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				Date of Report PRISCA	6/11/2019 5.0.2.37
Patient Data					
Name		Mrs Madhuri	Patient ID		011911040305
Birthday		30/03/1990	Sample ID		10572382
Age at delivery		29.6	Sample Date		04/11/19
Gestational age		11+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62 Diabetes		no	Pregnancies	
Smoker	no Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter Val	lue	Corr Mom	Gestational age	2	11+5
PAPP-A 3.	.65 mIU/ml	1.1	Method		CRL (<>Robinson)
fb-hCG 29	9.6 ng/ml	0.59	Scan Date		4/11/2019
Risks at sampling date			Crown Rump Length (mm) 49.3		
Age Risk		1:660	Nuchal translu	cency MoM	0.75
Biochemical Trisomy 21 Risk		<1:10000	Nasal Bone prese		present
Combined Trisomy 21 Risk		<1:10000	Sonographer DR. B		DR. B.A. WANI
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MD
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: 00 1:250 Cut off 1:1000 1:10000			After the result of the Trisomy 21 test (with NT) 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 <b>PRISCA</b> 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!		
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 translucency) is   <1:10000, which represents a low risk.					