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				Date of Report PRISCA	6/9/2019 5.0.2.37
Patient Data					
Name		Mrs Lalit	a Patient ID		011909030343
Birthday		5/2/200	0 Sample ID		10558411
Age at delivery		19.	6 Sample Date		05/09/19
Gestational age		11+0	6		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	42 Diabo	etes	no	Pregnancies	
Smoker	no Origi	n	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	11+4
PAPP-A	8.12 mIU/	ml 1.56	Method		CRL (<>Robinson)
fb-hCG	141.36 ng/ml	2.46	Scan Date		3/9/2019
Risks at sampling date	e		Crown Rump Length (mm) 48.3		
Age Risk		1:1068	Nuchal translu	thal translucency MoM 0.7	
Biochemical Trisomy 21 Risk		1:1749	Nasal Bone	Vasal Bone prese	
Combined Trisomy 2	1:8825	Sonographer Dr. B.A.W.		Dr. B.A.WANI	
Trisomy 13/18 + NT		<1:10000	Qualification i	n measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10			TThe calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.		
1:100 1:250 Cut off 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 Age			After the result of the Trisomy 21 test (with NT) it is expected that among 8825 women with the same data, there is one woman with a trisomy 21 pregnancy and 8824 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		
R	isk Above Cut Off		Risk above Ag	e Risk	Risk below Age risk