

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

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				Date of Report PRISCA	14-09-2023 5.2.0.13
Patient Data					0.210120
Name	MRS MAHIMA				012309130047
Birthday		22-11-1997	Sample ID		11805158
Age at Sample date		25.8	Sample Date		13-09-2023
Gestational age 13+2			2		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	40.7 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age		13+2
PAPP-A	5.48 mIU/n	nl 0.49	Method		CRL (<>Robinson)
fb-hCG	34.6 ng/ml	1	Scan date		13-09-2023
Risks at sampling date			Crown rump length in mm 71.1		
Age Risk		1:948	Nuchal translu	cency MoM	0.62
Biochemical T21 risk		1:1060	Nasal bone		PRESENT
Combined trisomy 21 risk		1:6356	Sonographer		DR SAHIL LOOMBA
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 00 1:250 00 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 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 6356 women with the same data, there is one woman with a trisomy 21 pregnancy and 6355 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				e Risk	Risk below Age risk