

*Free Home Sample Collection 9999 778 778

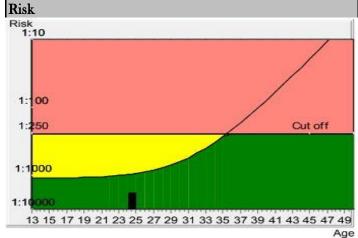


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Date of Report 6/12/2019 PRISCA 5.0.2.37

					FMSCA	3.0.2.37	
Patient Data							
Name		Mrs Rashmi		Patient ID		011912050153	
Birthday		5/6/1995		Sample ID		10507426	
Age at delivery		24.5		Sample Date		05/12/19	
Gestational age		12+1					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	75	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	arameter Value		Gestational age	12+0	
PAPP-A	$1.69~\mathrm{mIU/ml}$	0.57	Method	CRL (<>Robinson)	
fb-hCG	23.4 ng/ml	0.51	Scan Date	4/12/2019	
Risks at sampling date			Crown Rump Length (mm)	56	
Age Risk		1:968	Nuchal translucency MoM	1.15	
Biochemical Trisomy 21 Risk		1:6399	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. RAKHI BANSAL	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MBBS, DMRD	



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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