

*Free Home Sample Collection 9999 778 778



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Date of Report 12/4/2022PRISCA 5.1.0.17

Patient Data					
Name MRS. PRIYA RANAWAT			Patient ID		012204110053
Birthday	21/03/1995		Sample ID		11443777
Age at term 27.07		Sample Date		11/4/2022	
Gestational age		13+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+3
PAPP-A	$5.6 \mathrm{mIU/ml}$	0.84	Method		CRL (<>Robinson)
fb-hCG	30.1 ng/ml	0.72	Scan date		10/4/2022
Risks at sampling date			Nuchal Translucency 1.2		
Age Risk		1:886	Nuchal Translucency MoM 0.6		0.66
Biochemical T21 risk		1:7611	Nasal Bone Preser		
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:100 1:250 Cut off 1:1000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 The calculated risk for Trisomy 13/18 is <1:10000, which indicates a low risk			The calculated risk for Trisomy 21 is below the cut off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy 9999 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!		