

\*Free Home Sample Collection 9999 778 778

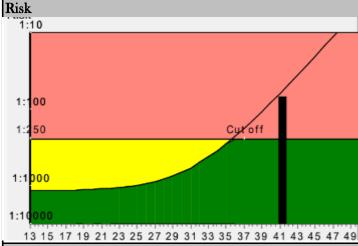


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Date of Report 15-10-2024 PRISCA 5.2.0.13

			PRISCA		3.2.0.13	
Patient Data						
Name		MR	S. RAMAN	Patient ID		12410140077
Birthday	26-06-1983			Sample ID		1197837 <i>5</i>
Age at Sample date	41.3			Sample Date		14-10-2024
Gestational age			13+6			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	13+5	
PAPP-A	8.1 mIU/ml	0.90	Method	CRL (<>Robinson)	
fb-hCG	91.5 ng/ml	3.66	Scan date	13-10-2024	
Risks at sampling date			Crown rump length in mm	78.6	
Age Risk		1:58	Nuchal translucency MoM	0.63	
Biochemical T21 risk		>1:50	Nasal bone	PRESENT	
Combined trisomy 21 risk		1:68	Sonographer	DR. SUBHASISH	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10			The calculated risk for Trisomy 21 (w	with NT) is above the	



## 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 above the cut off, which represents an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 68 women with the same data, there is one woman with a trisomy 21 pregnancy and 67 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