

*Free Home Sample Collection 9999 778 778

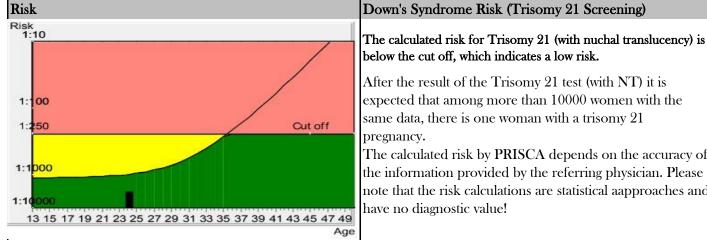


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03/01/2020 Date of Report PRISCA

					TRISCA	5.0.2.07	
Patient Data							
Name		Mrs Indu		Patient ID		052001020012	
Birthday		26/11/1995		Sample ID		10501429	
Age at delivery		24.1		Sample Date		02/01/2020	
Gestational age		12+3					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	56	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	meter Value		Gestational age	11+5	
PAPP-A	$3.21~\mathrm{mIU/ml}$	0.68	Method	CRL (<>Robinson)	
fb-hCG	36.8 ng/ml	0.76	Scan Date	28/12/2019	
Risks at sampling date			Crown Rump Length (mm)	49.3	
Age Risk		1:993	Nuchal translucency MoM	0.90	
Biochemical Trisomy 21 Risk		1:4645	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. KRISHNAN JAIN	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



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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