

which indicates a low risk

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Date of Report 04-07-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. KUNTI			Patient ID		12407040151
Birthday 01-01-2001				Sample ID		11860857
Age at Sample date 23.5				Sample Date		04-07-2024
Gestational age			11+5	5		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	41	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	11+5
PAPP-A	3.21	mIU/ml	0.53	Method		CRL (<>Robinson)
fb-hCG	45.3	ng/ml	0.86	Scan date		04-07-2024
Risks at sampling date				Crown rump length in mm 54.9		
Age Risk 1:986			1:986	Nuchal translucency MoM 0.89		
Biochemical T21 risk			1:1889	Nasal bone PRESEN		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer DR.DEEPIK		DR.DEEPIKA
Trisomy 13/18 + NT			<1:10000	Qualifications in measuring NT M		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				1990).		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk				The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		