

*Free Home Sample Collection 9999 778 778

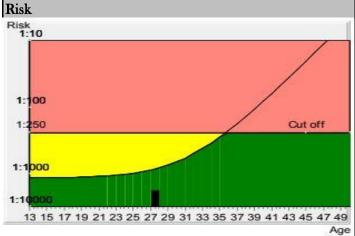


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Date of Report 16/07/19
PRISCA 5.0.2.37

					TMSCA	5.0.2.37	
Patient Data							
Name		Mrs Radha		Patient ID		011907150123	
Birthday		16/01/1992		Sample ID		10452185	
Age at delivery		27.5		Sample Date		15/07/19	
Gestational age		13+6					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	68	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	ameter Value		Gestational age	13+6	
PAPP-A	$5.96~\mathrm{mIU/ml}$	0.98	Method	CRL (<>Robinson)	
fb-hCG	21.5 ng/ml	0.56	Scan Date	15/07/19	
Risks at sampling date			Crown Rump Length (mm)	80.4	
Age Risk		1:866	Nuchal translucency MoM	0.78	
Biochemical Trisomy 21 Risk		<1:10000	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR.VARUN RAJ	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	DMC	



Down's Syndrome Risk (Trisomy 21 Screening)

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 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!

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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