

*Free Home Sample Collection 9999 778 778

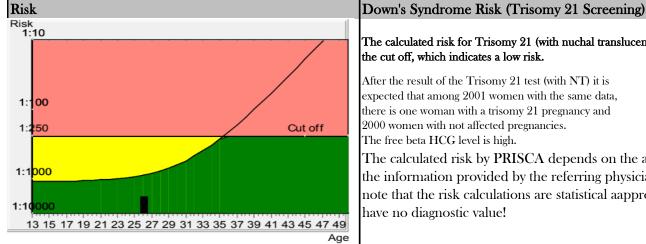


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12/02/2020 Date of Report PRISCA

				TRISCA	5.0.2.07
Patient Data					
Name]	Mrs Ambadi Mamtha	Patient ID		012002100194
Birthday		25/12/1993	Sample ID		10673320
Age at delivery		26.1			10/02/2020
Gestational age		12+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52 D iab	petes	no	Pregnancies	
Smoker	no Orig	in	Asian		

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	11+6	
PAPP-A	3.37 mIU/ml	0.78	Method	CRL (<>Robinson)	
fb-hCG	131.25 ng/ml	2.52	Scan Date	10/2/2020	
Risks at sampling date			Crown Rump Length (mm)	51.2	
Age Risk		1:891	Nuchal translucency MoM	0.80	
Biochemical Trisomy 21 Risk		1:334	Nasal Bone	present	
Combined Trisomy 21 Risk		1:2001	Sonographer	DR. VIKAS GOYAL	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	DMRD	



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 2001 women with the same data, there is one woman with a trisomy 21 pregnancy and 2000 women with not affected pregnancies.

The free beta HCG level is high.

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