

*Free Home Sample Collection 9999 778 778

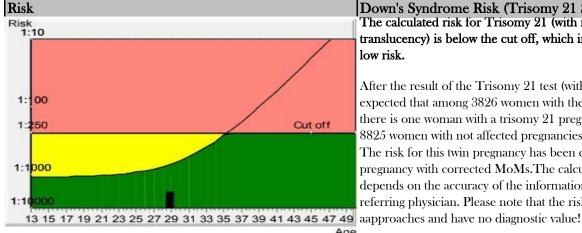


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Date of Report 17/12/19 PRISCA 5 0 9 37

			PRISCA		
Patient Data					
Name	Mrs Lalita l	Fetus B Patient ID		011912140199	
Birthday	4/3	3/1991 Sample ID		10303062	
Age at delivery		28.8 Sample Date		14/12/19	
Gestational age		11+6			
Correction factors					
Fetuses	2 IVF	unknown	Previous trisomy 21	unknown	
Weight in kg	75 Diabetes	no	Pregnancies		
Smoker	no Origin	Asian			

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	11+5	
PAPP-A	3.54 mIU/ml	0.72	Method	CRL (<>Robinson)	
fb-hCG	112.5 ng/ml	1.10	Scan Date	14/12/19	
Risks at sampling date			Crown Rump Length (mm)	45	
Age Risk		1:721		1.21	
Overall Population Risk		1:1699	Nasal Bone	present	
Trisomy 21 Risk		1:3826	Sonographer	DR. VIKASH GOYAL	
Trisomy 13/18		<1:10000	Qualification in measuring NT	DMRD	



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 3826 women with the same data, there is one woman with a trisomy 21 pregnancy and 8825 women with not affected pregnancies. 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

Trisomy 13/18 + NT The calculated risk for trisomy 13/18 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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