*Free Home Sample Collection 9999 778 778

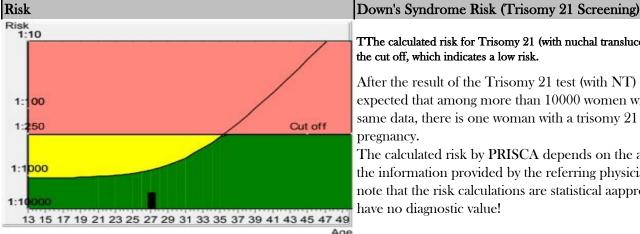


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Date of Report 7/6/2019 PRISCA 5.0.2.37

					TRISCIT	0.0.2.07	
Patient Data							
Name		Mrs Narmada Swain		Patient ID		021907050002	
Birthday		5/20/1992		Sample ID		10480235	
Age at delivery		27.1		Sample Date		05/06/2019	
Gestational age			12+0				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	55	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	ameter Value		Gestational age	12+0	
PAPP-A	$3.25~\mathrm{mIU/ml}$	0.8	Method	CRL (<>Robinson)	
fb-hCG	27.1 ng/ml	0.53	Scan Date	7/5/2019	
Risks at sampling date			Crown Rump Length (mm)	53.6	
Age Risk		1:836	Nuchal translucency MoM	0.7	
Biochemical Trisomy 21 Risk		<1:10000	Nasal Bone	present	
Combined Trisomy 21	Risk	<1:10000	Sonographer	DR.VENKAT REDDY M	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



TThe 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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