

*Free Home Sample Collection 9999 778 778



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 Date of Report
 30-01-2021

 PRISCA
 5.0.2.37

Patient Data				
Name MRS RINKI KUMARI		Patient ID		012101290003
Birthday 10-03-1997		Sample ID		10881412
Age at delivery 23.9		Sample Date		29/01/2021
Gestational age 13+1				
Correction factors				
Fetuses 1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg 55	Diabetes	unknown	Pregnancies	unknown
Smoker Unknown	Origin	Asian		
Biochemical Data		Ultrasound Data		
Parameter Value	Corr Mom	Gestational ago	2	13+0
PAPP-A 3.45	mIU/ml 0.56	Method		CRL(<>Robinson
fb-hCG 41.3	ng/ml 0.92	Scan date		28-01-2021
Risks at sampling date		Crown rump length in mm 67.2		
Age Risk	1:1024	Nuchal translucency MOM 0.89		
Biochemical T21 risk	1:1906	Nasal bone Prese		Present
Combined Trisomy 21 Risk	<1:10000	Sonographer DR.VARG		DR.VARUN
Trisomy 13/18 + NT <1:10000		Qualification in measuring NT MBBS,M		
Risk 1:10	Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:250 1:1000 13 15 17 19 21 23 25 27 29 31 33 Trisomy 13/18 + NT The calculated risk for Trisomy 1 <1:10000, which indicates a low ri	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 (with NT) 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			