

*Free Home Sample Collection 9999 778 778 Download "MOLQ" App on

Book a Test Online www.molq.in

					Date of Report PRISCA	08-11-2023 5.2.0.13
Patient Data						
Name MRS CHARU SACHDEVA				Patient ID		012311070140
Birthday	nday 25-11-199			Sample ID		11806879
Age at Sample date 29.			29.9	Sample Date		07-11-2023
Gestational age 13+0						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52 Diabetes			NO	Pregnancies	unknown
Smoker	NO Origin			Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+6
PAPP-A	4.93	mIU/ml	0.64	Method		CRL (<>Robinson)
fb-hCG	62.3	ng/ml	1.81	Scan date		06-11-2023
Risks at sampling date				Crown rump length in mm 64.5		
Age Risk			1:661	Nuchal translu	cency MoM	0.67
Biochemical T21 risk		1:368	Nasal bone PRESEN			
Combined trisomy 21 risk			1:2232	Sonographer		DR INDRAJEET
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MD
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				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 2232 women with the same data, there is one woman with a trisomy 21 pregnancy and 2231 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 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		
which indicates a low risk Risk	Above Cu	ıt Off	Risk above Ag	e Risk	Risk below Age risk	