

which indicates a low risk

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Date of Report 04-11-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name	N	IRS. KUSUM	Patient ID		12411030169
Birthday		09-11-1996	Sample ID		11877712
Age at Sample date		28.0	Sample Date		03-11-2024
Gestational age		12+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	50 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+2
PAPP-A	5.1 mIU/ml	0.83	Method		CRL (<>Robinson)
fb-hCG	27.6 ng/ml	0.65	Scan date		03-11-2024
Risks at sampling date			Crown rump length in mm 52.5		
Age Risk		1:790	Nuchal translu	cency MoM	0.86
Biochemical T21 risk		1:8268	Nasal bone		PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer		DR. MONIKA
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
1:10 1:100 1:250 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.		
1:1000 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 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! Calculated risks have no diagnostic values		