

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

					Date of Report PRISCA	4/9/2021 5.2.0.13	
Patient Data							
Name			MRS. JWALA	Patient ID		012109020129	
Birthday			19/10/199	4 Sample ID		11087415	
Age at delivery			27.	4 Sample Date		2/9/2021	
Gestational age			13+	1			
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	70	Diabetes		NO	Pregnancies	unknown	
Smoker	NO	Origin		Asian			
Biochemical Data				Ultrasound Da	ata		
Parameter	Value	;	Corr Mom	Gestational ago	e	13+0	
PAPP-A	1.78	3 mIU/ml	0.4	Method		CRL(<>Robinson	
fb-hCG	50.1	ng/ml	1.34	Scan date		2/9/2021	
Risks at sampling date				Nuchal translu	Nuchal translucency in mm 1.3		
Age Risk			1:1217	Nuchal translu	icency MOM	0.74	
Biochemical T21 risk			1:398	Nasal bone		Present	
Combined Trisomy 21 Risk			1:2629	Sonographer		DR. RUBY RAHUL	
Trisomy 13/18 + NT			<1:10000	Qualification i	n measuring NT	CR	
Risk Risk				Down's Syndr	ome Risk (Trisomy 21 S	Screening)	
				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 2629 women with the same data, there is one woman with a trisomy 21 pregnancy and 2628 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					atory cannot be hold responsible for their impact		

values

Risk above Age Risk