

\*Free Home Sample Collection 9999 778 778



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Date of Report 18-10-2024 PRISCA 5.2.0.13

					5.2.0.13	
Patient Data						
Name		MRS.	SUPRIYA	Patient ID		12410160118
Birthday			09-05-1997	Sample ID		11905608
Age at Sample date			27.4	Sample Date		16-10-2024
Gestational age	12+3					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	51	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data			Ultrasound Data			

Weight in kg	51 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data		Ultrasound Data			
Parameter	Value	Corr Mom	Gestational age	2	11+1
PAPP-A	$4.8~\mathrm{mIU/ml}$	0.76	Method		CRL (<>Robinson)
fb-hCG	42.6 ng/ml	1.05	Scan date		07-10-2024
Risks at sampling date			Crown rump le	ength in mm	42.9
Age Risk		1:829	Nuchal translucency MoM		0.94
Biochemical T21 risk		1:2483	Nasal bone		PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer		DR. INDRAJEET
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk 1:10		Down's Syndrome Risk (Trisomy 21 Screening)			
1:100		Cut off	cut off, which After the result of expected that an	I risk for Trisomy 21(vertical represents a low risk.) of the Trisomy 21 test (with a mong more than 10000 we man with a trisomy 21 pre-	th NT) it is omen with the same data,
1:10000 1:10000 13 15 17 19 21 23 25 27	29 31 33 35 37 39		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).		
Trisomy 13/18+NT	. 41 10000	The laboratory of	The laboratory cannot be hold responsible for their impact on the		

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