

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

				Date of Report PRISCA	28/7/2023 5.2.0.13
Patient Data					
Name		MRS.KAJAL	Patient ID		022307270006
Birthday			Sample ID		11792094
Age at term		31.7	Sample Date		27/7/2023
Gestational age		13+0			
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			12+6
PAPP-A	4.62 mIU/ml	0.59	Method		CRL (<>Robinson)
fb-hCG	41.8 ng/ml	1.21	Scan date		26/7/2023
Risks at sampling date			Crown rump length in mm 64		
Age Risk		1:563	Nuchal translucency MoM		1.04
Biochemical T21 risk		1:663	Nasal bone		Present
Combined trisomy 21 ris	k	1:2617			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 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 2617 women with the same data, there is one woman with a trisomy 21 pregnancy and 2616 women with not affected pregnancies.		
1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Risk	Above Cut Off		Risk above Ag	e Risk 🛛 🚺 R	isk below Age risk