

\*Free Home Sample Collection 9999 778 778

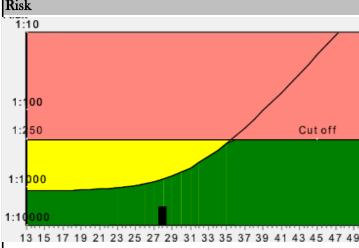


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Date of Report 23-09-2024 PRISCA 5 9 0 13

				PRISCA	
Patient Data					
Name	1	MRS. MEENAKSHI	F2 Patient ID		12409210299
Birthday		01-11-19	996 Sample ID	Sample ID	
Age at Sample date		2'	7.9 Sample Date	Sample Date	
Gestational age		12	2+6		
Correction factors					
Fetuses	2	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	65	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data		
arameter Value		Corr Mom	Gestational age	12+5	
PAPP-A	5.9 mIU/ml		Method	CRL (<>Robinson)	
fb-hCG	74.8 ng/ml	1.04	Scan date	21-09-2024	
Risks at sampling date			Crown rump length in mm	63.6	
Age Risk		1:812	Nuchal translucency MoM	0.80	
Biochemical T21 risk		1:1215	Nasal bone	PRESENT	
Combined trisomy 21 risk		1:7040	Sonographer	DR. DEEPIKA	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
Risk			Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21(with NT) is below the 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 7040 women with the same data, there is one woman with a trisomy 21 pregnancy and 7039 women with not affected pregnancies. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.

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