

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

which indicates a low risk

*Free Home Sample Collection 9999 778 778



risk assessment! Calculated risks have no diagnostic values

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Date of Report 03-01-2025 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. KAJAL			Patient ID		12501020048
Birthday	08-12-2024			Sample ID		11932874
Age at Sample date 20.1			Sample Date		02-01-2025	
Gestational age 12+5				5		
Correction factors						
Fetuses	1 I	VF		unknown	Previous trisomy 21	unknown
Weight in kg	51 Diabetes			NO	Pregnancies	unknown
Smoker	NO (Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	12+5
PAPP-A	5.3 n	nIU/ml	0.88	Method		CRL (<>Robinson)
fb-hCG	15.6 ng/ml		0.43	Scan date		02-01-2025
Risks at sampling date				Crown rump length in mm 62.9		
age Risk			1:1095	Nuchal translucency MoM		0.68
Biochemical T21 risk			<1:10000	Nasal bone		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer		DR. VIKASH GOYAL
Trisomy 13/18 + NT			<1:10000	Qualifications in measuring NT		MBBS
Risk				Down's Syndr	ome Risk (Trisomy 21	Screening)
1:100 1:100 1:100 1:1000				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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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		
13 15 17 19 21 23 25 2 Trisomy 13/18+NT The calculated risk for Triso				1998).	to accepted guidelines (Procannot be hold responsible	