

which indicates a low risk

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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	07-03-1998			Sample ID		11900453
Age at Sample date 31.2				Sample Date		29-05-2024
Gestational age 12+3						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	l unknown
Weight in kg	51	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	12+2
PAPP-A	4.6	mIU/ml	0.73	Method		CRL (<>Robinson)
fb-hCG	35.7	ng/ml	0.88	Scan date		28-05-2024
Risks at sampling date				Crown rump length in mm 57		
Age Risk	ge Risk 1:544			Nuchal translucency MoM 0.70		
Biochemical T21 risk	iochemical 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 Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				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; 1998).		
Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
which indicates a low risk					5	·