

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 03-11-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. INDU BALA RAWAT			Patient ID		12411020067
Birthday		24-04-1998				11877715
Age at Sample date 36.5				Sample Date		02-11-2024
Gestational age			13+4			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	66	66 Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	13+4
PAPP-A	7.1	mIU/ml	1.01	Method		CRL (<>Robinson)
fb-hCG	51.2	ng/ml	1.93	Scan date		02-11-2024
Risks at sampling date				Crown rump length in mm 74.6		
Age Risk			1:199	Nuchal translu	icency MoM	0.98
Biochemical T21 risk			1:264	Nasal bone		PRESENT
Combined trisomy 21 ri	sk		1:1110	Sonographer		DR. VIKASH GOYAL
Trisomy 13/18 + NT <1:10000			Qualifications	in measuring NT	MBBS	
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
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 1110 women with the same data, there is one woman with a trisomy 21 pregnancy and 1109 women with not affected pregnancies.  The calculated risk by PRISCA depends on the accuracy of the		
1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000.				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).  The laboratory cannot be hold responsible for their impact on the		