

*Free Home Sample Collection 9999 778 778



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Date of Report 30-05-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name			MRS. PINKI	Patient ID		12405290145
Birthday			07-03-1993	Sample ID		11900453
Age at Sample date			31.2	Sample Date		29-05-2024
Gestational age			12+3			
Correction factors						
Fetuses	1 I	VF		unknown	Previous trisomy 21	unknown
Weight in kg	51 I	Diabetes		NO	Pregnancies	unknown
Smoker	NO (Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	e	12+2
PAPP-A	4.6 n	nIU/ml	0.73	Method		CRL (<>Robinson)
fb-hCG	35.7 n	ıg/ml	0.88	Scan date		28-05-2024
Risks at sampling date				Crown rump l	ength in mm	57
Age Risk			1:544	Nuchal translu	cency MoM	1.33
Biochemical T21 risk			1:2180	Nasal bone		PRESENT
Combined trisomy 21 risk			1:3137	Sonographer		DR. HARENDER BHASKAR
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk				Down's Syndr	ome Risk (Trisomy	21 Screening)
1:100 1:250 Cut off 1:1000				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 3137 women with the same data, there is one woman with a trisomy 21 pregnancy and 3136 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;		
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				1998).		

Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

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