

*Free Home Sample Collection 9999 778 778



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Date of Report 14-03-19 PRISCA 5.0.2.37

					PRISCA	5.0.2.37
Patient Data						
Name			Mrs Naina	Patient ID		011903130036
Birthday			21-02-89	Sample ID		10443706
Age at delivery			30.5	Sample Date		13/03/2019
Gestational age by			14+4			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	56	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	13+3
AFP	27.5	ng/ml	0.86	Method		CRL
uE3	0.36 ng/ml		1.13	Scan Date		05-03-19
hCG	60145.30 mIU/ml		1.44	CRL measurments		73.1
Risks at sampling date				Nuchal translucency MoM 0.72		
Age Risk			1:905	Nasal bone		
Biochemical T21 Risk			1:1515	Sonographer		Dr Akshay Shukla
Combined Trisomy 21 R	lisk		1:9449	Qualifications	in measuring NT	MBBS, MD
Trisomy 13/18			<1:10000			
Risk				Down's Syndi	rome Risk (Trisomy 21	Screening)
1:10 1:250 1:1000 1:10000 1:10000 1:10000 1:10000 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 is < 1:10000, which represents a low risk.				The calculated risk for Trisomy 21 (is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 9449 women with the same data, there is one woman with a trisomy 21 pregnancy and 9448 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 risk calculations are statistical approaches and have no diagnostic value! The patient combined risk presumes the NT measurement was done according to accepted guidelines. The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!		