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Sample Collection
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				Date of Report PRISCA	21/8/2021 5.1.0.17
Patient Data					
Name	MRS. DIF	TY KUMARI	Patient ID		012108210050
Birthday		25/03/1996	Sample ID		11132508
Age at delivery		25.4	Sample Date		21/8/2021
Gestational age		12+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	81 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data		Ultrasound Da	ata		
Parameter	Value	Corr Mom	Gestational age	2	12+3
PAPP-A	5.3 mIU/ml	1.76	Method		CRL(<>Robinson
fb-hCG	22.4 ng/ml	0.52	Scan date		21/08/2021
Risks at sampling date			Crown rump length in mm 58.6		
Age Risk		1:940	Nuchal translucency MOM		0.72
Biochemical T21 risk		<1:10000	Nasal bone		Present
Combined Trisomy 21 Risk	X	<1:10000	Sonographer		DR. A. YADAV
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MBBS, MD
Risk		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. After the result of the Trisomy 21 test (with NT) it is		
1: 00 1:250 Cut off 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 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 approaches 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	oove Cut Off		Risk above Ag	e Risk	Risk below Age risk