

*Free Home Sample Collection 9999 778 778

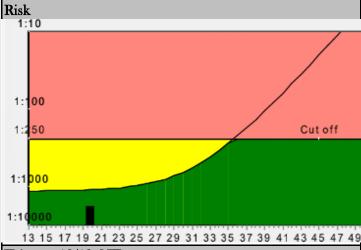


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Date of Report 11-09-2024 PRISCA 5.2.0.13

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Patient Data						
Name	MRS. SHALINI W/O ISHUVAR			Patient ID		12409100183
Birthday	27-12-2004			Sample ID		11966977
Age at Sample date	19.7			Sample Date		10-09-2024
Gestational age			13+2			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57.5	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Illtrasound De	ata	

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	13+1	
PAPP-A	$7.4~\mathrm{mIU/ml}$	0.98	Method	CRL (<>Robinson)	
fb-hCG	27.1 ng/ml	0.89	Scan date	09-09-2024	
Risks at sampling date			Crown rump length in mm	69.1	
Age Risk		1:1120	Nuchal translucency MoM	1.09	
Biochemical T21 risk		1:8692	Nasal bone	PRESENT	
Combined trisomy 21 risk		<1:10000	Sonographer	DR. AMENDA	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1.10			The calculated risk for Trisomy 21(v	vith NT) is below the	



Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

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 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 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 risk assessment! Calculated risks have no diagnostic values