

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 05-11-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13	
Patient Data							
Name		MRS. SU	J NEHA JAIN	Patient ID		22411040009	
Birthday			21-11-1987	Sample ID		11989383	
Age at Sample date 37.0				Sample Date		04-11-2024	
Gestational age 12+6							
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	57	57 Diabetes		NO	Pregnancies	unknown	
Smoker	NO	NO Origin		Asian			
Biochemical Data				Ultrasound Data			
Parameter	Value		Corr Mom	Gestational age	e	12+6	
PAPP-A	6.2	mIU/ml	0.95	Method		CRL (<>Robinson)	
fb-hCG	34.2	ng/ml	0.99	Scan date		04-11-2024	
Risks at sampling date				Crown rump length in mm 64.8			
Age Risk			1:175	Nuchal translu	icency MoM	0.97	
Biochemical T21 risk			1:1003	Nasal bone		PRESENT	
Combined trisomy 21 risl	ζ		1:4248	Sonographer		DR.	
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				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 4248 women with the same data, there is one woman with a trisomy 21 pregnancy and 4247 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			
1:10000 13 15 17 19 21 23 25 2 Trisomy 13/18+NT				done according 1998).	to accepted guidelines (P		
The calculated risk for Trise	omy 13/18	(with NT)	is <1·10000		10 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		