

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



The laboratory cannot be hold responsible for their impact

on the risk assessment! Calculated risks have no diagnostic

Risk below Age risk

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					Date of Report PRISCA	28/7/2021 5.1.0.17
Patient Data						
Name MRS. ANJU DEVI				Patient ID		012107270032
Birthday			1/1/1985	Sample ID		11004745
Age at delivery			36.6	Sample Date		27/07/2021
Gestational age			12+5			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	e	12+4
PAPP-A	4.28	mIU/ml	0.89	Method		CRL(<>Robinson
fb-hCG	58.8	ng/ml	1.29	Scan date		26/07/2021
Risks at sampling date				Crown rump length in mm 61.2		
Age Risk			1:191	Nuchal translu	icency MOM	0.88
Biochemical T21 risk			1:514	Nasal bone		Present
Combined Trisomy 21 Risk			1:2563	Sonographer		DR. NITIN YADAV
Trisomy 13/18 + NT			<1:10000	Qualification is	n measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:250 Cut off 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age				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 2563 women with the same data, there is one woman with a trisomy 21 pregnancy and 2562 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!		

values

Risk above Age Risk