

\*Free Home Sample Collection 9999 778 778

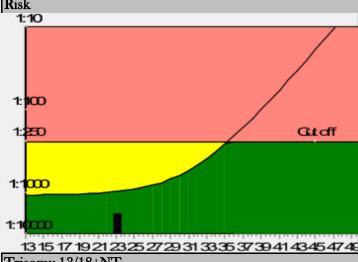


Book a Test Online www.molq.in

Date of Report 05-12-2024

				PRISCA		5.2.0.13
Patient Data						
Name	MRS. LAXMI W/O HITESH			Patient ID		12412030140
Birthday	05-11-2001			Sample ID		11956360
Age at Sample date	23.1			Sample Date		03-12-2024
Gestational age	11+4					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	54.7	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Da	ata	
Parameter	Value		Corr Mom	Gestational ago	e	11+4

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	11+4	
PAPP-A	3.1 mIU/ml	0.76	Method	CRL (<>Robinson)	
fb-hCG	98.5 ng/ml	2.02	Scan date	03-12-2024	
Risks at sampling date			Crown rump length in mm	47.4	
Age Risk		1:993	Nuchal translucency MoM	0.85	
Biochemical T21 risk		1:624	Nasal bone	present	
Combined trisomy 21 risk		1:3502	Sonographer	DR.AMENDA	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Gtoff			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 3502 women with the same data, there is one woman with a trisomy 21 pregnancy and 3501 women with not affected pregnancies.		



## Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

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 that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values