

which indicates a low risk

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Date of Report 05-12-2024

					PRISCA	5.2.0.13
Patient Data						
Name		MRS. RITU VERMA				12412030033
Birthday			06-05-1998	Sample ID		11895018
Age at Sample date 31.6				Sample Date		03-12-2024
Gestational age 13+5				5		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	65	65 Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian	ı	
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	13+2
PAPP-A	7.3	mIU/ml	0.97	Method		CRL (<>Robinson)
fb-hCG	17.5	ng/ml	0.69	Scan date		30-11-2024
Risks at sampling date				Crown rump length in mm 70.4		
Age Risk 1:540			1:540	Nuchal translucency MoM 0.79		
Biochemical T21 risk			1:7098	Nasal bone prese		present
Combined trisomy 21 ri	sk		<1:10000	Sonographer		DR.
Trisomy 13/18 + NT <1:10000			<1:10000	Qualifications in measuring NT MBH		MBBS
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  1:1000  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 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		