

*Free Home Sample Collection 9999 778 778



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Date of Report 20/12/2023 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name		YASHODA	Patient ID		012312190131
Birthday	ay 4/6/1998		Sample ID		11781025
Age at sample		30.5	Sample Date		19/12/2023
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age 12+3		
PAPP-A	5.6 mIU/ml	0.73	Method		CRL (<>Robinson)
fb-hCG	109.5 ng/ml	3.18	Scan date		15/12/2023
Risks at sampling date			Crown rump length in mm 60		
Age Risk 1:613		1:613	Nuchal translucency MoM 1.09		
Biochemical T21 risk	1:108		Nasal bone Present		
Combined trisomy 21 risk 1:35		1:352	Sonographer		DR.HARENDRA BHASKAR
Trisomy 13/18 + NT	risomy 13/18 + NT <1:10000		Qualifications in measuring NT MBF		
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 1:10000 1:10000 1:10000 Trisomy 13/18 + NT 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 352 women with the same data, there is one woman with a trisomy 21 pregnancy and 351 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					