

*Free Home Sample Collection 9999 778 778

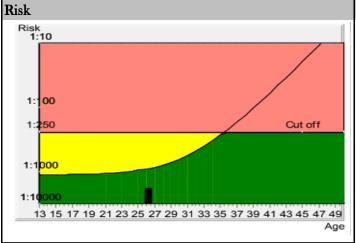


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Date of Report 28/10/19 PRISCA 5.0.2.37

					TMSCA	5.0.2.57	
Patient Data							
Name		Mrs Priyanka		Patient ID		011910260024	
Birthday		12/8/1993		Sample ID		10551494	
Age at delivery		26.2		Sample Date		26/10/19	
Gestational age		12+4					
Correction factors							
Fetuses	2	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	57	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	11+6	
PAPP-A	$5.12~\mathrm{mIU/ml}$	0.57	Method	CRL (<>Robinson)	
fb-hCG	131.2 ng/ml	1.28	Scan Date	21/10/19	
Risks at sampling date			Crown Rump Length (mm)	51.6	
Age Risk		1:906	Nuchal translucency MoM	0.58	
Biochemical T21 Risk		1:836	Nasal Bone	Present	
Combined Trisomy 21 Risk		1:4998	Sonographer	Dr Divya Agarwal	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	C/R	



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 4998 women with the same data, there is one woman with a trisomy 21 pregnancy and 4997 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!

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 is <1:10000, which represents a low risk.

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