

*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. UPMA			Patient ID		12407280122
Birthday 01-01-1991		Sample ID 1186		11861371	
Age at Sample date 33.6		Sample Date 28-07-20		28-07-2024	
Gestational age		13+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70 Diabet	es	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+3
PAPP-A	6.6 mIU/n	nl 1.01	Method		CRL (<>Robinson)
fb-hCG	19.5 ng/ml	0.75	Scan date		27-07-2024
Risks at sampling date			Crown rump length in mm 72.5		
Age Risk 1:379			Nuchal translucency MoM 0.78		
Biochemical T21 risk	emical T21 risk 1:4527		Nasal bone PRESI		PRESENT
Combined trisomy 21 risk <1:10000		Sonographer DR. VIKASH GO'		DR. VIKASH GOYAL	
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:10 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, 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 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 risks have no diagnostic values		



Risk Above Cut Off

