

Trisomy 13/18+NT

which indicates a low risk

The calculated risk for Trisomy 13/18 (with NT) is 1:5873,

\*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

The laboratory cannot be hold responsible for their impact on the

risk assessment! Calculated risks have no diagnostic values

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. ANSHU YADAV			Patient ID		12408040129
Birthday	05-07-1992			Sample ID		11860607
Age at Sample date 32.1				Sample Date		04-08-2024
Gestational age 12+5						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	44	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+5
PAPP-A	3.1	mIU/ml	0.37	Method		CRL (<>Robinson)
fb-hCG	19.4	ng/ml	0.49	Scan date		04-08-2024
Risks at sampling date				Crown rump l	ength in mm	63.2
Age Risk			1:481	Nuchal translu	cency MoM	0.85
Biochemical T21 risk			1:1075	Nasal bone		PRESENT
Combined trisomy 21 risk			1:6142	Sonographer		DR. NAMIT
Trisomy 13/18 + NT 1:5873			Qualifications	in measuring NT	MD	
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000				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 6142 women with the same data, there is one woman with a trisomy 21 pregnancy and 6141 women with not affected pregnancies. The PAPA-A level is low.  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;		
10 10 11 10 2 20 20 21 20 01 00 00 01 00 11 10 10 11				1998).		