

*Free Home Sample Collection 9999 778 778



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 Date of Report
 21-03-2024

 PRISCA
 5.2.0.13

Patient Data						
Name	MRS. 1	KRAT	TRIPATHI	Patient ID		12403200164
Birthday			19-11-1991	Sample ID		11850041
Age at Sample date			32.3	Sample Date		20-03-2024
Gestational age 12+0						
Correction factors						
Fetuses	1 IV	F		unknown	Previous trisomy 21	unknown
Weight in kg	63 D i	abetes		NO	Pregnancies	unknown
Smoker	NO Or	rigin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+0
PAPP-A	5.9 mI	U/ml	1.42	Method		CRL (<>Robinson)
fb-hCG	31.5 ng/	/ml	0.75	Scan date		20-03-2024
Risks at sampling date				Crown rump length in mm 53.9		
Age Risk			1:449	Nasal transluce	ency MoM	0.84
Biochemical T21 risk			<1:10000	Nasal bone		present
Combined trisomy 21 risk			<1:10000	Sonographer		DR. NAVEEN
Trisomy 13/18 + NT			<1:10000	Qualification is	n measuring NT	MBBS
Risk 1:10				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:1000 1:1000 1:1000 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 it is expected that among 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 comboned risk presumes the NT measurement was done according to accepted guidelines (prenat Diagn 18:511-523(1998)). The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!		