

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	20/4/2022 5.1.0.17
Patient Data						
Name MRS.NISHA W/O PREM				Patient ID		012204190100
Birthday 11/8/1988				Sample ID		11449200
Age at term 34.01				Sample Date		19/4/2022
Gestational age 12+5						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	63.2	Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin			Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	e	12+5
PAPP-A	2.62	m <b>I</b> U/ml	0.58	Method		CRL (<>Robinson)
fb-hCG	28.6	ng/ml	0.64	Scan date		19/4/2022
Risks at sampling date				Nuchal Translucency 1.1		
Age Risk			1:360	Nuchal Translucency MoM 0.67		
Biochemical T21 risk			1:1613	Nasal Bone Presen		
Combined T21 risk			1:8671			
Trisomy 13/18+NT			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 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 it is expected that among 8671 women with the same data, there is one woman with a trisomy 21 pregnancy 8670 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