

The calculated risk for Trisomy 13/18 (with NT) is < 1:10000,

Risk Above Cut Off

Trisomy 13/18 + NT

which indicates a low risk





The laboratory cannot be hold responsible for their impact on the risk

Risk below Age risk

assessment! Calculated risks have no diagnostic values

Risk above Age Risk

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				Date of Report PRISCA	5/2/2024 5.2.0.13
Patient Data					
Name		SHAADMA	Patient ID		012402030133
Birthday		4/1/1993	Sample ID		11818249
Age at sample		31.1	Sample Date		3/2/2024
Gestational age 13		13+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	65 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+5
PAPP-A	6.80 mIU/ml	0.87	Method		CRL (<>Robinson)
fb-hCG	34.2 ng/ml	1.42	Scan date		2/2/2024
Risks at sampling date			Crown rump length in mm 77		
Age Risk 1:585		1:585	Nuchal translucency MoM 1.0		
Biochemical T21 risk		1:1182	Nasal bone Pre		Present
Combined trisomy 21 risk		1:4674	Sonographer DR.SANGEI		DR.SANGEETA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT		MBBS
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:1000 1:10000 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 test (with NT) it is expected that among 4674 women with the same data, there is one woman with a trisomy 21 pregnancy and 4673 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! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).		