

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 20-12-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
n : . D :				IMSCA	3.2.0.13
Patient Data	) (DC CI		D. d ID		10410100100
Name	MRS. SU	JMAN LATA			12412190130
Birthday			Sample ID		11996145
Age at Sample date		30.3	Sample Date		19-12-2024
Gestational age		12+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	47 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age		12+1
PAPP-A	3.2  mIU/ml	0.52	Method		CRL (<>Robinson)
fb-hCG	24.45  ng/ml	0.54	Scan date		19-12-2024
Risks at sampling date			Crown rump le	ngth in mm	57.8
Age Risk		1:612	Nuchal transluc	ency MoM	0.99
Biochemical T21 risk		1:2793	Nasal bone		PRESENT
Combined trisomy 21 ris	sk	<1:10000	Sonographer		DR. DEEPIKA
Trisomy 13/18 + NT		<1:10000	Qualifications in	n measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Gtdff			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.		
1:1000 1:1000 13 15 17 19 21 23 25 2 Trisomy 13/18+NT The calculated risk for Tri			information provi the risk calculatio diagnostic value! The patient comb done according to 1998).	sk by PRISCA depends of ided by the referring phy ons are statistical aapproachined risk presumes that to accepted guidelines (Prunnot be hold responsible)	rsician. Please note that ches and have no  NT measurement was