

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



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	30/4/2022 5.1.0.17
Patient Data						
Name MRS. MILKA YADAV				Patient ID		012204290041
Birthday 7/8/1993				Sample ID		11605983
Age at term 29.01				Sample Date		29/4/2022
Gestational age 13+5						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57	Diabetes		NO	Pregnancies	unknown
Smoker	moker NO Origin			Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	13+2
PAPP-A	7.28	mIU/ml	1.02	Method		CRL (<>Robinson)
fb-hCG	175.8	ng/ml	4.24	Scan date		26/4/2022
Risks at sampling date				Nuchal Translucency 1.15		
Age Risk			1:774	Nuchal Translucency MoM 0.64		
Biochemical T21 risk			1:151	Nasal Bone Preser		Present
Combined T21 risk			1:901			
Trisomy 13/18+NT			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:1000 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 test it is expected that among 901 women with the same data, there is one woman with a trisomy 21 pregnancy 900 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