

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

Book a Test Online www.molq.in

Patient Data				PRISCA	5.1.0.17
Name		MRS. POOJA	Patient ID		012212290211
Birthday		1/1/1995	Sample ID		11523555
Age at term		28.06	Sample Date		29/12/2022
Gestational age		11+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	50 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			11+6
PAPP-A	4.12 mIU/ml	0.8	Method		CRL (<>Robinson)
fb-hCG	66.4 ng/ml	1.41	Scan date		29/12/2022
Risks at sampling date			Nuchal translucency 1.1		
Age Risk	sk 1:776		Nuchal translucency MoM 0.8		
Biochemical T21 risk		1:1343	Nasal bone		Present
Combined trisomy 21 risk		1:7404			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:100 1:250 Cut off 1:1000 1:1			The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21with NT test it is expected that among 7404 women with the same data, there is one woman with a trisomy 21 pregnancy and 7403 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		