

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

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				Date of Report PRISCA	29-07-2024 5.2.0.13
Patient Data					
Name	MRS. RUCHI JINDAL				12407280172
Birthday	23-02-1991		Sample ID		11905728
Age at Sample date	33.4		Sample Date		28-07-2024
Gestational age		13+0			
Correction factors				1	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	61 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origi	n	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	13+0
PAPP-A	5.7 mIU,	/ml 0.90	Method		CRL (<>Robinson)
fb-hCG	18.3 ng/m	1 0.56	Scan date		28-07-2024
Risks at sampling date			Crown rump length in mm 67.03		
Age Risk 1:382		Nuchal translucency MoM 0.61			
Biochemical T21 risk	1:6381		Nasal bone		present
Combined trisomy 21 risk	<1:10000		Sonographer		DR. INDRAJEET
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:100 1:1000 1:1000 1:10000 1:15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18+NT 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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 risk page no diagnostic values		
which indicates a low risk			risk assessment! Calculated risks have no diagnostic values		



Risk Above Cut Off

