

which indicates a low risk

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Date of Report 26-07-2024

					PRISCA	5.2.0.13
Patient Data						
Name . NISHA YADAV W/O AISHWARY				Patient ID		12407250365
Birthday 20-06-1999				Sample ID		11874885
Age at Sample date 25.1				Sample Date		25-07-2024
Gestational age 12+5						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	50	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	0.73	Method		CRL(<>Robinson)
fb-hCG	29.1	ng/ml	0.77	Scan date		25-07-2024
Risks at sampling date				Crown rump length in mm 62.7		
Age Risk			1:963	Nuchal translu	icency MoM	1.36
Biochemical T21 risk			1:5281	Nasal bone		present
Combined trisomy 21 risk 1:6943				Sonographer		DR. VIKASH
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MD
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:100 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000				The calculated risk for Trisomy 21 test (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 amog 6943 women with the same data, there is one woman with a trisomy 21 pregnancy and 6942 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;		
Trisomy 13/18 + NT				1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		