

*Free Home Sample Collection 9999 778 778

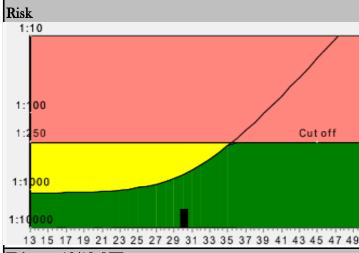


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Date of Report 20-10-2024

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. NANCY W/O RAJAT			Patient ID		12410190139
Birthday	13-09-1994			Sample ID		11978140
Age at Sample date	30.1			Sample Date		19-10-2024
Gestational age	12+4					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	64	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data	Pata U				ata	
Parameter	Value		Corr Mom	Gestational age	e	12+4
PAPP-A	4.9	mIU/ml	0.96	Method		CRL (<>Robinson)
fb-hCG	56.4	ng/ml	1.56	Scan date		19-10-2024

60.1 Risks at sampling date Crown rump length in mm Age Risk 1:639 Nuchal translucency MoM 0.45 Biochemical T21 risk 1:1279 Nasal bone **PRESENT** Combined trisomy 21 risk 1:6855 Sonographer DR. Trisomy 13/18 + NT <1:10000 Qualifications in measuring NT MBBS



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)

After the result of the Trisomy 21 test (with NT) it is expected that among 6855 women with the same data, there is one woman with a trisomy 21 pregnancy and 6854 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

Trisomv 13/18+NT

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