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 Date of Report
 25-01-2021

 PRISCA
 5.0.2.37

Patient Data						
Name MRS JHARANA RANI F1			Patient ID		012101220163	
Birthday			01-01-1985	Sample ID		10793849
Age at delivery			36.1	Sample Date		22/01/2021
Gestational age 11+6						
Correction factors						
Fetuses	2	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59	Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	11+5
PAPP-A	3.4	mIU/ml	0.52	Method		CRL(<>Robinson
fb-hCG	35.2	ng/ml	0.32	Scan date		22-01-2021
Risks at sampling date				Crown rump length in mm 49		
Age Risk			1:208	Nuchal translucency MOM 1.		
Biochemical T21 risk		1:2433	Nasal bone Pres		Present	
Combined Trisomy 21 Risk			1:5995	Sonographer DR.SAG.		DR.SAGAR
Trisomy 13/18 + NT 1:1190			1:1190	Qualification in measuring NT MBBS,MD		
Risk 1:10				Down's Syndrome Risk (Trisomy 21 Screening)		
1:1000 1:1000 1:10000 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 NT) is 1:1190, which indicates a low risk			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among more than 5995 women with the same data, there is one woman with a trisomy 21 pregnancy and 5994 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! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values			