

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

Book a Test Online www.molq.in

				Date of Report PRISCA	26/4/2023 5.2.0.13
Patient Data					
Name MRS. NEHA NIWAS			Patient ID		012304250227
Birthday		19/10/1988	Sample ID		11502444
Age at term		34.1	Sample Date		25/4/2023
Gestational age		12+5			
Correction factors				<b>1</b>	
Fetuses	1 IVF		unknown	Previous trisomy	21 unknown
Weight in kg	66 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
<b>Biochemical Data</b>		Ultrasound Data			
Parameter	Value	Corr Mom			
PAPP-A	2.87 mIU/ml	0.55	Method		CRL (<>Robinson)
fb-hCG	28.9 ng/ml	0.84	Scan date		
Risks at sampling date			Nuchal translucency 0.9		
Age Risk 1:304		1:304	Nuchal translucency MoM 0.55		
Biochemical T21 risk 1:674		1:674	Nasal bone Present		
Combined trisomy 21 risk 1:3823					
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 1:250 1:10 1:10000 1:10000 1:10			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 3823 women with the same data, there is one woman with a trisomy 21 pregnancy and 3822 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 values		
	isk Above Cut Off		Risk above Ag	e Risk	Risk below Age risk