

*Free Home Sample Collection 9999 778 778

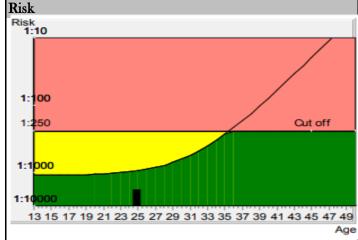


Book a Test Online www.molq.in

Date of Report 28/10/19 PRISCA 5.0.2.37

			PRISCA		5.0.2.37	
Patient Data						
Name		Mrs	Renu Mishra	Patient ID		011910260060
Birthday		1/1/1995		Sample ID		10530432
Age at delivery		24.8		Sample Date		26/10/19
Gestational age			12 + 2			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	67.4	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data			Ultrasound Data			

Parameter Value		Corr Mom	Gestational age	12+2	
PAPP-A	3.65 mIU/ml	1.02	Method	CRL (<>Robinson)	
fb-hCG	23.6 ng/ml	0.51	Scan Date	26/10/19	
Risks at sampling date			Crown Rump Length (mm)	57.6	
Age Risk		1:961	Nuchal translucency MoM	0.70	
Biochemical Trisomy 2	1 Risk	<1:10000	Nasal Bone	present	
Combined Trisomy 21	Risk	<1:10000	Sonographer	DR.AKSHAY SHUKLA	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	C/R	



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.

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

Trisomy 13/18 + NT

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

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