

which indicates a low risk

\*Free Home Sample Collection 9999 778 778



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Date of Report 31-07-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. SALOCHANA			Patient ID		12407300169
Birthday	04-08-1996			Sample ID		11672948
Age at Sample date 28.0				Sample Date		30-07-2024
Gestational age			11+4			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	11+3
PAPP-A	3.4	mIU/ml	0.87	Method		CRL (<>Robinson)
fb-hCG	33.9	ng/ml	0.7	Scan date		29-07-2024
Risks at sampling date				Crown rump length in mm 49.9		
ge Risk 1:768			1:768	Nuchal translucency MoM 0.82		
Biochemical T21 risk		1:7575	Nasal bone		present	
Combined trisomy 21 risk			<1:10000	Sonographer DR. DEEPIN		DR. DEEPIKA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT		MD	
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
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.  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!		
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49  Trisomy 13/18+NT				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		