

*Free Home Sample Collection 9999 778 778

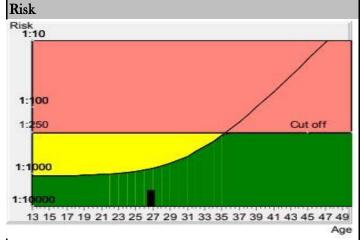


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Date of Report 22/11/19
PRISCA 5.0.2.37

					FMSCA	3.0.2.37	
Patient Data							
Name		Mrs Shweta		Patient ID		011911210131	
Birthday		18/02/1993		Sample ID		10607437	
Age at delivery		26.8		Sample Date		21/11/1	
Gestational age		12+0					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	68.8	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
arameter Value		Corr Mom	Gestational age	11+6	
PAPP-A	2-A 3.61 mIU/ml		Method	CRL (<>Robinson)	
b-hCG 66.08 ng/ml		1.39	Scan Date	20/11/19	
Risks at sampling date			Crown Rump Length (mm)	52	
Age Risk		1:857	Nuchal translucency MoM	0.65	
Biochemical Trisomy 21 Risk		1:3362	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. PRAKASH LAL CHANDANI	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



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!

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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