

which indicates a low risk

\*Free Home Sample Collection 9999 778 778



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Date of Report 07-09-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13	
Patient Data							
Name	MRS. DIVYA			Patient ID		12409050117	
Birthday	22-08-2000			Sample ID		11988983	
Age at Sample date 24.0				Sample Date		05-09-2024	
Gestational age			12+4				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	66	Diabetes		NO	Pregnancies	unknown	
Smoker	NO	Origin		Asian			
Biochemical Data				Ultrasound Data			
Parameter	Value		Corr Mom	Gestational ago	e	12+4	
PAPP-A	4.9	mIU/ml	0.99	Method		CRL (<>Robinson)	
fb-hCG	62.5 ng/ml		1.74	Scan date		05-09-2024	
Risks at sampling date				Crown rump length in mm 62			
Age Risk 1:			1:1000	Nuchal translucency MoM 0.59			
Biochemical T21 risk		1:1646	Nasal bone		PRESENT		
Combined trisomy 21 risk			1:8873	Sonographer D		DR.	
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT MBBS				
Risk 1:10				Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:250 Cut off 1:1000 1:1000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18+NT				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 8873 women with the same data, there is one woman with a trisomy 21 pregnancy 8872 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			
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk				risk assessment!	risk assessment! Calculated risks have no diagnostic values		