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Date of Report 23-09-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name IRS. SONIA W/O MANOJ KUMAR				Patient ID		12409220191
Birthday	nday 17-12-2009			Sample ID		11886196
Age at Sample date 21.8				Sample Date		22-09-2024
Gestational age			12+4	Į.		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	12+3
PAPP-A	6.3	mIU/ml	1.07	Method		CRL (<>Robinson)
fb-hCG	19.6	ng/ml	0.52	Scan date		21-09-2024
Risks at sampling date				Crown rump length in mm 58.8		
Age Risk			1:1061	Nuchal translu	icency MoM	0.84
Biochemical T21 risk			<1:10000	Nasal bone		PRESENT
Combined trisomy 21 risk <1:1000			< 1:10000	Sonographer D		DR.
Trisomy 13/18 + NT <1:3			- <1:10000	Qualifications in measuring NT M		MBBS
Risk 1:10				Down's Syndrome Risk (Trisomy 21 Screening) 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.		
1:1000 1:1000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18+NT				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 aapproaches 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 laboratory cannot be hold responsible for their impact on the		

The calculated risk for Trisomy 13/18 (with NT) is <1:10000,

which indicates a low risk

risk assessment! Calculated risks have no diagnostic values