

which indicates a low risk

*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report 06-07-2024

					PRISCA	5.2.0.13
Patient Data						
Name		MRS. SAUMYA				12407050175
Birthday	rthday 10-02-1998			Sample ID		11903847
Age at Sample date 36.4				Sample Date		05-07-2024
Gestational age 13+6						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	75.8	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	13+2
PAPP-A	6.8	mIU/ml	1.04	Method		CRL (<>Robinson)
fb-hCG	32.4	ng/ml	1.41	Scan date		01-07-2024
Risks at sampling date				Crown rump length in mm 72.2		
Age Risk 1:207			Nuchal translucency MoM 1.00			
Biochemical T21 risk			1:643	Nasal bone		PRESENT
Combined trisomy 21 risk 1:2516			1:2516	Sonographer DR. MANE		DR. MANEESH
Trisomy 13/18 + NT <1:10000				Qualifications in measuring NT .		
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 2516 women with the same data, there is one woman with a trisomy 21 pregnancy and 2515 women with not affected pregnancies.		
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		