

*Free Home Sample Collection 9999 778 778



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Date of Report 11-03-19
PRISCA 5 0 2 37

					PRISCA	5.0.2.37
Patient Data						
Name Mrs Megha Kularia				Patient ID		011903090267
Birthday			18-03-89	Sample ID		10436337
Age at delivery			30.0	Sample Date		09/03/19
Gestational age by			13+4			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	76.4	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	13+3
PAPP-A	2.33	mIU/ml	0.57	Method		CRL (<>Hadlock)
fb-hCG	32.05	ng/ml	0.87	Scan Date		09-03-19
				CRL measurm	ents	72.9
Risks at sampling date				Nuchal translucency MoM 0.50		
Age Risk			1:671	Nasal bone	·	Present
Biochemical T21 Risk			1:1506	Sonographer		DR.SAHIL LOOMBA
Combined Trisomy 21 Risk			1:8509	Qualifications	in measuring NT	MBBS,DNB
Trisomy 13/18			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10				The calculated risk for Trisomy 21 (with nuchal		
				translucency) is below the cut off, which indicates a low risk.		
					t of the Trisomy 21 test (w	gith NT) it is
				After the result of the Trisomy 21 test (with NT) it is expected that among 8509 women with the same data,		
1:100				there is one woman with a trisomy 21 pregnancy and		
1:250 Out off				8508 women with not affected pregnancies.		
				The calculated risk by PRISCA depends on the accuracy of		
1.1000				the information provided by the referring physician. Please		
e e constante de la constante				note that risk calculations are statistical approach and have no diagnostic values!		
1.1900				The patient combined risk presumes the NT		
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age				measurement was done according to accepted guidelines		
Trisomy 13/18 + NT						
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.						
ansideency is > 1.10000, W	mich re	presents a	10 W 1 15K.			