

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 PRISCA	27/4/2022 5.1.0.17
Patient Data					
Name	N	MRS. SHRUTI	Patient ID		012204260309
Birthday		30/03/1994	Sample ID		11169489
Age at term		29.09	Sample Date		26/4/2022
Gestational age		13+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	48 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ago	e	13+2
PAPP-A	3.29 mIU/ml	0.49	Method		CRL (<>Robinson)
fb-hCG	33.9 ng/ml	0.7	Scan date		26/4/2022
Risks at sampling date			Nuchal Translucency 1.03		
Age Risk		1:721	Nuchal Transl	ucency MoM	0.58
Biochemical T21 risk		1:1658	Nasal Bone		Present
Combined T21 risk		1:9611			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 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 9611 women with the same data, there is one woman with a trisomy 21 pregnancy 9610 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!		
Trisomy 13/18+NT	D: 19/10 / :/1	VICEA :	The laboratory cannot be hold responsible for their impact		

values

Risk above Age Risk