

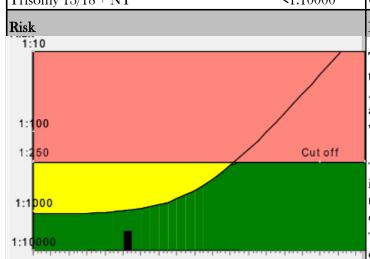
*Free Home Sample Collection 9999 778 778



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					Date of Report	25-07-2024
					PRISCA	5.2.0.13
Patient Data						
Name	MRS. DIVYA W/O NILESH			Patient ID		12407240202
Birthday	31-12-2000			Sample ID		11884224
Age at Sample date	23.6			Sample Date		24-07-2024
Gestational age	12+2					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	78.3	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Da	ata	
Parameter	Value		Corr Mom	Gestational age	e	12+1
PAPP-A	5. 3	m I U/ml	1.48	Method		CRL(<>Robinson)
fb-hCG	21.8	ng/ml	0.59	Scan date		23-07-2024
Risks at sampling date				Crown rump l	ength in mm	54.3

Age Risk Nuchal translucency MoM 1.39 1:1005 Biochemical T21 risk Nasal bone <1:10000 present Combined trisomy 21 risk <1:10000 Sonographer DR. VIKASH Trisomy 13/18 + NT <1:10000 Qualifications in measuring NT MD



13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18 + NT

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

Down's Syndrome Risk (Trisomy 21 Screening)

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 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