

\*Free Home Sample Collection 9999 778 778

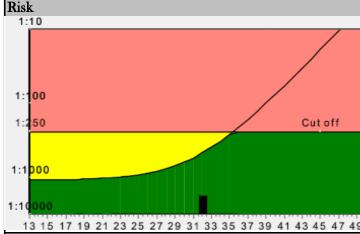


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Date of Report 27.06-2024

				PRISCA		5.2.0.13
Patient Data						
Name	]	MRS. NA	VI JAISWAL	Patient ID		12406260028
Birthday	17-05-1992			Sample ID		11814093
Age at Sample date	32.1			Sample Date		26-06-2024
Gestational age	12+0					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	71	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Da	ata	
Parameter	Value		Corr Mom	Gestational age	2	11+6

Parameter	Value	Corr Mom	Gestational age	11+6
PAPP-A	4.1 mIU/ml	1.14	Method	CRL (<>Robinson)
fb-hCG	45.4 ng/ml	1.12	Scan date	25-06-2024
Risks at sampling date			Crown rump length in mm	51
Age Risk		1:466	Nuchal translucency MoM	0.58
Biochemical T21 risk		1:2948	Nasal bone	PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer	DR. Abhishek
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
1:10			The calculated risk for Trisomy 2 cut off, which represents a low risk	



## Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates 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 the risk calculations are statistical approaches 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 risk assessment! Calculated risks have no diagnostic values