

which indicates a low risk

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Date of Report 10-03-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name	MRS. SUMITRA YADAV		Patient ID		12405240113
Birthday	07-04-1994		Sample ID		12000216
Age at Sample date	mple date 30.1		Sample Date		24-05-2024
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	67.1 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian	1	
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	ge	13+1
PAPP-A	6.3 mIU/ml	1.06	Method		CRL (<>Robinson)
fb-hCG	158.6 ng/ml	5.24	Scan date		24-05-2024
Risks at sampling date			Crown rump length in mm 69.8		
Age Risk		1:649	Nuchal translu	icency MoM	0.57
Biochemical T21 risk		1:136	Nasal bone		PRESENT
Combined trisomy 21 risk		1:805	Sonographer		DR.
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:1000 1:10000 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 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 805 women with the same data, there is one woman with a trisomy 21 pregnancy and 804 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 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 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		