

*Free Home Sample Collection 9999 778 778

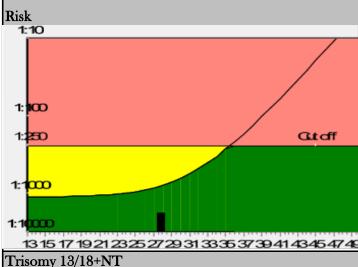


Book a Test Online www.molq.in

Date of Report 92 11 9094

					Date of Report	23-11-2024
					PRISCA	5.2.0.13
Patient Data						
Name	MRS. UPASANA W/O INDRAJ S.			Patient ID		12411220148
Birthday			06-02-1997	Sample ID		11966723
Age at Sample date	27.8			Sample Date		22-11-2024
Gestational age	12+5					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound D	ata	
Parameter	Value		Corr Mom	Gestational ago	e	12+4
PAPP-A	6.9	m I U/ml	1.42	Method		CRL (<>Robinson)

Parameter	Value	Corr Mom	Gestational age	12+4	
PAPP-A	$6.9~\mathrm{mIU/ml}$	1.42	Method	CRL (<>Robinson)	
fb-hCG	38.2 ng/ml	1.13	Scan date	21-11-2024	
Risks at sampling date			Crown rump length in mm	61.5	
Age Risk		1:815	Nuchal translucency MoM	1.01	
Biochemical T21 risk		1:7806	Nasal bone	PRESENT	
Combined trisomy 21 risk		<1:10000	Sonographer	DR.	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		



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 calculated risk for Trisomy 21(with NT) is below the

expected that among more than 10000 women with the same data,

cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is

there is one woman with a trisomy 21 pregnancy.

The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523;

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