

Book a Test Online www.molq.in

 Date of Report
 14/9/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. PRAGATI		Patient ID		012209130134	
Birthday	1/1/1990		Sample ID		11581448
Age at term 33.01		Sample Date 13/9/202		13/9/2022	
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	51.6 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+1
PAPP-A	4.29 mIU/ml	0.53	Method		CRL (⇔Robinson)
fb-hCG	27.2 ng/ml	0.82	Scan date		13/9/2022
Risks at sampling date			Nuchal translucency 0.9		
Age Risk 1:4		1:439	Nuchal translucency MoM 0.4		0.52
Biochemical T21 risk		1:908	Nasal bone		Present
Combined trisomy 21 ris	sk	1:5192			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 Cut off 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 Age Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, 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 21with NT test it is expected that among more than 5192 women with the same data, there is one woman with a trisomy 21 pregnancy and 5191 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		