

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

which indicates a low risk

*Free Home Sample Collection 9999 778 778



risk assessment! Calculated risks have no diagnostic values

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Date of Report 23-11-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name	MR	S. SHAHEEN	Patient ID		12411220134
Birthday		29-08-1996	Sample ID		11871754
Age at Sample date		28.2	Sample Date		22-11-2024
Gestational age 12+			_		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	61 Diabetes	S	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound D	ata	
Parameter	Value	Corr Mom	Gestational age	e	12+0
PAPP-A	5.4 mIU/ml	1.11	Method		CRL (<>Robinson)
fb-hCG	36.6 ng/ml	0.92	Scan date		20-11-2024
Risks at sampling date			Crown rump length in mm 53.6		
Age Risk 1:		1:772	Nuchal translucency MoM		0.63
Biochemical T21 risk		1:7170	Nasal bone		PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer		DR. POONAM DHALL
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Gutoff			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 calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that		
1:1000 1:1000 13 15 17 19 21 23 25 Trisomy 13/18+NT The calculated risk for T	527293133353733 Prisomy 13/18 (with NT)		the risk calculate diagnostic value The patient con done according 1998).	ons are statistical aapproac	ches and have no NT measurement was enat Diagn 18:511-523;