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					Date of Report PRISCA	31-01-2024 5.2.0.13
Patient Data					TRISCH	0.2.0.10
Name MRS NEHA 1				Patient ID		012401300045
Birthday 24-08-1993				Sample ID		11827022
Age at Sample date 30.4			Sample Date		30-01-2024	
Gestational age 13+5				í		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	67 Diabetes			NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	13+1
PAPP-A	5.86	mIU/ml	0.81	Method		CRL (<>Robinson)
fb-hCG	18	ng/ml	0.72	Scan date		26-01-2024
Risks at sampling date				Crown rump length in mm 67.8		
Age Risk			1:636	Nuchal translu	cency MoM	0.99
Biochemical T21 risk	chemical T21 risk		1:5116	Nasal bone PRESEN		
Combined trisomy 21 risk	K		<1:10000	Sonographer		DR RAKESH
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
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		
	Above Cu	ıt Off		Risk above Ag	e Risk	Risk below Age risk