

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

Risk Above Cut Off

<1:10000, which indicates a low risk



on the risk assessment! Calculated risks have no diagnostic

Risk below Age risk

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Date of Report 15-02-2021 PRISCA 5.0.2.37

				Date of Report	13-02-2021
				PRISCA	5.0.2.37
Patient Data					
Name	MI	RS ANJALI F2	Patient ID		012102120143
Birthday		01-08-1997	Sample ID		10872621
Age at delivery		23.5	Sample Date		12/02/2021
Gestational age					
Correction factors					
Fetuses	2 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70 Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ago	e	13+0
PAPP-A	$5.1  \mathrm{mIU/ml}$	0.62	Method		CRL(<>Robinson
fb-hCG	124.5 ng/ml	1.37	Scan date		29-01-2021
Risks at sampling date					
Age Risk		1:1031			
Biochemical T21 risk		1:600			
Combined Trisomy 21 I	Risk	1:1032			
Trisomy 13/18 + NT		<1:10000			
Risk		1	Down's Syndrome Risk (Trisomy 21 Screening)		
1:10				d risk for Trisomy 21 (wit represents a low risk.	h NT) is below the
1:100 1:250 Cut off			After the result of the Trisomy 21 test (with NT) it is expected that among more than 1032 women with the same data, there is one woman with a trisomy 21		
1:1000 1:1000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			pregnancy and 1031 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 13 17 18 21 23 25 2	29 31 33 33 37 39 4	Age			
Trisomy 13/18 + NT				y cannot be hold responsib	_
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values

Risk above Age Risk