

*Free Home Sample Collection 9999 778 778



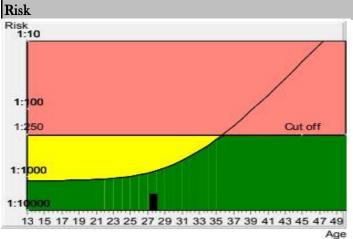
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 Date of Report
 22/08/19

 PRISCA
 5.0.2.37

					TRISCIT	3.0.2.07	
Patient Data							
Name		Mrs Vaishnavi		Patient ID		011908200200	
Birthday		31/12/1991		Sample ID		10574902	
Age at delivery		27.6		Sample Date		20/08/19	
Gestational age			13+5				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	47	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	nrameter Value		Gestational age	13+1	
PAPP-A	3.91 mIU/ml		Method	CRL (<>Robinson)	
fb-hCG	29.15 ng/ml	0.66	Scan Date	17/08/19	
Risks at sampling date	2		Crown Rump Length (mm)	69.5	
Age Risk		1:852	Nuchal translucency MoM	0.63	
Biochemical Trisomy	21 Risk	1:1706	Nasal Bone	present	
Combined Trisomy 21 Risk		1:9916	Sonographer	DR. P.GROVER	
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 9916 women with the same data, there is one woman with a trisomy 21 pregnancy and 9915 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

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