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				Date of Report PRISCA	2/1/2020 5.0.2.37
Patient Data					
Name		Mrs Jyoti	Patient ID		051912510030
Birthday		10/9/1992	Sample ID		10501471
Age at delivery		27.3	Sample Date		31/12/19
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	71 Diabetes	5	no	Pregnancies	
Smoker	no Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+4
PAPP-A	3.75 mIU/ml	0.86	Method		CRL (<>Robinson)
fb-hCG	36.3 ng/ml	0.87	Scan Date		28/12/2019
Risks at sampling date			Crown Rump Length (mm) 62.3		
Age Risk		1:855	Nuchal translucency MoM (		0.75
Biochemical Trisomy 21 Risk		1:5277	Nasal Bone		present
Combined Trisomy 21 Risk		<1:10000	Sonographer		DR. KRISHNAN JAIN
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MD
Risk			Down's Syndro	ome Risk (Trisomy	21 Screening)
Risk 1:10				risk for Trisomy 21 (w ff, which indicates a lo	ith nuchal translucency) is w risk.
1:100 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.			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values Risk above Age Risk Risk below Age risk		