

*Free Home Sample Collection 9999 778 778

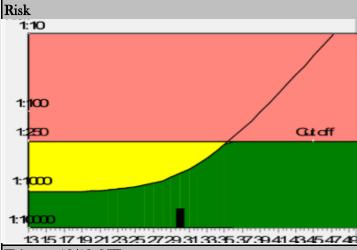


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Date of Report 21-12-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13	
Patient Data							
Name	N	IRS. SHRIY.	A GUPTA	Patient ID		12412200112	
Birthday			02-12-1994	Sample ID		11863580	
Age at Sample date			30.1	Sample Date		20-12-2024	
Gestational age			12+2				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	68.5	Diabetes		NO	Pregnancies	unknown	
Smoker	NO	Origin		Asian			
Biochemical Data				Ultrasound Data			
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Biochemical Data			Ultrasound Data		
Parameter Value		Corr Mom	Gestational age	12+0	
PAPP-A	3.1 mIU/ml	0.73	Method	CRL (<>Robinson)	
fb-hCG	18.94 ng/ml	0.5	Scan date	18-12-2024	
Risks at sampling date			Crown rump length in mm	56.6	
Age Risk		1:636	Nuchal translucency MoM	0.94	
Biochemical T21 risk	_	1:8511	Nasal bone	PRESENT	
Combined trisomy 21 risk	_	<1:10000	Sonographer	DR. DHRUV TANEJA	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10					



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.

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; 1008)

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