

*Free Home Sample Collection 9999 778 778

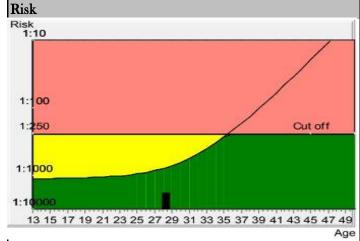


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Date of Report 10/11/2019
PRISCA 5 0 9 37

				5.0.2.37			
Patient Data							
Name	Mrs. Preeti		Patient ID		011911080286		
Birthday		13/07/1991		Sample ID		10525764	
Age at delivery		28.2		Sample Date		08/11/19	
Gestational age	12+3						
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	65	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	ameter Value		Gestational age	12+2	
PAPP-A	$4.22~\mathrm{mIU/ml}$	1.07	Method	CRL (<>Robinson)	
fb-hCG	70.8 ng/ml	1.54	Scan Date	8/11/2019	
Risks at sampling date			Crown Rump Length (mm)	58.1	
Age Risk		1:770	Nuchal translucency MoM	0.66	
Biochemical Trisomy 21 Risk		1:2016	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	Dr. B.A. WANI	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



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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