

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

which indicates a low risk

*Free Home Sample Collection 9999 778 778



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Date of Report 05-08-2024

risk assessment! Calculated risks have no diagnostic values

				PRISCA	5.2.0.13
Patient Data					
Name	MRS. R	REKHA DEVI	Patient ID		12408040160
Birthday		29-06-1994	Sample ID		11883928
Age at Sample date		30.1	Sample Date		04-08-2024
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58.8 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ago	2	12+4
PAPP-A	4.1 mIU/ml	0.69	Method		CRL (<>Robinson)
fb-hCG	42.4 ng/ml	1.19	Scan date		03-08-2024
Risks at sampling date			Crown rump l	ength in mm	60.3
Age Risk		1:642	Nuchal translu	cency MoM	1.02
Biochemical T21 risk		1:1145	Nasal bone		PRESENT
Combined trisomy 21 risk		1:4566	Sonographer		DR. VIKASH
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 1:1000	Cut off, which rep After the result of the expected that amond there is one woman 4565 women with n The calculated risk information provide the risk calculations diagnostic value! The patient combine done according to a			I risk for Trisomy 21 (we represents a low risk.) of the Trisomy 21 test (with nong 4566 women with the nan with a trisomy 21 pregath not affected pregnancies lisk by PRISCA depends of wided by the referring physions are statistical aapproach abined risk presumes that I to accepted guidelines (Presumes 1) and the property of the presumes that I to accepted guidelines (Presumes 2) and the presumes (Presumes 2) and the presu	h NT) it is e same data, gnancy and s. on the accuracy of the sician. Please note that thes and have no
13 15 17 19 21 23 25 27 Trisomy 13/18+NT	29 31 33 35 37 39	41 43 45 47 49	1998). The laboratory o	cannot be hold responsible	e for their impact on the