

\*Free Home Sample Collection 9999 778 778

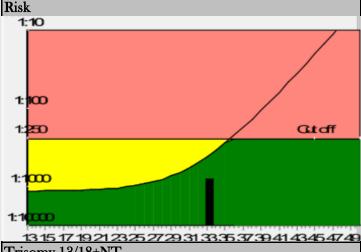


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Date of Report 16-12-2024 DDICCA 59019

				PRISCA		5.2.0.13
Patient Data						
Name	M	RS. KIRAN I	KILOLIYA	Patient ID		12412140263
Birthday			04-09-1991	Sample ID		11969941
Age at Sample date			33.3	Sample Date		14-12-2024
Gestational age			13+2			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Da	ata.	

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Parameter Value		Corr Mom	Gestational age	13+1	
PAPP-A	6.9 mIU/ml	0.96	Method	CRL (<>Robinson)	
fb-hCG	93.4 ng/ml	3.11	Scan date	14-12-2024	
Risks at sampling date			Crown rump length in mm	67.8	
Age Risk		1:397	Nuchal translucency MoM	0.81	
Biochemical T21 risk		1:137	Nasal bone	PRESENT	
Combined trisomy 21 risk		1:792	Sonographer	DR. DEEPIKA	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
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(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 792 women with the same data, there is one woman with a trisomy 21 pregnancy and 791 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;

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