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				Date of Report PRISCA	22-12-18 5.0.2.37
Patient Data				TRISCA	5.0.2.37
Name Mrs Savita Devi			Patient ID		011812190216
Birthday		06-01-97	Sample ID		10415708
Age at delivery		22.0	Sample Date		21/12/2018
Gestational age		12+1			
<b>Correction factors</b>					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	55 Diabetes		no	Pregnancies	
Smoker	no Origin		Asian		
<b>Biochemical Data</b>		Ultrasound Data			
Parameter	Value	Corr Mom	Gestational ag	e	11+4
PAPP-A	3.26 mIU/ml	0.89	Method		CRL (<>Hadlock)
fb-hCG	54.85 ng/ml	1.14	Scan Date		17-12-18
Risks at sampling date			CRL		47
Age Risk		1:1042	Nuchal translucency MoM		1.25
Biochemical T21 Risk		1:3720	Nasal Bone		Present
Combined Trisomy 21 Ri	sk	1:7170	Sonographer		DR.ANKUSH DHANADIA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT		MD
Risk Risk 1:10 1:100 1:250 Cut off 1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			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 7170 women with the same data, there is one woman with a trisomy 21 pregnancy and 7169 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 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!		

