

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

which indicates a low risk

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Date of Report 03-09-2024

risk assessment! Calculated risks have no diagnostic values

			P	PRISCA	5.2.0.13
Patient Data					
Name]	MRS. POOJA	Patient ID		12409020087
Birthday		06-10-1999	Sample ID		11860479
Age at Sample date		24.9	Sample Date		02-09-2024
Gestational age		12+4			
Correction factors					
Fetuses	1 IVF		unknown P	Previous trisomy 21	unknown
Weight in kg	60 Diabetes		NO P	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data	a	
Parameter	Value	Corr Mom	Gestational age		12+3
PAPP-A	6.4 mIU/ml	1.16	Method		CRL (<>Robinson)
fb-hCG	51.3 ng/ml	1.39	Scan date		01-09-2024
Risks at sampling date			Crown rump len	ngth in mm	61.7
Age Risk		1:967	Nuchal transluce	ency MoM	0.69
Biochemical T21 risk		1:3802	Nasal bone		PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer		DR. DEEPIKA
Trisomy 13/18 + NT		<1:10000	Qualifications in	n measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000		Cut off	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 than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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 Trisomy 13/18+NT	29 31 33 35 37 39	41 43 45 47 49	1998).	nnot be hold responsible	_