

The calculated risk for Trisomy 13/18 (with NT) is

Risk Above Cut Off

<1:10000 , which indicates a low risk

*Free Home Sample Collection 9999 778 778



on the risk assessment! Calculated risks have no diagnostic

Risk below Age risk

Risk above Age Risk

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					Date of Report PRISCA	14/8/2023 5.2.0.13
Patient Data						
Name		SWA	TI MANGLA	Patient ID		012308130139
Birthday			18/12/1994	Sample ID		11781367
Age at term			29.1	Sample Date		13/8/2023
Gestational age			12+5			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	44	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom			12+3
PAPP-A	3.87	mIU/ml	0.46	Method		CRL (<>Robinson)
fb-hCG	38.2	ng/ml	0.96	Scan date		11/8/2023
Risks at sampling date				Crown rump length in mm 59		
Age Risk			1:754	Nuchal translu	cency MoM	1.23
Biochemical T21 risk			1:786	Nasal bone		Present
Combined trisomy 21 risk			1:1710			
Trisomy 13/18 + NT			<1:10000			
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 1710 women with the same data, there is one woman with a trisomy 21 pregnancy and 1709 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!		
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				The laboratory cannot be hold responsible for their impact		