

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

Book a Test Online www.molq.in

				Date of Report PRISCA	2/3/2023 5.2.0.13
Patient Data					
Name		MRS. ERA	Patient ID		012303010252
Birthday		30/03/1999	Sample ID		11487662
Age at term		24.3	Sample Date		1/3/2023
Gestational age		12+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy	21 unknown
Weight in kg	56 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+1
PAPP-A	2.28 mIU/ml	0.45	Method		CRL (<>Robinson)
fb-hCG	10.6 ng/ml	0.25	Scan date		1/3/2023
Risks at sampling date			Nuchal translucency 1.18		
Age Risk 1:989		1:989	Nuchal translucency MoM 0.8		
Biochemical T21 risk	iochemical T21 risk 1:7848		Nasal bone Presen		
Combined trisomy 21 ris	sk	<1:10000			
Trisomy 13/18 + NT		1:5034			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 Cut off 1:1000 1:250 Cut off 1:1000 1:1000 1:10000 1:10000 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 with NT test it is expected that among more than 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic		
1:5034 , which indicates a low risk Risk Above Cut Off			values Risk above Ag	e Risk	Risk below Age risk