

which indicates a low risk

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Date of Report 07-11-2024
PRISCA 5.2.0.13

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Patient Data						
Name	e MRS. MONIKA SHARMA			Patient ID		12411070048
Birthday 11-10-1996				Sample ID		11890277
Age at Sample date			28.1	Sample Date		07-11-2024
Gestational age 13+1						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	44.7	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	12+6
PAPP-A	6.3	mIU/ml	0.66	Method		CRL (<>Robinson)
fb-hCG	25.61	ng/ml	0.73	Scan date		05-11-2024
Risks at sampling date				Crown rump length in mm 65.7		
Age Risk			1:807	Nuchal translu	ncency MoM	0.66
Biochemical T21 risk			1:3739	Nasal bone		PRESENT
Combined trisomy 21 ris	k		<1:10000	Sonographer		DR.
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
				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 risk assessment! Calculated risks have no diagnostic values		