



Date of Report 23-11-2023
PRISCA 5.2.0.13

Patient Data			
Name	MRS SADHYA	Patient ID	012311220029
Birthday	14-02-1991	Sample ID	11854831
Age at Sample date	32.8	Sample Date	22-11-2023
Gestational age	12+3		

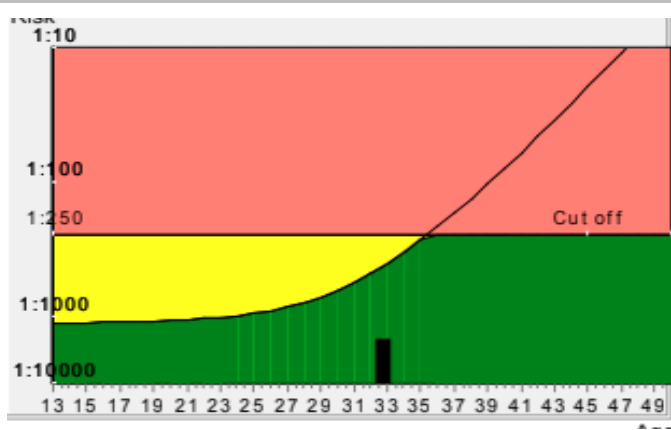
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	67.9	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	3.83 mIU/ml	0.85	Gestational age	12+1
fb-hCG	57.7 ng/ml	1.56	Method	CRL (<>Robinson)
			Scan date	20-11-2023

Risks at sampling date			Ultrasound Data	
Age Risk		1:423	Crown rump length in mm	55
Biochemical T21 risk		1:645	Nuchal translucency MoM	1.20
Combined trisomy 21 risk		1:1466	Nasal bone	PRESENT
Trisomy 13/18 + NT		<1:10000	Sonographer	DR.NIDHI
			Qualifications in measuring NT	.

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 1466 women with the same data, there is one woman with a trisomy 21 pregnancy and 1465 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