

which indicates a low risk

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Date of Report 11-12-2024

				PRISCA	5.2.0.13
Patient Data					
Name	MRS. PRIYANKA YADAV		Patient ID		12412100046
Birthday	05-01-2000		Sample ID		11956832
Age at Sample date		24.6	Sample Date		10-12-2024
Gestational age		12+6	6		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ago	e	12+5
PAPP-A	5.3 mIU/ml	1.04	Method		CRL (<>Robinson)
fb-hCG	23.5 ng/ml	0.72	Scan date		09-12-2024
Risks at sampling date			Crown rump length in mm 63.8		
Age Risk		1:975	Nuchal translu	icency MoM	0.79
Biochemical T21 risk		<1:10000	Nasal bone		present
Combined trisomy 21 risk <1:10000		<1:10000	Sonographer DR. MEENU SOLAR		DR. MEENU SOLANKI
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT .		
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Gutoff			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 Trisomy 13/18+NT The calculated risk for Tri		414345474	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 risk assessment! Calculated risks have no diagnostic values		