

*Free Home Sample Collection 9999 778 778



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 Date of Report
 30-01-2024

 PRISCA
 5.2.0.13

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Patient Data						
Name		M	RS PRABHA	Patient ID		012401280104
Birthday			16-11-1997	Sample ID		11813165
Age at Sample date			26.2	Sample Date		28-01-2024
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	55	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	2	12+2
PAPP-A	4.6	mIU/ml	0.84	Method		CRL (<>Robinson)
fb-hCG	18.4	ng/ml	0.45	Scan date		07-01-2024
Risks at sampling date						
Age Risk			1:897			
Biochemical T21 risk			1:600			
Combined trisomy 21 risk			<1:10000			
Trisomy 13/18 + N T			<1:10000			
Risk				Down's Syndro	ome Risk (Trisomy 2	1 Screening)
1:10 1:250 Cut off 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies. 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		
which indicates a low risk	(with 1 V 1) 1	· · · · · · · · · · · · · · · · · · ·	risk assessment!	Calculated risks have no	diagnostic values	
Risk A	Above Cu	ıt Off		Risk above Ag	e Risk	Risk below Age risk