

*Free Home Sample Collection 9999 778 778



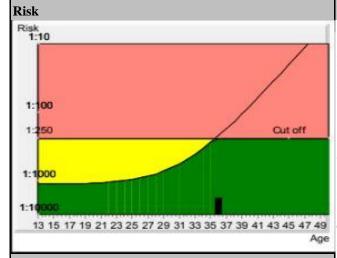
Book a Test Online www.molq.in

Date of Report 05-12-18

					z and or respond	00 12 10
					PRISCA	5.0.2.37
Patient Data						
Name			Mrs Sweety	Patient ID		011812020166
Birthday	25-12-82			Sample ID		10316156
Age at delivery	35.9			Sample Date		02/12/18
Gestational age	13+1					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknowr
Weight in kg	81	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound D	ata	
Parameter	Value		Corr Mom	Gestational ag	e	13+0
PAPP-A	3.14	mIU/ml	0.95	Method		CRL (<>Hadlock)
Biochemical Data Parameter	Value			Ultrasound D Gestational ag	ata	

PAPP-A	3.14 mIU/ml	0.95	Method	CRL (<>Hadlock)
fb-hCG	54.48 ng/ml	1.42	Scan Date	02-12-18
Risks at sampling date			CRL	67
Age Risk		1:255	Nuchal translucency MoM	0.59
Biochemical T21 Risk		1:549	Nasal Bone	Present

Combined Trisomy 21 Risk 1:2914 Sonographer DR.PRAKASH LALCHANDANI Trisomy 13/18 + NT<1:10000 Qualifications in measuring NT



Trisomy 13/18 + NTThe 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 2914 women with the same data, there is one woman with a trisomy 21 pregnancy and 2913 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!