

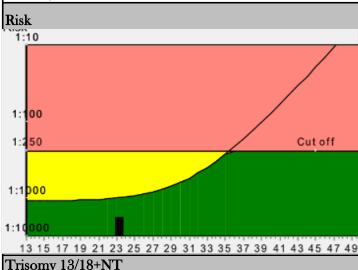
*Free Home Sample Collection 9999 778 778



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					Date of Report PRISCA	09-07-2024 5.2.0.13
Patient Data						
Name			MRS. NEHA	Patient ID		12407080107
Birthday	06-04-2001			Sample ID		11903864
Age at Sample date	23.3			Sample Date		08-07-2024
Gestational age			12+1			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	66.9	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound D	ata	
Parameter	Value		Corr Mom	Gestational ag	e	12+1
PAPP-A	4.1	m I U/ml	1.00	Method		CRL (<>Robinson)
fb-hCG	35.6	ng/ml	0.89	Scan date		08-07-2024
Risks at sampling date				Crown rump l	ength in mm	56.2
Age Risk			1:1009	Nuchal translucency MoM		1.15
Biochemical T21 risk			1:8063	Nasal bone		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer		DR. Manish chaudhary

<1:10000



Trisomy 13/18 + NT

Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

MD

Qualifications in measuring NT

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 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;

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