

*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

 Date of Report
 1/1/2022

 PRISCA
 5.2.0.13

Patient Data					
Name	nme MRS. KOMAL		Patient ID		012112300116
Birthday		10/1/1994	Sample ID		11272335
Age at delivery		28.47	Sample Date		30/12/2021
Gestational age		13+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	67 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter V	alue	Corr Mom	Gestational age	2	13+3
PAPP-A	3.04 mIU/ml	0.52	Method		CRL(<>Robinson
fb-hCG	19.5 ng/ml	0.57	Scan date		28/12/2021
Risks at sampling date			Nuchal translucency 1.5		
Age Risk		1:1121	Nuchal translu	cency MoM	0.8
Biochemical T21 risk		1:4773	Nasal bone presen		
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:250 Cut off 1:1000 1:10000 1:15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 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 21 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 values Risk above Age Risk Risk below Age risk		