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					Date of Report PRISCA	02-01-2024 5.2.0.13
Patient Data					THISEIT	0.2.0.10
Name				Patient ID		012312310133
Birthday	10-10-			6 Sample ID		11827831
Age at Sample date			27.	2 Sample Date		31-12-2023
Gestational age 13+6				6		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62 Diabetes			NO	Pregnancies	unknown
Smoker	NO Origin			Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+6
PAPP-A	5.34	mIU/ml	0.64	Method		CRL (<>Robinson)
fb-hCG	18.2	ng/ml	0.74	Scan date		25-12-2023
Risks at sampling date				Crown rump length in mm 65.3		
Age Risk			1:884	Nuchal translu	Nuchal translucency MoM 0.54	
Biochemical T21 risk			1:3758	Nasal bone PRESENT		
Combined trisomy 21 risk			<1:10000	Sonographer DR ASHISH GAR		
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:			A	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 Ag	e Risk	Risk below Age risk