

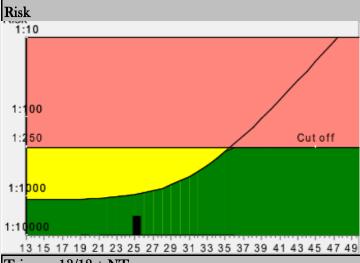
*Free Home Sample Collection 9999 778 778



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				Date of Report	17-06-2024
			1	PRISCA	5.2.0.13
Patient Data					
Name		MRS. GEETA	Patient ID		12407160227
Birthday		13-05-1999	Sample ID		11867661
Age at Sample date		25.2	Sample Date		16-07-2024
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	50 Diabete	es	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Da	ata	
Parameter	Value	Corr Mom	Gestational ago	e	13+0
PAPP-A	6.1 mIU/m	0.72	Method		CRL(<>Robinson)
fb-hCG	22.5 ng/ml	0.67	Scan date		15-07-2024
Risks at sampling date			Crown rump l	ength in mm	77
Age Risk		1:974	Nuchal translu	icency MoM	1.22

Nasal bone Biochemical T21 risk 1:6821 present Combined trisomy 21 risk <1:10000 Sonographer DR. SHRUTI SANGWAN Trisomy 13/18 + NT <1:10000 Qualifications in measuring NT MBBS Risk Down's Syndrome Risk (Trisomy 21 Screening)



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 test (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 amog 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 approaches 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