

*Free Home Sample Collection 9999 778 778



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 Date of Report
 6/2/2024

 PRISCA
 5.2.0.13

				TMSCA	5.2.0.13
Patient Data					
Name	JAGRUTI		Patient ID		012402050061
Birthday	thday 2/9/1990		Sample ID 1		11827455
Age at sample 33.4		Sample Date 5/2/2024			
Gestational age 12+6					
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	65.2 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ago	e	11+5
PAPP-A	4.80 mIU/ml	0.86	Method CRL (<>Robinson)		
fb-hCG	58.2 ng/ml	1.75	Scan date		28/1/2024
Risks at sampling date			Crown rump length in mm 51		
Age Risk 1:380		1:380	Nuchal translucency MoM 0.8		
biochemical T21 risk 1:455		Nasal bone Present			
Combined trisomy 21 risk 1:2531		1:2531	Sonographer DR.ABHISHEK		DR.ABHISHEKH
Trisomy 13/18 + NT <1:10000		<1:10000	Qualifications in measuring NT M		MBBS
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 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 (with NT) it is expected that among 2531 women with the same data, there is one woman with a trisomy 21 pregnancy and 2530 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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		