

which indicates a low risk

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Date of Report 26-11-2024

				PRISCA	5.2.0.13
Patient Data				TRISCA	0.2.0.10
Name	MR	MRS. LAKSHMI			12411250132
Birthday		14-08-1994			11981171
Age at Sample date	ge at Sample date 30.8				25-11-2024
Gestational age		13+1			
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 ag	e	13+1
PAPP-A	4.8 mIU/ml	0.57	Method		CRL (<>Robinson)
fb-hCG	31.5 ng/ml	0.94	Scan date		25-11-2024
Risks at sampling date			Crown rump length in mm 70.1		
Age Risk 1:637			Nuchal translucency MoM 0.63		
Biochemical T21 risk	ochemical T21 risk 1:1197		Nasal bone PRESEN		PRESENT
Combined trisomy 21 ris	sk	1:6839	Sonographer		DR. SANJEEV KUMAR
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Cut off			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 6839 women with the same data, there is one woman with a trisomy 21 pregnancy and 6838 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the		
1:1000 1:10000 13 15 17 19 21 23 25 Trisomy 13/18+NT The calculated risk for Tri			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		