

\*Free Home Sample Collection 9999 778 778



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Date of Report 18-10-2024 PRISCA 5.2.0.13

			PRISCA		5.2.0.13
Patient Data					
Name		MRS. FIRDOSH	Patient ID		12410160136
Birthday		16-06-1996	Sample ID		11863903
Age at Sample date		28.3	Sample Date		16-10-2024
Gestational age		13+4			
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	53	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Weight in kg	53 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data		Ultrasound Data			
Parameter	Value	Corr Mom	Gestational age	e	13+4
PAPP-A	7.9 mIU/ml	0.87	Method		CRL (<>Robinson)
fb-hCG	66.5 ng/ml	2.33	Scan date		16-10-2024
Risks at sampling date			Crown rump length in mm 74.5		
Age Risk		1:800	Nuchal translu	cency MoM	0.65
Biochemical T21 risk		<b>1:47</b> 3	Nasal bone		PRESENT
Combined trisomy 21 risk		1:2743	Sonographer		DR. SANJEEV KUMAR
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:250 Cut off 1:1000			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 2743 women with the same data, there is one woman with a trisomy 21 pregnancy and 2742 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		

## Trisomy 13/18+NT

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

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

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