

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 15-11-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	M	IRS. KIR	AN KUMARI	Patient ID		12411130121
Birthday			01-06-2005	Sample ID		11862880
Age at Sample date	ge at Sample date 19.5					13-11-2024
Gestational age 12+3				3		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	46	46 Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	11+6
PAPP-A	4.6	mIU/ml	0.65	Method		CRL (<>Robinson)
fb-hCG	28.6	ng/ml	0.68	Scan date		09-11-2024
Risks at sampling date				Crown rump l	ength in mm	51.1
Age Risk			1:1092	Nuchal translucency MoM 0.6		0.66
Biochemical T21 risk			1:5700	Nasal bone		PRESENT
Combined trisomy 21 risk	,		<1:10000	Sonographer		DR. ALKA JAIN
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	
				Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Gutoff				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		
1:1000 1:1000 1:1000 13:15:17:19:21:23:25:2 Trisomy 13/18+NT The calculated risk for Trisc			04143454740 is <1:10000	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:		