

*Free Home Sample Collection 9999 778 778

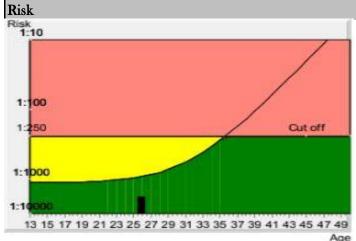


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Date of Report 18/5/2019 DDICCA 50927

				PRISCA		5.0.2.37	
Patient Data							
Name		I	Mrs PREETI	Patient ID		011905160164	
Birthday		20/7/1993		Sample ID		10631951	
Age at delivery		25.8		Sample Date		16/05/2019	
Gestational age			13+1				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	74	Diabetes		no	Pregnancies	unknown	
Smoker	no	Origin		Asian			
Biochemical Data			Ultrasound Da	nta.			

Biochemical Data			Ultrasound Data		
Parameter	rameter Value		Gestational age	13+0	
PAPP-A	5.19 mIU/ml	1.2	Method	CRL (<>Robinson)	
fb-hCG	60.7 ng/ml	1.49	Scan Date	16/05/2019	
Risks at sampling date			Crown Rump Length (mm)	67.2	
Age Risk		1:943	Nuchal translucency MoM	1.02	
Biochemical Trisomy 21 Risk		1:3356	Nasal Bone	present	
Combined Trisomy 21	Risk	<1:10000	Sonographer	DR.ANKIT BHARGAVA	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



After the result of the Trisomy 21 test (with NT) it is

the cut off, which indicates a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

expected that among 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies.

TThe calculated risk for Trisomy 21 (with nuchal translucency) is below

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 approaches 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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