

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

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				Date of Report PRISCA	08-072024 5.2.0.13
Patient Data					
Name MRS. PRIYA			Patient ID		12407070177
Birthday	02-04-1999		Sample ID		11860824
Age at Sample date	ge at Sample date 25.3		Sample Date 07-07-		07-07-2024
Gestational age 12+6					
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	45.8 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	12+6
PAPP-A	10.2 mIU/m	ıl 1.21	Method		CRL (<>Robinson)
fb-hCG	181.5 ng/ml	4.83	Scan date		07-07-2024
Risks at sampling date			Crown rump length in mm 64.9		
Age Risk 1:961		Nuchal translucency MoM 0.78			
Biochemical T21 risk	<u>x</u> 1:264		Nasal bone		PRESENT
Combined trisomy 21 risk	ombined trisomy 21 risk 1:1511		Sonographer DR.		DR. DR.Deepika
Trisomy 13/18 + NT		<1:10000		in measuring NT	MD
Risk 1:10			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:100 1:250 1:1000 1:1000 1:10000 13 15 17 19 21 23 25 27 Trisomy 13/18+NT The calculated risk for Trisor 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 1511 women with the same data, there is one woman with a trisomy 21 pregnancy and 1510 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

