

which indicates a low risk

*Free Home Sample Collection 9999 778 778



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Date of Report 15-12-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. GEETANJALI			Patient ID		12412140121
Birthday	22-10-1988			Sample ID		11981156
Age at Sample date 36.1				Sample Date		14-12-2024
Gestational age			11+6	5		
Correction factors		•				
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	65	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	11+6
PAPP-A	2.74	mIU/ml	0.73	Method		CRL (<>Robinson)
fb-hCG	11.2	ng/ml	0.26	Scan date		14-12-2024
Risks at sampling date				Crown rump length in mm 52.4		
Age Risk	Risk 1:204			Nuchal translucency MoM 0.71		
Biochemical T21 risk	al T21 risk		1:5699	Nasal bone		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer DR. SANJEEV KUMA		DR. SANJEEV KUMAR
Frisomy 13/18 + NT 1		1:6509	Qualifications in measuring NT		MD	
				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:20 Gt off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The free beta HCG level is low. 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		
The calculated risk for Trisomy 13/18 (with NT) is 1:6509, which indicates a low risk				risk assessment! Calculated risks have no diagnostic values		