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					Date of Report PRISCA	17-09-2023 5.2.0.13
Patient Data					PRISCA	3.2.0.18
Name MRS POOJA ARORA				Patient ID		012309160078
Birthday						11805821
Age at Sample date) Sample Date		16-09-2023
Gestational age 12+5				5		
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	9	12+5
PAPP-A	4.9	mIU/ml	0.81	Method		CRL (<>Robinson)
fb-hCG	19.8	ng/ml	0.55	Scan date		16-09-2023
Risks at sampling date				Crown rump length in mm 65.4		
Age Risk			1:651	Nuchal translucency MoM 0.98		
Biochemical T21 risk			1:8843	Nasal bone PRESENT		
Combined trisomy 21 risk			<1:10000	Sonographer DR BHAWAN		
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Cut off 1:100 1:250 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000				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 among10000 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		
which indicates a low risk Image: Constraint of the second seco					e Risk	Risk below Age risk