

*Free Home Sample Collection 9999 778 778



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Date of Report 26-04-2021 PRISCA 5 0 2 37

				PRISCA	5.0.2.37
Patient Data					
Name	MRS. PO	OJA SHARMA	Patient ID		052104200036
Birthday		24-11-1992	Sample ID		10870066
Age at delivery		28.4	Sample Date		20/04/2021
Gestational age 12+3					
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	67 Diabetes	3	No	Pregnancies	unknown
Smoker	No Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+2
PAPP-A	2.63 mIU/ml	0.69	Method		CRL(<>Robinson
fb-hCG	33.9 ng/ml	0.74	Scan date		20-04-2021
Risks at sampling date			Crown rump length in mm 58.3		
Age Risk 1:7		1:764	Nuchal translucency MOM		0.52
Biochemical T21 risk		1:3891	Nasal bone		Present
Combined Trisomy 21 Risk		<1:10000	Sonographer		Dr
Trisomy 13/18 + NT		<1:10000	Qualification is	n measuring NT	MBBS, MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.		
1:1000 1:250 Cut off 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			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. 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		
Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			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		