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 Date of Report
 25/10/2021

 PRISCA
 5.2.0.13

Patient Data					
Name	MRS. MAMTA		. Patient ID		012110240068
Birthday	7/5/1995		Sample ID		11040475
Age at delivery	ge at delivery 27.07		Sample Date 24/10/20		24/10/2021
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	67 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Da	ata	
Parameter	Value	Corr Mom	Gestational age	2	11+5
PAPP-A	3 mIU/ml	0.73	Method		CRL(<>Robinson
fb-hCG	58.1 ng/ml	1.44	Scan date		17/10/2021
Risks at sampling date			Nuchal translucency 1.3		
Age Risk		1:1243	Nuchal translucency MoM		0.93
Biochemical T21 risk		1:1663	Nasal bone		Present
Combined T21 risk	mbined T21 risk 1:8134		Qualification in measuring NT CR		
Trisomy 13/18 + NT <1:10000					
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 1:3 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 <1:10000, 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 it is expected that among 8134 women with the same data, there is one woman with a trisomy 21 pregnancy and 8133 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		