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				Date of Report PRISCA	7/11/2019 5.0.2.37
Patient Data					
Name		Mrs Pooja Dev	i Patient ID		011911050428
Birthday		7/5/198	9 Sample ID		10249541
Age at delivery		30.	5 Sample Date		06/11/19
Gestational age		12+	4		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60 Diabe	tes	no	Pregnancies	
Smoker	no Origin	l	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	12+2
PAPP-A	2.29 mIU/r	nl 0.50	Method		CRL (<>Robinson)
fb-hCG	39.5 ng/ml	0.85	Scan Date		4/11/2019
Risks at sampling date			Crown Rump	Crown Rump Length (mm) 58.8	
Age Risk		1:607	Nuchal translucency MoM		0.87
Biochemical Trisomy 21 Risk		1:1014	Nasal Bone		present
Combined Trisomy 21 Risk		1:5644	Sonographer		DR. GOURAV JAIN
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT		C/R
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.		
1:100 1:250 1:1000 1:10000 1:10000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:250 Cut off Age			After the result of the Trisomy 21 test (with NT) it is expected that among 5644 women with the same data, there is one woman with a trisomy 21 pregnancy and 5643 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 the risk calculations are statistical aapproaches and have no diagnostic value!		
Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Risk	Above Cut Off		Risk above Ag	e Risk	Risk below Age risk