

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

0/1/9093

					Date of Report PRISCA	9/1/2023 5.1.0.17
Patient Data						
Name	MRS. PUJA KUMARI			Patient ID		012301070173
Birthday			11/7/1993	Sample ID		11494769
Age at term			29.11	Sample Date		7/1/2023
Gestational age			13+1			
Correction factors						
Fetuses	1 IV	VF		unknown	Previous trisomy 21	unknown
Weight in kg	51 D	iabetes		NO	Pregnancies	unknown
Smoker	NO C	Prigin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom			13+0
PAPP-A	3.46 m	nIU/ml	0.42	Method		CRL (<>Robinson)
fb-hCG	17.2 ng	g/ml	0.52	Scan date		7/1/2023
Risks at sampling date				Nuchal translu	cency	1.5
Age Risk			1:701	Nuchal translu	cency MoM	0.87
Biochemical T21 risk			1:1988	Nasal bone		Present
Combined trisomy 21 risk			<1:10000			
Trisomy 13/18 + NT			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 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 with NT test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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				The laboratory cannot be hold responsible for their impact		

values

Risk above Age Risk