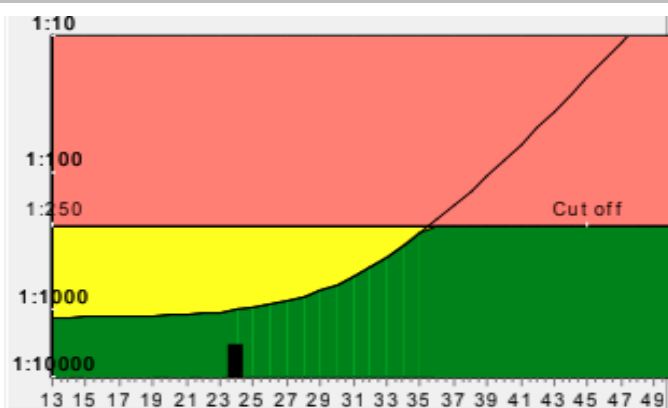
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	100.4	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom	
PAPP-A	4.32 mIU/ml	1.15	Method CRL (<>Robinson)
fb-hCG	25.2 ng/ml	0.97	Scan date 09-12-2023

Risks at sampling date		
Age Risk	1:1029	Crown rump length in mm 69
Biochemical T21 risk	1:9253	Nuchal translucency MoM 0.98
Combined trisomy 21 risk	<1:10000	Nasal bone PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer DR ADIYA
		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 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