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	Sample Collection 9999 778 778

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				Date of Report PRISCA	17/8/2021 5.1.0.17
Patient Data					
Name	MRS	5. PRIYANKA	Patient ID		012108150037
Birthday		26/10/1990	Sample ID		11066727
Age at delivery		30.8	Sample Date		15/08/2021
Gestational age		13+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	48 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data		Ultrasound Data			
Parameter	Value	Corr Mom	Gestational age	e	13+1
PAPP-A	5.6 mIU/ml	0.62	Method		CRL(<>Robinson
fb-hCG	27.9 ng/ml	0.64	Scan date		11/8/2021
Risks at sampling date			Crown rump length in mm 70.4		
Age Risk		1:608	- Nuchal translu	cency MOM	1.08
Biochemical T21 risk		1:3124	Nasal bone		Present
Combined Trisomy 21 Risk	X	<1:10000	Sonographer		DR. ARSHBIR SINGH
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	C/R
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
Risk 1:10 1:700 1:250 Cut off			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 10000 women with the same data, there is one woman with a trisomy 21		
1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			pregnancy and 9999 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			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Risk Al	bove Cut Off		Risk above Ag	e Risk	Risk below Age risk