

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 above Age Risk

Risk below Age risk

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					Date of Report PRISCA	31/5/2023 5.2.0.13
Patient Data						
Name		MRS.	MEENAKSHI	Patient ID		012305290173
Birthday			1/1/1994	Sample ID		11650584
Age at term			30	Sample Date		29/5/2023
Gestational age			13+1			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg		Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom			12+1
PAPP-A	4.57	mIU/ml	0.61	Method		CRL (<>Robinson)
fb-hCG	50.8	ng/ml	1.42	Scan date		29/5/2023
Risks at sampling date				Crown rump length in mm 69		
Age Risk			1:708	Nuchal translu	cency MoM	0.81
Biochemical T21 risk			1:631	Nasal bone		Present
Combined trisomy 21 risk	_		1:3718			
Trisomy 13/18 + NT			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 1:10000				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 with NT test it is expected that among 3718 women with the same data, there is one woman with a trisomy 21 pregnancy and 3717 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!		
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