

which indicates a low risk

*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report 05-08-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. MONIKA SRIVASTAV			Patient ID		12408040175
Birthday		14-06-1991				11863768
Age at Sample date 33.1				Sample Date		04-08-2024
Gestational age			13+0)		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	69	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	12+6
PAPP-A	4.62 1	mIU/ml	0.84	Method		CRL (<>Robinson)
fb-hCG	27.1 1	ng/ml	0.86	Scan date		03-08-2024
Risks at sampling date				Crown rump length in mm 65.2		
Age Risk			1:403	Nuchal translucency MoM		0.70
Biochemical T21 risk			1:2370	Nasal bone		PRESENT
Combined trisomy 21 ris	sk		<1:10000	Sonographer		DR. INDRAJEET
Trisomy 13/18 + NT <1:1000			<1:10000	Qualifications in measuring NT MI		
Risk				Down's Syndr	ome Risk (Trisomy 21	Screening)
1:100				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 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				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, which indicates a low risk				The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		