

which indicates a low risk

\*Free Home Sample Collection 9999 778 778



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Date of Report 11-08-2024

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. AARZOO YADAV			Patient ID		12408100189
Birthday	11-12-2001			Sample ID		11870743
Age at Sample date 22.7				Sample Date		10-08-2024
Gestational age			13+5			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	e	13+4
PAPP-A	7.3	mIU/ml	0.88	Method		CRL (<>Robinson)
fb-hCG	27.5	27.5 ng/ml 1.06		Scan date		09-08-2024
Risks at sampling date				Crown rump length in mm 75.6		
Age Risk	Risk 1:108			Nuchal translucency MoM 0.59		
Biochemical T21 risk			1:4537	Nasal bone PRESE		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer DR. DHRUV TANK		DR. DHRUV TANEJA
Trisomy 13/18 + NT			<1:10000	Qualifications in measuring NT MI		MBBS
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.  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)		
Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		