

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

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				Date of Report PRISCA	12-07-2024 5.2.0.13
Patient Data					
Name	MRS. RUHITA				12407110081
Birthday	08-04-1993		Sample ID		11664260
Age at Sample date	31.3		Sample Date		11-07-2024
Gestational age 12+0					
Correction factors				1	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	11+6
PAPP-A	4.3 mIU/n	nl 1.01	Method		CRL (<>Robinson)
fb-hCG	53.2 ng/ml	1.26	Scan date		10-07-2024
Risks at sampling date			Crown rump length in mm 51.		
Age Risk	ge Risk 1:533		Nuchal translucency MoM 0.6		
Biochemical T21 risk	1:2020		Nasal bone		PRESENT
Combined trisomy 21 risk	ombined trisomy 21 risk <1:10000		Sonographer DR. Adi		DR. Aditi Dixit
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT		C/R
Risk 1:10			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:100 1:250 1:1000 1:1000 1:10000 13 15 17 19 21 23 25 27 Trisomy 13/18+NT The calculated risk for Trison which indicates a low risk		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 amog 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 risks have no diagnostic values			



Risk Above Cut Off

