

which indicates a low risk

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Date of Report	20-05-2024
PRISCA	5.2.0.13

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Patient Data						
Name	MR	S. NEET	U SHARMA	Patient ID		12405190186
Birthday			22-08-1997	Sample ID		12001233
Age at Sample date			26.7	Sample Date		19-05-2024
Gestational age 13+6						
Correction factors						
Fetuses	1 Γ	VF		unknown	Previous trisomy 21	unknown
Weight in kg	65 I	Diabetes		NO	Pregnancies	unknown
Smoker	NO C	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	e	13+6
PAPP-A	6.8 n	nIU/ml	0.87	Method		CRL (<>Robinson)
fb-hCG	52.8 n	g/ml	2.19	Scan date		19-05-2024
Risks at sampling date				Crown rump l	ength in mm	80
Age Risk			1:914	Nuchal translu	icency MoM	0.73
Biochemical T21 risk			1:632	Nasal bone		PRESENT
Combined trisomy 21 risk 1:3640			Sonographer		DR.DEEPIKA	
Trisomy 13/18 + NT	+ NT <1:10000			Qualifications	in measuring NT	MD
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 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 13/18 (with NT) is <1:10000,			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 3640 women with the same data, there is one woman with a trisomy 21 pregnancy and 3639 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 laboratory cannot be hold responsible for their impact on the			
which indicates a low risk	/1U1 1N 1 / 1	· 1110000 ,	risk assessment! Calculated risks have no diagnostic values			