

which indicates a low risk

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Date of Report 03-07-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. VIJAY LAXMI			Patient ID		12407020157
Birthday		15-07-1996				11860891
Age at Sample date 28.0				Sample Date		02-07-2024
Gestational age 13+6						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	13+5
PAPP-A	6.8	mIU/ml	0.77	Method		CRL (<>Robinson)
fb-hCG	55.6	55.6 ng/ml 2.5		Scan date		01-07-2024
Risks at sampling date				Crown rump length in mm 78.6		
Age Risk 1:834			Nuchal translucency MoM 0.75			
Biochemical T21 risk 1:422			1:422	Nasal bone PRESEN		PRESENT
Combined trisomy 21 risk 1:2510			1:2510	Sonographer D		DR.
Trisomy 13/18 + NT <1:10000			Qualifications	in measuring NT	MBBS	
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 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 2510 women with the same data, there is one woman with a trisomy 21 pregnancy and 2509 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		
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		