

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 11-09-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name				Patient ID		12409100187
Birthday	28-03-1998			Sample ID		11966975
Age at Sample date	e at Sample date 26.5					10-09-2024
Gestational age 13+2				!		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	83 Diabetes			NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	13+2
PAPP-A	7.1	mIU/ml	1.47	Method		CRL (<>Robinson)
fb-hCG	67.3	ng/ml	2.47	Scan date		10-09-2024
Risks at sampling date				Crown rump l	ength in mm	71.2
Age Risk				Nuchal translucency MoM 0.90		
Biochemical T21 risk			1:1353	Nasal bone		PRESENT
Combined trisomy 21 risk			1:6255	Sonographer		DR. AMENDA
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MD
Risk				Down's Syndr	ome Risk (Trisomy 21	Screening)
1:100 1:250 Cut off 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				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 6255 women with the same data, there is one woman with a trisomy 21 pregnancy and 6254 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 done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).		
Trisomy 13/18+NT				1998).	cannot be hold responsible	e for their impact on the
The calculated risk for Trison	with NT)	is <1:10000.				