

*Free Home Sample Collection 9999 778 778

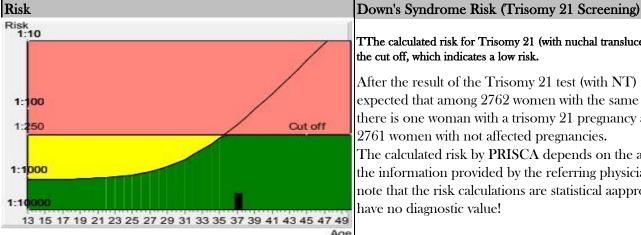


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Date of Report 6/5/2019 PRISCA 50937

			PRISCA		3.0.2.37	
Patient Data						
Name			Mrs Geeta	Patient ID		011905050037
Birthday			22/03/1982	Sample ID		10479847
Age at delivery		37.1		Sample Date		05/04/19
Gestational age			12+6			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59	Diabetes		no	Pregnancies	unknown
Smoker	no	Origin		Asian		

Biochemical Data			Ultrasound Data		
Parameter	meter Value		Gestational age	12+6	
PAPP-A	3.65 mIU/ml	0.71	Method	CRL (<>Robinson)	
fb-hCG	28.3 ng/ml	0.63	Scan Date	5/5/2019	
Risks at sampling date	:		Crown Rump Length (mm)	64.8	
Age Risk		1:168	Nuchal translucency MoM	0.78	
Biochemical Trisomy 21 Risk		1:1286	Nasal Bone	absent	
Combined Trisomy 21 Risk		1:2762	Sonographer	DR.RUBY RAHUL	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	CON. RADIOLOGIT	



TThe 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 2762 women with the same data, there is one woman with a trisomy 21 pregnancy and 2761 women with not affected pregnancies.

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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