

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

Book a Test Online www.molq.in

					Date of Report PRISCA	12-12-2023 5.2.0.13
Patient Data					ТКІЗСЛ	3.2.0.10
Name		M	RS DEEPIKA	Patient ID		012312110076
Birthday	23-02-1993			3 Sample ID		11659801
Age at Sample date			30.	8 Sample Date		11-12-2023
Gestational age 12+2				2		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52 Diabetes			NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+1
PAPP-A	5.14	mIU/ml	0.88	Method		CRL (<>Robinson)
fb-hCG	53.5	ng/ml	1.28	Scan date		10-12-2023
Risks at sampling date				Crown rump length in mm 52		
Age Risk			1:576	Nuchal translu	cency MoM	0.72
Biochemical T21 risk			1:1534	Nasal bone		PRESENT
Combined trisomy 21 risk			1:8178	Sonographer		DR.RAJESH
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk				Down's Syndro	ome Risk (Trisomy 21	Screening)
The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk				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 test (with NT) it is expected that among 8178 women with the same data, there is one woman with a trisomy 21 pregnancy and 8177 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 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		
					e Risk	Risk below Age risk