

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 18-10-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name	MRS. POO	NAM JUYAL	Patient ID		12410170027
Birthday		04-12-1988	Sample ID		11895481
Age at Sample date		35.9	Sample Date		17-10-2024
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	68 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+6
PAPP-A	6.8 mIU/ml	1.22	Method		CRL (<>Robinson)
fb-hCG	21.4 ng/ml	0.68	Scan date		16-10-2024
Risks at sampling date			Crown rump length in mm 65.8		
Age Risk	1:228		Nuchal translucency MoM 0.		0.78
Biochemical T21 risk		1:4948	Nasal bone		PRESENT
Combined trisomy 21 ris	sk	<1:10000	Sonographer		DR.
Trisomy 13/18 + NT		< 1:10000	Qualifications	in measuring NT	MBBS
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:10 1:100 1:250 Cut off			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		
1:1000 1:10000 13 15 17 19 21 23 25 3 Trisomy 13/18+NT The calculated risk for Tris			information pro the risk calculati diagnostic value! The patient con done according 1998).	vided by the referring physions are statistical aapproac	sician. Please note that ches and have no NT measurement was enat Diagn 18:511-523;