

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

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				Date of Report PRISCA	23-10-2024 5.2.0.13
Patient Data					
Name	I	MRS. LAXMI	Patient ID		12410210188
Birthday		06-07-1994	4 Sample ID		11892664
Age at Sample date	30.3		3 Sample Date		21-10-2024
Gestational age		12+3	3		
Correction factors				_	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	51 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	12+2
PAPP-A	5.1 mIU/ml	0.81	Method		CRL (<>Robinson)
fb-hCG	84.47 ng/ml	2.08	Scan date		21-10-2024
Risks at sampling date			Crown rump l	ength in mm	59
Age Risk		1:620	Nuchal translu	icency MoM	1.13
Biochemical T21 risk		1:415	Nasal bone		present
Combined trisomy 21 risk		1:1174	Sonographer		DR. SHRUTI SANGWAN
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 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:100 1:10000 1:10000 1:10000 1:100		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 1174 women with the same data, there is one woman with a trisomy 21 pregnancy and 1173 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 Ag	e Risk	Risk below Age risk