

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

Book a Test Online www.molq.in

				Date of Report PRISCA	31-12-2023 5.2.0.13
Patient Data				Тизел	3.2.0.10
Name	MRS ARCHNA				012312300068
Birthday		22-01-1994	Sample ID		11837938
Age at Sample date		29.9	Sample Date		31-12-2023
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+2
PAPP-A	3.47 mIU/ml	0.68	Method		CRL (<>Robinson)
fb-hCG	13.8 ng/ml	0.42	Scan date		26-12-2023
Risks at sampling date			Crown rump length in mm 58		
Age Risk		1:659	Nuchal translu	cency MoM	1.32
Biochemical T21 risk		1:9800	Nasal bone		PRESENT
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk		Down's Syndro	ome Risk (Trisomy 21	Screening)	
1:10 1:100 1:250 1:1000 1:1000 1:100000 1:100000 1:100000 1:100000 1:100000 1:10000		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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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			
Risk Above Cut Off Risk above Age Risk     Risk below Age risk					