

*Free Home Sample Collection 9999 778 778 Download "MOLQ" App on Book a Test Online www.molq.in

				Date of Report PRISCA	30/5/2023 5.2.0.13
Patient Data					
Name	MRS.N	EHA SINGH	Patient ID		012305290236
Birthday		1/7/1992	Sample ID		11475100
Age at term		31.5	Sample Date		29/5/2023
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	54 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+5
PAPP-A	3.26 mIU/ml	0.49	Method		CRL (<>Robinson)
fb-hCG	25.4 ng/ml	0.69	Scan date		28/5/2023
Risks at sampling date			Crown rump length in mm 55		
Age Risk 1:576		1:576	Nuchal translucency MoM 0.8		
Biochemical T21 risk		1:1442	Nasal bone		Present
Combined trisomy 21 risk 1:7884					
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:10 1:100 1:350 Cut off		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 with NT test it is expected that among 7884 women with the same data, there is one woman with a trisomy 21 pregnancy and 7883 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please			
1:1 0000 13 15 17 19 21 23 25 2 The calculated risk for <1:10000 , which indica	Trisomy 13/18 (with	1	have no diagno The laboratory	y cannot be hold respon essment! Calculated risl	sible for their impact