

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

Book a Test Online www.molq.in

				Date of Report PRISCA	22/8/2022 5.1.0.17
Patient Data				1100011	
Name	MRS. KO	MAL YADAV	Patient ID		012208210030
Birthday		28.8.1994	Sample ID		11245947
Age at term		28.06	Sample Date		21/8/2022
Gestational age		11+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	11+4
PAPP-A	2.65 mIU/ml	0.75	Method		CRL (<>Robinson)
fb-hCG	42.9 ng/ml	0.92	Scan date		21/8/2022
Risks at sampling date			Nuchal translucency 1.3		
Age Risk	Risk 1:768		Nuchal translucency MoM 0.99		
Biochemical T21 risk		1:3040	Nasal bone		Present
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:100 1:250		Cut off	<b>cut off, which</b> After the resul expected that a same data, the	<b>d risk for Trisomy 21(w.</b> <b>represents a low risk.</b> t of the Trisomy 21with among more than 10000 re is one woman with a t 19999 women with not a	NT test it is ) women with the trisomy 21
1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			The calculated risk by <b>PRISCA</b> 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!		
Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Risk A	Above Cut Off		Risk above Ag	e Risk 🛛 📕 F	Risk below Age risk