

\*Free Home Sample Collection 9999 778 778

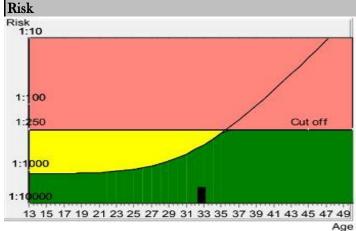


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Date of Report 10/12/2019 PRISCA 5.0.2.37

					TMSCA	3.0.2.37	
Patient Data							
Name		Mrs Shazia		Patient ID		011912090003	
Birthday		9/3/1987		Sample ID		10530232	
Age at delivery		32.8		Sample Date		09/12/19	
Gestational age		13+0					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	64.8	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	neter Value		Gestational age	12+6	
PAPP-A	3.75  mIU/ml	0.77	Method	CRL (<>Robinson)	
fb-hCG	39.2 ng/ml	0.91	Scan Date	8/12/2019	
Risks at sampling da	ute		Crown Rump Length (mm)	64.4	
Age Risk		1:433	Nuchal translucency MoM	1.10	
Biochemical Trisomy 21 Risk		1:1860	Nasal Bone	present	
Combined Trisomy 21 Risk		1:5910	Sonographer	DR VARUN SHARMA	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT		



## 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 5910 women with the same data, there is one woman with a trisomy 21 pregnancy and 5909 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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