

*Free Home Sample Collection 9999 778 778

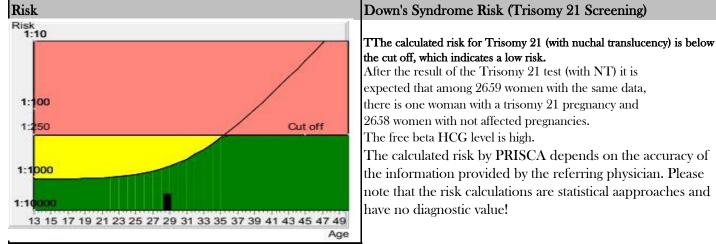


Book a Test Online www.molq.in

Date of Report 21/08/19 5.0.2.37 PRISCA

					TMSCA	5.0.2.07	
Patient Data							
Name	Ms Jyoti		Patient ID		011908190351		
Birthday		25/11/1990		Sample ID		10598034	
Age at delivery		28.7		Sample Date		19/08/19	
Gestational age	12+1						
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	75	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	arameter Value		Gestational age	12+0	
PAPP-A	APP-A 4.36 mIU/ml		Method	CRL (<>Robinson)	
b-hCG 135.4 ng/ml		2.97 Scan Date		19/08/2019	
Risks at sampling date	;		Crown Rump Length (mm)	54.3	
Age Risk		1:733	Nuchal translucency MoM	0.97	
Biochemical Trisomy 21 Risk		1:655	Nasal Bone	present	
Combined Trisomy 21 Risk		1:2659	Sonographer	DR.RUBY RAHUL	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	CON. RADIOLOGIT	



After the result of the Trisomy 21 test (with NT) it is expected that among 2659 women with the same data, there is one woman with a trisomy 21 pregnancy and 2658 women with not affected pregnancies.

The free beta HCG level is high.

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!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

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