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Date of Report 13-01-19 PRISCA 5.0.2.37

				PRISCA	5.0.2.37
Patient Data					
ame Mrs PREETI YADAV			Patient ID		011901110108
Birthday		10-10-96	Sample ID		10396789
Age at delivery		22.3	Sample Date		11/01/2019
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	Diabetes		no	Pregnancies	
Smoker	no Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	12+5
PAPP-A	3.55 mIU/ml	0.91	Method		CRL (⇔Hadlock)
fb-hCG	95.36 ng/ml	2.17	Scan Date		27-11-18
Risks at sampling date			CRL		
Age Risk 1:10		1:1056	Nuchal translucency MoM		
Biochemical T21 Risk			Nasal Bone		Present
Combined Trisomy 21 Risk		1:826	Sonographer		
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	
Risk 1:10 1:100 1:250 Cut off			Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with nuchal translucency) 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 826 women with the same data, there is one woman with a trisomy 21 pregnancy and 825 women with no affected pregnancy. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please		
1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal			note that risk calculations are statistical approach and have no diagnostic values! The patient combined risk presumes the NT measurement was done according to accepted guidelines The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!		

risk.

The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low