

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

					Date of Report PRISCA	8/9/2021 5.2.0.13
Patient Data						
Name	me MRS. JYOTI DEVI					012109080024
Birthday			28/11/1991	Sample ID		11040834
Age at delivery 30.32				Sample Date		8/9/2021
Gestational age 11+6						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	48.6	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	e	11+5
PAPP-A	4.63	mIU/ml	1	Method		CRL(<>Robinson
fb-hCG	101	ng/ml	1.94	Scan date		7/9/2021
Risks at sampling date				Nuchal translucency in mm 1.4		
Age Risk			1:929	Nuchal translu	cency MOM	1.01
Biochemical T21 risk			1:1209	Nasal bone		Present
Combined Trisomy 21 Ris	k		1:4760			
Trisomy 13/18 + NT			<1:10000	Qualification in measuring NT C		CR
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:1000 Cut off				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 4760 women with the same data, there is one woman with a trisomy 21 pregnancy and 4759 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