

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

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				Date of Report PRISCA	03-09-2024 5.2.0.13
Patient Data					
Name MRS. RANJANA PATEL			Patient ID		12409020085
Birthday	thday 09-08-1996			Sample ID 1186048	
Age at Sample date	ge at Sample date 28.1		Sample Date 02-09-2		02-09-2024
Gestational age 13+1					
Correction factors				•	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	51 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	13+0
PAPP-A	6.9 mIU/m	d 0.84	Method		CRL (<>Robinson)
fb-hCG	47.5 ng/ml	1.43	Scan date		01-09-2024
Risks at sampling date			Crown rump length in mm 58.		
Age Risk 1:808			Nuchal translucency MoM 0.65		
Biochemical T21 risk	cal T21 risk 1:1482		Nasal bone P		PRESENT
Combined trisomy 21 risk 1:8119		Sonographer DR. DEEP		DR. DEEPIKA	
Γrisomy 13/18 + NT <1:10000		Qualifications in measuring NT		MD	
Risk 1:10			Down's Syndr	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 Triso 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 8119 women with the same data, there is one woman with a trisomy 21 pregnancy and 8118 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			



Risk Above Cut Off

