

*Free Home Sample Collection 9999 778 778

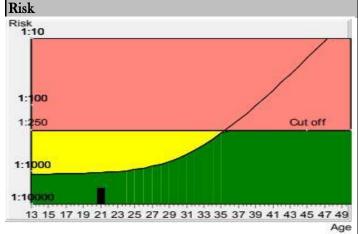


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Date of Report 17/10/19
PRISCA 5.0.2.37

					3.0.2.37		
Patient Data							
Name		Mrs Pooja		Patient ID		011910150193	
Birthday		30/09/1998		Sample ID		10554637	
Age at delivery		21		Sample Date		15/10/1	
Gestational age		12+3					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	34	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	meter Value		Gestational age	12+1	
PAPP-A	4.12 mIU/ml	0.52	Method	CRL (<>Robinson)	
fb-hCG	24.6 ng/ml	0.42	Scan Date	14/10/19	
Risks at sampling date			Crown Rump Length (mm)	55.1	
Age Risk		1:1069	Nuchal translucency MoM	1.10	
Biochemical Trisomy 21 Risk		1:7872	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. RUBY RAHUL	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	CON. RADIOLOGIST	



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!

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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