

Trisomy 13/18 + NT

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



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					Date of Report PRISCA	03-02-2020 5.0.2.37
Patient Data					TRISCIT	3.0.2.07
			MRS VIDHI	Patient ID		012002020039
Birthday			02-09-1981 Sample ID			10607122
Age at delivery			38.4	Sample Date		02/02/2020
Gestational age 12+2						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	75	Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+1
PAPP-A	4.65	mIU/ml	1.48	Method		CRL(<>Robinson
fb-hCG	110.2 ng/ml		2.46	Scan date		01-02-2020
Risks at sampling date				Crown rump le	ength in mm	55
Age Risk			1:119	Nuchal translucency MOM 0.9		0.96
Biochemical T21 risk			1:179	Nasal bone		Present
Combined Trisomy 21 Risk			1:734	Sonographer		DR.PRAKASH LALCHANDANI
Trisomy 13/18 + NT			<1:10000	Qualification in measuring NT		MD
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10				The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.		
1: 00 1:250				After the result of the Trisomy 21 test (with NT) it is expected that among more than 734 women with the same data, there is one woman with a trisomy 21		
1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				pregnancy and 733 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!		

Age

Risk above Age Risk

The laboratory cannot be hold responsible for their impact on

the risk assessment! Calculated risks have no diagnostic values

Risk below Age risk