

*Free Home Sample Collection 9999 778 778



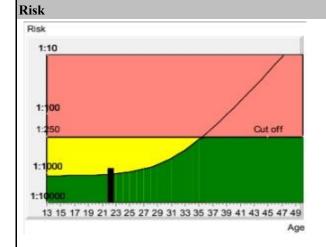
Book a Test Online www.molq.in

Date of Report 13-01-19
PRISCA 5 0 2 37

				_	PRISCA		5.0.2.37
Patient Data							
Name		Mrs PRI	EETI YADAV	Patient ID			011901110108
Birthday			10-10-96	Sample ID			10396789
Age at delivery			22.3	Sample Date			11/01/2019
Gestational age			12+5				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21		unknown
Weight in kg		Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			
Biochemical Data				Ultrasound D	ata		
Parameter	Value		Corr Mom	Gestational ago	e	12+5	

			\mathcal{E}	
PAPP-A	3.55 mIU/ml	0.91	Method	CRL (<>Hadlock)
fb-hCG	95.36 ng/ml	2.17	Scan Date	27-11-18
Risks at sampling date			CRL	
Age Risk		1:1056	Nuchal translucency MoM	
Biochemical T21 Risk			Nasal Bone	Present
Combined Trisomy 21 Ris	k	1:826	Sonographer	

<1:10000



Trisomy 13/18 + NT

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

Down's Syndrome Risk (Trisomy 21 Screening)
The calculated risk for Trisomy 21 (with nuchal
translucency) is below the cut off, which indicates a
low risk.

Qualifications in measuring NT

After the result of the Trisomy 21 test (with NT) it is expected that among more than 826 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 risk calculations are statistical approach and have no diagnostic values!

The patient combined risk presumes the NT measurement was done according to accepted guidelines. The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!