Date of Report 13-04-19
PRISCA 5 0 2 37

					PRISCA	5.0.2.37
Patient Data						
Name		Mr	s. Priyanka	Patient ID		021904100007
Birthday			08-02-91	Sample ID		10470317
Age at delivery			28.2	Sample Date		10/04/2019
Gestational age by			13+3			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	68	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	12+0
PAPP-A	2.61	mIU/ml	0.58	Method		CRL (⇔Robinson)
fb-hCG	29.38	ng/ml	0.76	SCAN Date		02-04-19
				CRL measurm	ents	53.8
Risks at sampling date				Nuchal translucency MoM 0.70		
Age Risk			1:808	Nasal bone		present
Biochemical T21 risk	1:2532		Sonographer		DR.APARNA	
Trisomy 21 Risk			<1:10000	Qualifications	in measuring NT	C/G
Trisomy 13/18			<1:10000	-		
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:250 Cut off				The calculated risk for Trisomy 21 (is below the cut off, which indicates 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. 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 calculated risk for trisomy 13/18 is <1:10000, which

Trisomy 13/18 + NT

represents a low risk.