



Date of Report 20-10-2023
PRISCA 5.2.0.13

Patient Data			
Name	MRS MEENA	Patient ID	012310190182
Birthday	17-05-2001	Sample ID	11843894
Age at Sample date	22.4	Sample Date	19-10-2023
Gestational age	13+0		

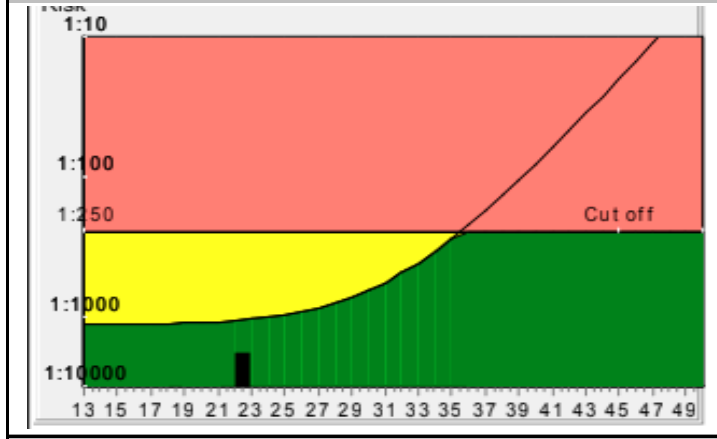
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	59	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom		
PAPP-A	3.87 mIU/ml	0.59	Gestational age	12+6
fb-hCG	38.8 ng/ml	1.18	Method	CRL (<>Robinson)
			Scan date	18-10-2023

Risks at sampling date		Ultrasound Data	
Age Risk	1:1062	Crown rump length in mm	65
Biochemical T21 risk	1:1297	Nuchal translucency MoM	1.08
Combined trisomy 21 risk	1:4475	Nasal bone	PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer	DR HARENDRA
		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 4475 women with the same data, there is one woman with a trisomy 21 pregnancy and 4474 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