

*Free Home Sample Collection 9999 778 778

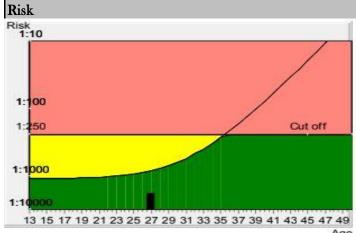


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Date of Report 10/9/2019 PRISCA 5.0.2.37

					FMSCA	3.0.2.37	
Patient Data							
Name		Mrs Deepika		Patient ID		011909080248	
Birthday		11/10/1992		Sample ID		10477041	
Age at delivery		26.9		Sample Date		08/09/19	
Gestational age		12+1					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	63	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter Value		Corr Mom	Gestational age	11+6	
PAPP-A	$1.61 \mathrm{mIU/ml}$	0.44	Method	CRL (<>Robinson)	
fb-hCG	33.43 ng/ml	0.70	Scan Date	7/9/2019	
Risks at sampling date			Crown Rump Length (mm)	43	
Age Risk		1:853	Nuchal translucency MoM	0.58	
Biochemical Trisomy 2	21 Risk	1:1517	Nasal Bone	present	
Combined Trisomy 21	Risk	1:8892	Sonographer	Dr. Shruti Sangwan	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



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The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

After the result of the Trisomy 21 test (with NT) it is expected that among 8892 women with the same data, there is one woman with a trisomy 21 pregnancy and 8891 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!

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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