

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

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					Date of Report PRISCA	26-04-2021 5.0.2.37
Patient Data						
Name	N	ARS. SH	IV RANJANI	Patient ID		012104190262
Birthday			23-06-1994	Sample ID		10883741
Age at delivery			26.8	Sample Date		19/04/2021
Gestational age			13+3			
Correction factors						
Fetuses	1 I	VF		unknown	Previous trisomy 21	unknown
Weight in kg	58.3 I	Diabetes		No	Pregnancies	unknown
Smoker	No C	Drigin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	13+3
PAPP-A	4.58 n	nIU/ml	0.72	Method		CRL(<>Robinson
fb-hCG	52.9 n	g/ml	1.25	Scan date		20-04-2021
Risks at sampling date				Crown rump length in mm 73.2		
Age Risk			1:896	Nuchal translucency MOM 0.		0.88
Biochemical T21 risk			1:1609	Nasal bone		Present
Combined Trisomy 21 R	lisk		1:8394	Sonographer		Dr Vishakha Mittal
Trisomy 13/18 + NT			<1:10000	Qualification in	n measuring NT	MD
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:10				The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.		
1: 00 1:250 Cut off 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				After the result of the Trisomy 21 test (with NT) it is expected that among 8394 women with the same data, there is one woman with a trisomy 21 pregnancy and 8393 women without 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!		
Age Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk				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		
	Above Cut			Risk above Ag	_	Risk below Age risk