

*Free Home Sample Collection 9999 778 778

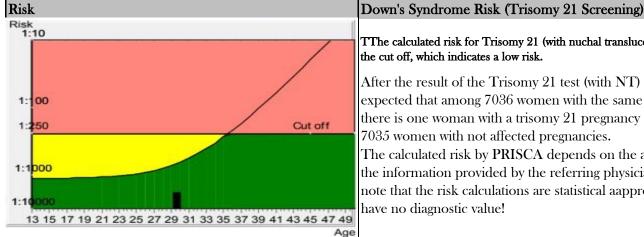


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Date of Report 7/12/2019 PRISCA 5.0.2.37

					IMSCA	5.0.2.37	
Patient Data							
Name		Mrs Neha		Patient ID		011907110175	
Birthday		11/22/1989		Sample ID		10629509	
Age at delivery		29.6		Sample Date		11/07/19	
Gestational age	12+5						
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	66	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	rameter Value		Gestational age	12+2	
PAPP-A	3.86 mIU/ml	0.9	Method	CRL (<>Robinson)	
fb-hCG	66.5 ng/ml	1.50	Scan Date	7/8/2019	
Risks at sampling dat	ie e		Crown Rump Length (mm)	58	
Age Risk		1:679	Nuchal translucency MoM	0.66	
Biochemical Trisomy 21 Risk		1:1298	Nasal Bone	present	
Combined Trisomy 2	21 Risk	1:7036	Sonographer	DR.NIDHI AGARWAL	
Trisomy 13/18 + N T		<1:10000	Qualification in measuring NT	MD	



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 7036 women with the same data, there is one woman with a trisomy 21 pregnancy and 7035 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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