

\*Free Home Sample Collection 9999 778 778

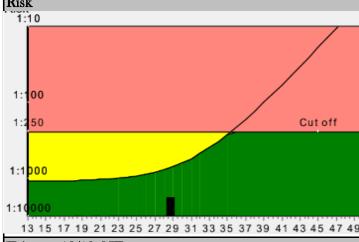


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Date of Report 13-06-2024 PRISCA 5.2.0.13

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					5.2.0.13	
Patient Data						
Name	M	RS. PRIT	TKA DIMAN	Patient ID		12406120198
Birthday	02-10-1995			Sample ID		11905852
Age at Sample date	28.7			Sample Date		12-06-2024
Gestational age	12+0					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	72	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound D	ata	
Parameter	Value		Corr Mom	Gestational ag	e	12+0

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+0	
PAPP-A	5.7 mIU/ml	1.61	Method	CRL (<>Robinson)	
fb-hCG	21.9 ng/ml	0.54	Scan date	12-06-2024	
Risks at sampling date			Crown rump length in mm	53.5	
Age Risk		1:732	Nuchal translucency MoM	0.81	
Biochemical T21 risk		<1:10000	Nasal bone	PRESENT	
Combined trisomy 21 risk		<1:10000	Sonographer	DR. INDRAJEET	
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 (with NT) is below the cut off, which represents a low risk.  After the result of the Trisomy 21 test (with NT) it is		



## 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 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; 1998).

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