MOLQ Express Lab
*Free Home Sample Collection 9999 778 778
Download "MOLQ" App on
Book a Test Online www.molq.in

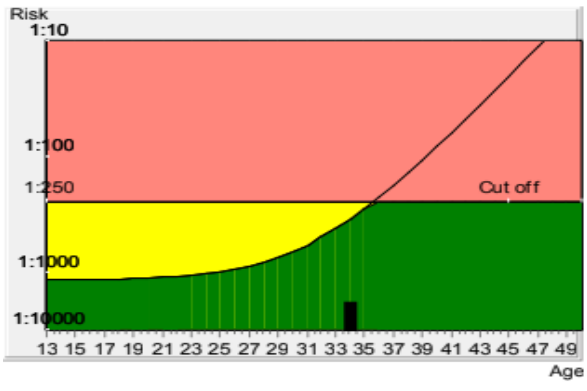
Date of Report	17/3/2023
PRISCA	5.2.0.13

Patient Data			
Name	**MRS. PREETI**	Patient ID	012303160128
Birthday	22/02/1989	Sample ID	11507648
Age at term	34.7	Sample Date	16/3/2023
Gestational age	13+6		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	78	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	**Value**	**Corr Mom**		13+5
PAPP-A	3.89 mIU/ml	0.62	Method	CRL (<>Robinson)
fb-hCG	26.5 ng/ml	1.16	Scan date	15/3/2023
Risks at sampling date			Nuchal translucency	1.5
Age Risk		1:347	Nuchal translucency MoM	0.8
Biochemical T21 risk		1:508	Nasal bone	Present
Combined trisomy 21 risk		1:2918		
Trisomy 13/18 + NT		<1:10000		

Risk



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

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.
After the result of the Trisomy 21with NT test it is expected that among 2918 women with the same data, there is one woman with a trisomy 21 pregnancy and 2917 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 the risk calculations are statistical aapproaches and have no diagnostic value!

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

Risk Above Cut Off Risk above Age Risk Risk below Age risk