

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

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				Date of Report PRISCA	11-03-2024 5.2.0.13
Patient Data					
Name	Ĩ	MRS. SAVITA	Patient ID	01	12403100093
Birthday		11-12-1991	Sample ID		11846983
Age at Sample date		32.2	Sample Date		10-03-2024
Gestational age		13+4			
Correction factors				Γ	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	65 Diabetes	5	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	13+3
PAPP-A	6.9 mIU/ml	0.96	Method		CRL (<>Robinson)
fb-hCG	13.8 ng/ml	0.52	Scan date		09-03-2024
Risks at sampling date			Crown rump l	rump length in mm 73.8	
Age Risk		1:482	Nuchal translu	cency MoM	0.60
Biochemical T21 risk		<1:10000	Nasal bone		present
combined trisomy 21 risk		<1:10000	Sonographer		DR.
Trisomy 13/18+NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk 1:10			Down's Syndr	ome Risk (Trisomy 2	1 Screening)
1:100 1:250 1:1000 1:1000 1:1000 1:1000 13 15 17 19 21 23 25 27 Trisomy 13/18+NT The calculated risk for Tris 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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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			
	Above Cut Off		Risk above Ag	e Risk	isk below Age risk