

*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report 14/3/2022PRISCA 5.1.0.17

Patient Data					
Name MRS. VANDNA PANDEY			Patient ID		012203130072
Birthday 15/07/1988		Sample ID		10964705	
Age at term 34.01		Sample Date 13/3/20		13/3/2022	
Gestational age		13+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	68 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+4
PAPP-A	$2.69~\mathrm{mIU/ml}$	0.49	Method		CRL (<>Robinson)
fb-hCG	24.7 ng/ml	0.62	Scan date		13/03/2022
Risks at sampling date			Nuchal translucency 1.4		
Age Risk		1:373	Nuchal translucency MoM 0.7		
Biochemical T21 risk		1:1110	Nasal Bone Preser		
Combined trisomy 21 risk 1:62		1:6225			
Trisomy 13/18 +NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
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 6225 women with the same data, there is one woman with a trisomy 21 pregnancy and 6224 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		