



Date of Report 19-12-2023
PRISCA 5.2.0.13

Patient Data			
Name	MRS SAGARIKA	Patient ID	012312180066
Birthday	22-07-1988	Sample ID	11806838
Age at Sample date	35.4	Sample Date	18-12-2023
Gestational age	13+0		

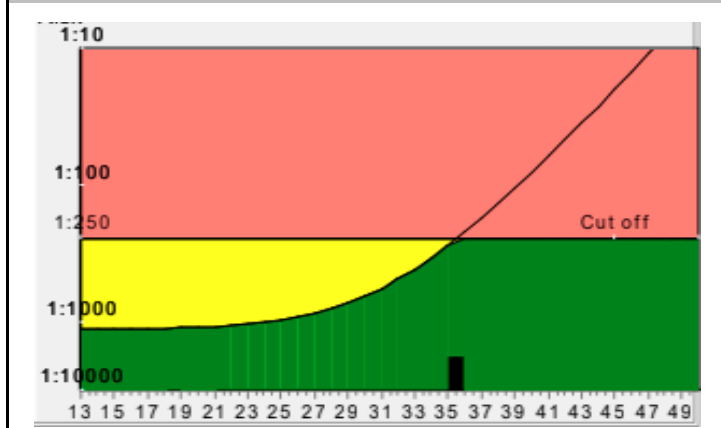
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	67	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom		
PAPP-A	4.5 mIU/ml	0.79	Gestational age	13+0
fb-hCG	41.2 ng/ml	1.3	Method	CRL (<>Robinson)
			Scan date	18-12-2023

Risks at sampling date		Ultrasound Data	
Age Risk	1:253	Crown rump length in mm	68.1
Biochemical T21 risk	1:510	Nuchal translucency MoM	0.87
Combined trisomy 21 risk	1:2649	Nasal bone	PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer	DRA.SHARMA
		Qualifications in measuring NT	MD

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 2649 women with the same data, there is one woman with a trisomy 21 pregnancy and 2648 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 held 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