

*Free Home Sample Collection 9999 778 778



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Date of Report 25/1/2022PRISCA 5.2.0.13

Patient Data					
Name MRS. RUCHI			Patient ID		012201240013
Birthday		1/1/1990	Sample ID		10907921
Age at delivery		32.1	Sample Date		24/1/2022
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	63 Diabetes		NO	Pregnancies	unknown
Smoker 1	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter Va	lue	Corr Mom	Gestational age	2	12+4
PAPP-A	2.85 mIU/ml	0.63	Method		CRL (<>Robinson)
fb-hCG	20.2 ng/ml	0.45	Scan date		23/01/2022
Risks at sampling date			Nuchal translucency 1.6		
Age Risk		1:483	Nuchal translucency MoM 1.0		1.03
Biochemical T21 risk		1:5287	Nasal Bone Preser		Present
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:1000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 (with 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		