

Risk Above Cut Off

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Risk below Age risk

Date of Report 17/12/2023 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name PRABHA KANTI			Patient ID		012312160151
Birthday 10/9/1993			Sample ID 11836501		
Age at sample 30.3		Sample Date 16/12/2		16/12/2023	
Gestational age		13+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58 Diabetes	3	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+6
PAPP-A	4.45 mIU/ml	0.49	Method		CRL (<>Robinson)
fb-hCG	17.7 ng/ml	0.71	Scan date		16/12/2023
Risks at sampling date			Crown rump length in mm 70		
Age Risk		1:653	Nuchal translucency MoM		0.91
Biochemical T21 risk		1:1566	Nasal bone		Present
Combined trisomy 21 risk		1:8108	Sonographer DR S		DR SANJEEV KUMAR
Γrisomy 13/18 + NT <1:10000		<1:10000	Qualifications in measuring NT		MBBS
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 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 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 8108 women with the same data, there is one woman with a trisomy 21 pregnancy and 8107 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					

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