

The calculated risk for Trisomy 13/18 (with NT) is <1:10000,

which indicates a low risk

*Free Home Sample Collection 9999 778 778



risk assessment! Calculated risks have no diagnostic values

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Date of Report 19-07-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13	
Patient Data							
Name	MRS. PRIYA SONI			Patient ID		12407180080	
Birthday	29-04-1999			Sample ID		11877795	
Age at Sample date 25.2				Sample Date		18-07-2024	
Gestational age 11+1							
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	71	71 Diabetes		NO	Pregnancies	unknown	
Smoker	NO Origin			Asian			
Biochemical Data				Ultrasound Data			
Parameter	Value		Corr Mom	Gestational age	e	11+1	
PAPP-A	3.1	mIU/ml	1.26	Method		CRL(<>Robinson)	
fb-hCG	29.4	ng/ml	0.6	Scan date		18-07-2024	
Risks at sampling date				Crown rump length in mm 42.			
Age Risk			1:903	Nuchal translu	icency MoM	0.84	
Biochemical T21 risk			<1:10000	Nasal bone		present	
Combined trisomy 21 risk	X.		<1:10000	Sonographer		DR. RAHUL	
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 1:10000 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 test (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 amog 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! 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 Tris	omy 19/19 /	with NT	is <1.10000		The laboratory cannot be hold responsible for their impact on the		