

*Free Home Sample Collection 9999 778 778

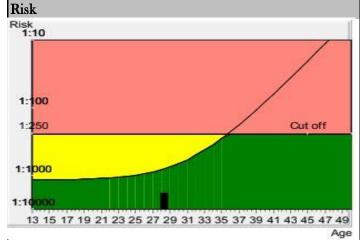


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Date of Report 11/9/2019 PRISCA 5.0.2.37

					TRISCIT	0.0.2.07	
Patient Data							
Name		Mrs Shruti Sahu		Patient ID		011909090286	
Birthday		17/05/1991		Sample ID		10541793	
Age at delivery		28.3		Sample Date		09/09/19	
Gestational age			13+6				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	60	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	arameter Value		Gestational age	13+3	
PAPP-A	3.56 mIU/ml	0.51	Method	CRL (<>Robinson)	
fb-hCG	26.3 ng/ml	0.66	Scan Date	7/9/2019	
Risks at sampling date			Crown Rump Length (mm)	72.5	
Age Risk		1:808	Nuchal translucency MoM	0.52	
Biochemical Trisomy 21 Risk		1:2405	Nasal Bone	present	
Combined Trisomy 21	Risk	<1:10000	Sonographer	DR. A. DHANADIA	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



Trisomy 13/18 + NT

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

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

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