

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

Book a Test Online www.molq.in

Date of Report 18-09-2024 PRISCA 5 9 0 13

<u> </u>	09160125 11966942
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Birthday 20-01-1990 Sample ID	11966942
	11000012
Age at Sample date 34.7 Sample Date 1	4-09-2024
Gestational age 11+1	
Correction factors	
Fetuses 1 IVF unknown Previous trisomy 21	unknown
Weight in kg 65 Diabetes NO Pregnancies	unknown
Smoker NO Origin Asian	

Sillokei	NO Oligili		Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+1
PAPP-A	3.4 mIU/ml	1.24	Method	CRL (<>Robinson)
fb-hCG	55.8 ng/ml	1.1	Scan date	14-09-2024
Risks at sampling date			Crown rump length in mm	42
Age Risk		1:277	Nuchal translucency MoM	0.52
Biochemical T21 risk		1:2156	Nasal bone	PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer	DR. RAVINDER KUMAR
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 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 1:1000 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			The laboratory cannot be hold responsible for their impact on the	