

*Free Home Sample Collection 9999 778 778



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Date of Report 28-09-2024

				Date of Keport	20-09-2024
				PRISCA	5.2.0.13
RS. RITU YADAV W/O MANJEET			Patient ID		12409270195
		07-01-2002	Sample ID		11966908
22.7			Sample Date		27-09-2024
12+0					
1	IVF		unknown	Previous trisomy 21	unknown
54.5	Diabetes		NO	Pregnancies	unknown
NO	Origin		Asian		
			Ultrasound D	ata	
Value		Corr Mom	Gestational ago	2	12+0
5.1	mIU/ml	1.03	Method		CRL (<>Robinson)
	1 54.5 NO Value	1 IVF 54.5 Diabetes NO Origin	07-01-2002 22.7 12+0 1 IVF 54.5 Diabetes NO Origin Value Corr Mom	54.5 Diabetes NO NO Origin Asian Ultrasound Description Value Corr Mom Gestational age	RS. RITU YADAV W/O MANJEET 07-01-2002 22.7 12+0 1 IVF unknown Previous trisomy 21 54.5 Diabetes NO Origin Asian Value Corr Mom Patient ID Sample ID Sample Date NO Pregnancies Vultrasound Data Gestational age

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+0	
PAPP-A	5.1 mIU/ml	1.03	Method	CRL (<>Robinson)	
fb-hCG	51.4 ng/ml	1.16	Scan date	27-09-2024	
Risks at sampling date			Crown rump length in mm	52.2	
Age Risk		1:1018	Nuchal translucency MoM	0.93	
Biochemical T21 risk		1:4812	Nasal bone	PRESENT	
Combined trisomy 21 risk		<1:10000	Sonographer	DR.	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS	
Risk 1:10 1:100 1:250 1:1000		Cut off	Down's Syndrome Risk (Trisomy The calculated risk for Trisomy Scut off, which represents a low risk After the result of the Trisomy 21 test expected that among more than 1000 there is one woman with a trisomy 21. The calculated risk by PRISCA depending or the risk calculations are statistical applications are statistical applications are statistical applications are combined risk presumes done according to accepted guidelines.	21(with NT) is below the sk. t (with NT) it is 0 women with the same data, pregnancy. Inds on the accuracy of the physician. Please note that proaches and have no that NT measurement was	

13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 1998).

Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off

Risk above Age Risk

Risk below Age risk