

*Free Home Sample Collection 9999 778 778

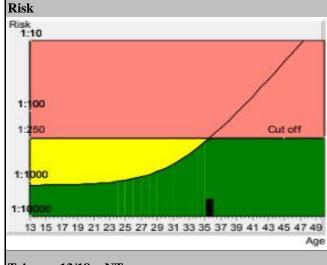


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Date of Report 22-12-18 PRISCA 50237

				PRISCA	3.0.2.37
Patient Data					
Name		Mrs Arti Katy	al Patient ID		011812210102
Birthday		05-06-8	33 Sample ID		10151397
Age at delivery		35.5 Sample Date			21/12/2018
Gestational age		13+	-2		
Correction factors					
Fetuses	1	IVF	Yes	Previous trisomy 21	unknown
Weight in kg	62	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		
Biochemical Data			Ultrasound D	ata	

Diochemical Data			Citi asouna Data		
Parameter	Value	Corr Mom	Gestational age	13+1	
PAPP-A	2.44 mIU/ml	0.51	Method	CRL (<>Hadlock)	
fb-hCG	35.41 ng/ml	0.87	Scan Date	20-12-18	
Risks at sampling date			CRL	67.9	
Age Risk		1:248	Nuchal translucency MoM	0.81	
Biochemical T21 Risk		1:406	Nasal Bone	Present	
Combined Trisomy 21 Risl	ζ	1:2353	Sonographer	DR.ANANT ANUPAM	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS, DMRD	



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

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 2353 women with the same data, there is one woman with a trisomy 21 pregnancy and 2352 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approach and have no diagnostic values!

The patient combined risk presumes the NT measurement was done according to accepted guidelines The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!