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					Date of Report PRISCA	28-01-2024 5.2.0.13
Patient Data					IMSCA	5.2.0.13
Name	MRS PRACHI					012401270087
Birthday				3 Sample ID		11812178
Age at Sample date			20.	5 Sample Date		27-01-2024
Gestational age 13+3				3		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	63 Diabetes			NO	Pregnancies	unknown
Smoker	NO	NO Origin				
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	9	13+2
PAPP-A	6.5	mIU/ml	0.92	Method		CRL (<>Robinson)
fb-hCG	17.5	ng/ml	0.62	Scan date		26-01-2024
Risks at sampling date				Crown rump length in mm 71		
Age Risk			1:1113	Nuchal translu	cency MoM	1.13
Biochemical T21 risk			<1:10000	Nasal bone		PRESENT
Combined trisomy 21 risk	Σ.		<1:10000	Sonographer		DR HARENDRA
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Kisk   1:100   1:250   Cut off   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   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 (with NT) it is expected that among 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 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 Risk below Age risk						