

*Free Home Sample Collection 9999 778 778



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Date of Report 14 - 12 - 2023PRISCA 5.2.0.13

			TRIBUTI	0.2.0.10	
atient Data					
Name MRS RUDHARAKSHI		Patient ID		012312120372	
Birthday 15-10-2004		Sample ID 1184829		11848292	
Age at Sample date 19.2		Sample Date 12-12-202		12-12-2023	
Gestational age 13+5					
orrection factors					
etuses 1	₹	unknown	Previous trisomy 21	unknown	
Veight in kg 60.5	abetes	NO	Pregnancies	unknown	
moker NO	igin	Asian			
ochemical Data		Ultrasound Da	ata		
rameter Value	Corr Mom	Gestational age	e	13+4	
APP-A 4.79	U/ml 0.59	Method		CRL (<>Robinson)	
-hCG 23.7	ml 0.91	Scan date		12-12-2023	
Risks at sampling date			Crown rump length in mm 78		
ge Risk	1:1141	Nuchal translucency MoM 0.		0.84	
Biochemical T21 risk		Nasal bone PRESE		PRESENT	
ombined trisomy 21 risk	<1:10000	Sonographer		DR JIGAR MAKWANA	
Trisomy 13/18 + NT		Qualifications in measuring NT		DNB	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 1:1			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 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 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		
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