

*Free Home Sample Collection 9999 778 778

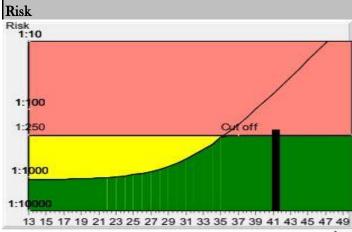


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Date of Report 19/11/19

			PRISCA		5.0.2.37		
Patient Data							
Name		Mrs Prabhjot Kaur F1		Patient ID		051911180030	
Birthday			10/8/1978	Sample ID		10536558	
Age at delivery			41.3	Sample Date		18/11/19	
Gestational age			11+6				
Correction factors							
Fetuses	2	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	56.7	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			
Biochemical Data			Ultrasound Da	ata			

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Parameter	rameter Value		Gestational age	11+6	
PAPP-A	A 8.65 mIU/ml		Method	CRL (<>Robinson)	
fb-hCG	283.5 ng/ml	2.55	Scan Date	18/11/19	
Risks at sampling date			Crown Rump Length (mm)	51.2	
Age Risk		1:54	Nuchal translucency MoM	1.02	
Biochemical Trisomy 2	21 Risk	1:54	Nasal Bone	present	
Combined Trisomy 21 Risk		1:201	Sonographer	DR. PAWAN JOON	
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 above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 201 women with the same data, there is one woman with a trisomy 21 pregnancy and 200 women with not affected pregnancies.

The free beta HCG level is high.

The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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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