

which indicates a low risk

\*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report 26-11-2024

				PRISCA	5.2.0.13
Patient Data					
Name	MRS. ISHIKA V	V/O NARESH	Patient ID		12411250102
Birthday		05-02-2005	Sample ID		11966709
Age at Sample date 19.8		Sample Date		25-11-2024	
Gestational age		13+4	1		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58.7 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	13+4
PAPP-A	6.9 mIU/ml	0.85	Method		CRL (<>Robinson)
fb-hCG	15.7 ng/ml	0.57	Scan date		25-11-2024
Risks at sampling date			Crown rump length in mm 74.5		
Age Risk		1:1128	Nuchal translu	icency MoM	0.92
Biochemical T21 risk		<1:10000	Nasal bone		PRESENT
Combined trisomy 21 risk <1:10000			Sonographer DR. AMENI		DR. AMENDA
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
1:100		Cut off	<b>cut off, which</b> After the result of expected that an	d risk for Trisomy 21(we represents a low risk.) of the Trisomy 21 test (with a nong more than 10000 we man with a trisomy 21 presented.)	th NT) it is omen with the same data,
1:10000 1:10000 13 15 17 19 21 23 25 2 Trisomy 13/18+NT The calculated risk for Tris		41 43 45 47 49	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		