

*Free Home Sample Collection 9999 778 778



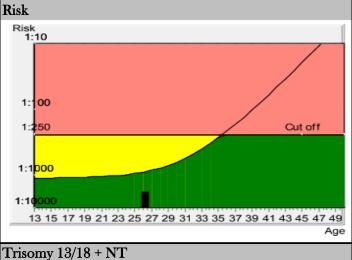
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 Date of Report
 28/10/19

 PRISCA
 5.0.2.37

					TRISCIT	0.0.2.07	
Patient Data							
Name		Mrs Priyanka		Patient ID		011910260024	
Birthday		12/8/1993		Sample ID		10551494	
Age at delivery		26.2		Sample Date		26/10/19	
Gestational age			12+3				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	57	Diabetes		no Pregnan			
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	nrameter Value		Gestational age	11+5	
PAPP-A	5.12 mIU/ml	0.6	Method	CRL (<>Robinson)	
fb-hCG	131.2 ng/ml	1.26	Scan Date	21/10/19	
Risks at sampling date			Crown Rump Length (mm)	50.6	
Age Risk		1:901	Nuchal translucency MoM	0.37	
Biochemical T21 Risk		1:989	Nasal Bone	Present	
Combined Trisomy 21 Risk	·	1:5815	Sonographer	Dr. Divya Agarwal	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	C/R	



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 5815 women with the same data, there is one woman with a trisomy 21 pregnancy and 5814 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!

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

low risk.

The calculated risk for trisomy 13/18 is <1:10000, which represents a

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