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Date of Report 01 - 03 - 2021PRISCA 5.0.2.37

Patient Data						
Name			MRS AARTI	Patient ID		012102280109
Birthday			07-08-1988	Sample ID		10848623
Age at delivery			32.6	Sample Date		28/02/2021
Gestational age			12+1			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	74	Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+0
PAPP-A	4.35	mIU/ml	1.44	Method		CRL(<>Robinson
fb-hCG	102.4	ng/ml	2.24	Scan date		27-02-2021
Risks at sampling date				Crown rump length in mm 52.4		
Age Risk			1:434	Nuchal translucency MOM 0.57		
Biochemical T21 risk			1:793	Nasal bone Prese		Present
Combined Trisomy 21 Risk			1:4026	Sonographer DR.APARN		DR.APARNA
Trisomy 13/18 + NT			<1:10000	Qualification in measuring NT M		
Risk 1:10				Down's Syndrome Risk (Trisomy 21 Screening)		
				The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.		
1:1000 1:10000 1:10000 1:10 1:10000				After the result of the Trisomy 21 test (with NT) it is expected that among more than 4026 women with the same data, there is one woman with a trisomy 21 pregnancy and 4025 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 NT) is <1:10000, which indicates a low risk Risk Above Cut Off				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		