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	Sample Collection 9999 778 778

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				Date of Report PRISCA	13-11-2020 5.0.2.37	
Patient Data						
Name		MRS RAJNI	Patient ID		052011110020	
Birthday		29-03-1992	Sample ID		10825140	
Age at delivery		28.6	Sample Date		11/11/2020	
Gestational age		13+1				
Correction factors				-		
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	54 Diabetes		NO	Pregnancies	unknown	
Smoker	NO Origin		Asian			
Biochemical Data			Ultrasound Data			
Parameter	Value	Corr Mom	Gestational age	2	13+1	
PAPP-A	2.83 mIU/ml	0.41	Method		CRL(<>Robinson	
fb-hCG	15.88 ng/ml	0.36	Scan date		09-11-2020	
Risks at sampling date			Crown rump length in mm 67.7			
Age Risk		1:775	Nuchal translu	cency MOM	0.71	
Biochemical T21 risk		1:3894	Nasal bone		Present	
Combined Trisomy 21 Ris	šk	<1:10000	Sonographer		DR.PRINCY SETHI	
Trisomy 13/18 + NT		1:8049	Qualification in	n measuring NT	MBBS	
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: 00 1:250 1:1000 1:1000 13 15 17 19 21 23 25 27 2 Trisomy 13/18 + NT		Age	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 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! The laboratory cannot be hold responsible for their impact			
The calculated risk for Trisomy 13/18 (with NT) is <1:8049, which indicates a low risk			on the risk assessment! Calculated risks have no diagnostic values			
Risk A	bove Cut Off		Risk above Ag	e Risk 📃 R	lisk below Age risk	