

*Free Home Sample Collection 9999 778 778

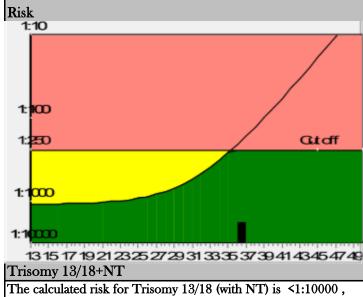


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Date of Report 15-02-2025 PRISCA 5.2.0.13

					TRISCIT	0.2.0.10
Patient Data						
Name	N	ARS. GAUT	CAMI DESI	Patient ID		12502140062
Birthday	31-07-1988			Sample ID		11908452
Age at Sample date	36.5			Sample Date		14-02-2025
Gestational age			11+5			
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 D	ata	

Biochemical Data			Olu asoulid Data	
Parameter Value		Corr Mom	Gestational age	11+4
PAPP-A	3.2 mIU/ml	0.68	Method	CRL (<>Robinson)
fb-hCG	51.6 ng/ml	1.07	Scan date	13-02-2025
Risks at sampling date			Crown rump length in mm	52.6
Age Risk		1:185	Nuchal translucency MoM	0.64
Biochemical T21 risk		1:409	Nasal bone	PRESENT
Combined trisomy 21 risk		1:2271	Sonographer	DR.DEEPIKA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD



Down's Syndrome Risk (Trisomy 21 Screening)

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 2271 women with the same data, there is one woman with a trisomy 21 pregnancy and 2270 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).

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

which indicates a low risk