

*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report 30-10-18
PRISCA 5 0 2 37

				PRISCA		5.0.2.37	
Patient Data							
Name Mrs Farhat Nisha			Patient ID		0118	011810290148	
Birthday	(02-01-92	Sample ID			10352078	
Age at delivery		26.8	Sample Date			29/10/18	
Gestational age		12+2					
Correction factors							
Fetuses 1	IVF		unknown	Previous trisomy 21		unknown	
Weight in kg 74.4	Diabetes		unknown	Pregnancies		unknown	
Smoker Unknown	Origin		Asian				
Biochemical Data			Ultrasound Data				
Parameter Value	Corr	Mom	Gestational age	2	12+2		
PAPP-A 1.57	mIU/ml	0.58	Method		CRL (<>Robin	ison)	
fb-hCG 33.19	ng/ml	0.77	Scan Date			29-10-18	
Risks at sampling date			Crown rump le	ength (mm)	58.2		
Age Risk	1:862	2	Nuchal translu	cency MoM	0.66		
Biochemical T21 Risk	1:263	33	Nasal Bone		Present		
Combined Trisomy 21 Risk <1:10000		0000	Sonographer		DR.SAHIL LOOMBA		
Trisomy 13/18 + NT	<1:10	0000	Qualifications	in measuring NT	MBBS,DNB		
Risk			Down's Syndr	ome Risk (Trisomy	21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 with the same data, there is one women 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines.				

The calculated risk for Trisomy 13/18 (with NT) is

<1:10000, which indicates a low risk

Trisomy 13/18 + NT

value!

The laboratory can not be held responsible for their impact on

the risk assessment! Calculated value has no diagnostic