

*Free Home Sample Collection 9999 778 778



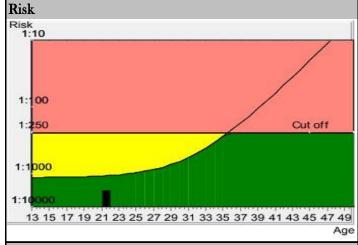
Book a Test Online www.molq.in

 Date of Report
 13/12/19

 PRISCA
 5.0.2.37

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Patient Data						
Name			Mrs Pooja	Patient ID		061912120006
Birthday			10/5/1998	Sample ID		60001232
Age at delivery			21.6	Sample Date		12/12/19
Gestational age			13+2			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	45	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound Da	ata	
Parameter	Value		Corr Mom	Cestational age	2	19+6

Diocricifficat Data			Citasound Data		
Parameter	ameter Value		Gestational age	12+6	
PAPP-A	5.12 mIU/ml	0.63	Method	CRL (<>Robinson)	
fb-hCG	21.5 ng/ml	0.45	Scan Date	9/12/2019	
Risks at sampling date			Crown Rump Length (mm)	65	
Age Risk		1:1090	Nuchal translucency MoM	0.78	
Biochemical Trisomy 21 Risk		<1:10000	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. ROHIT CHANDAK	
Trisomy 13/18 + N T		<1:10000	Qualification in measuring NT	MBBS, MD	



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

The calculated risk for trisomy 13/18 (with nuchal translucency) is

Trisomy 13/18 + NT

<1:10000, which represents a low risk.

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