

*Free Home Sample Collection 9999 778 778

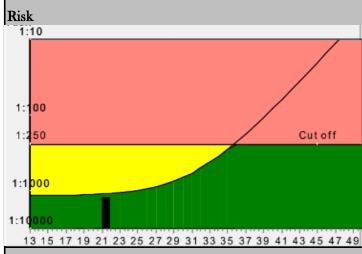


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Date of Report 11-10-2024 PRISCA 59012

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. SAPNA	W/O DH	ARAMVEER	Patient ID		12410090140
Birthday	24-05-2003			Sample ID		11966774
Age at Sample date	21.4			Sample Date		09-10-2024
Gestational age			13+6			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	43.6	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Da	ata	

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	13+4	
PAPP-A	7.3 mIU/ml	0.58	Method	CRL (<>Robinson)	
fb-hCG	73.2 ng/ml	2.63	Scan date	07-10-2024	
Risks at sampling date			Crown rump length in mm	76.6	
Age Risk		1:1113	Nuchal translucency MoM	0.75	
Biochemical T21 risk		1:189	Nasal bone	PRESENT	
Combined trisomy 21 risk		1:1239	Sonographer	DR. AMENDA	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
Risk 1:10			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.		



Trisomy 13/18+NT

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

After the result of the Trisomy 21 test (with NT) it is expected that among 1239 women with the same data, there is one woman with a trisomy 21 pregnancy and 1238 women with not affected pregnancies. The free beta HCG level is high.

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