

which indicates a low risk

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Date of Report 08-072024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13	
Patient Data							
Name	MRS. SONI YADAV			Patient ID		12407070175	
Birthday	08-02-2000			Sample ID		11860818	
Age at Sample date 24.4				Sample Date		07-07-2024	
Gestational age		13+5					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	63	Diabetes		NO	Pregnancies	unknown	
Smoker	NO	Origin		Asian			
Biochemical Data				Ultrasound Data			
Parameter	Value		Corr Mom	Gestational age	e	13+5	
PAPP-A	6.8	mIU/ml	0.87	Method		CRL (<>Robinson)	
fb-hCG	41.3	ng/ml	1.61	Scan date		07-07-2024	
Risks at sampling date				Crown rump length in mm 76.9			
Age Risk	Risk 1:1024			Nuchal translucency MoM 0.85			
Biochemical T21 risk			1:1549	Nasal bone		PRESENT	
Combined trisomy 21 risk 1:8211			1:8211	Sonographer		DR. DR.Deepika	
Trisomy 13/18 + NT	ny 13/18 + NT <1:10000		Qualifications in measuring NT		MD		
Risk				Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 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 8211 women with the same data, there is one woman with a trisomy 21 pregnancy and 8210 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			
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		