

*Free Home Sample Collection 9999 778 778



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Date of Report 28/09/2023 PRISCA 5.2.0.13

			PRISCA		5.2.0.13
Patient Data					
Name	MONIKA		Patient ID		012309270163
3 irthday 3/1/1990		Sample ID		11805787	
Age at sample 33.7		Sample Date 27/09/20		27/09/2023	
Gestational age 13+0					
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	87 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age 13+0		
PAPP-A	3.67 mIU/ml	0.89	Method		CRL (<>Robinson)
fb-hCG	35.5 ng/ml	1.21	Scan date		27/09/2023
Risks at sampling date			Crown rump length in mm 66		
Age Risk		1:360	Nuchal translucency MoM		0.59
Biochemical T21 risk 1:114		1:1142	Nasal bone Pres		Present
Combined trisomy 21 risk 1		1:6012	Sonographer		DR RAVINDER KUAMAR
Trisomy 13/18 + NT <		<1:10000	Qualifications in measuring NT		MBBS
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
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 6012 women with the same data, there is one woman with a trisomy 21 pregnancy and 6011 women with not affected pregnancies. 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		