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				Date of Report PRISCA	06-12-2023 5.2.0.13
Patient Data					
Name		MRS PREETI	Patient ID		012312050080
Birthday		05-08-2000	Sample ID		11792253
Age at Sample date		23.3	Sample Date		05-12-2023
Gestational age		13+3	3		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+2
PAPP-A	4.76 mIU/m	ol 0.61	Method		CRL (<>Robinson)
fb-hCG	39.8 ng/ml	1.37	Scan date		04-12-2023
Risks at sampling date			Crown rump length in mm 71		
Age Risk	ge Risk 1:1052		Nuchal translucency MoM 0.50		
Biochemical T21 risk	ochemical T21 risk		Nasal bone PRESEN		PRESENT
Combined trisomy 21 risk 1:5890		1:5890	Sonographer DR PRAKAS		DR PRAKASH
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
1:10 1:100 1:250 1:1000 1:	•	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 5890 women with the same data, there is one woman with a trisomy 21 pregnancy and 5889 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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values			
Risk Above Cut Off Risk above Age Risk					