

*Free Home Sample Collection 9999 778 778



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Date of Report 28-02-19
PRISCA 5 0 2 37

					PRISCA	5.0.2.37
Patient Data						
Name	me Mrs Monika					011902260331
Birthday	02-05-86			Sample ID		10433700
Age at delivery			32.8	Sample Date		26/02/19
Gestational age by			13+5			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	69.8	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	13+4
PAPP-A	5.63	mIU/ml	1.17	Method		CRL (⇔Hadlock)
fb-hCG	19.4	ng/ml	0.52	Scan Date		26-02-19
				CRL measurm	ents	76.2
Risks at sampling date				Nuchal translucency MoM 0.48		
Age Risk			1:438	Nasal bone		Present
Biochemical T21 Risk	<1:10000		Sonographer		DR.SAHIL LOOMBA	
Combined Trisomy 21 Risk			<1:10000	Qualifications	in measuring NT	MBBS,DNE
Trisomy 13/18			<1:10000			
Risk				Down's Syndi	rome 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.		
					. C.1 T.: 21	(: (1) ITE) : (:
1:100				After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21		
				The calculated risk by PRISCA depends on the accuracy of		
				the information provided by the referring physician. Please		
				note that risk calculations are statistical approach and have no		
The state of the s				diagnostic values! 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				The patient combined risk presumes the NT measurement was done according to accepted guidelines		
Trisomy 13/18 + NT			0.00000			
The calculated risk for trison						
translucency) is < 1:10000, which represents a low risk.						