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Sample Collection
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				Date of Report PRISCA	11/3/2022 5.1.0.17	
Patient Data						
Name	MRS. MANI	NA DEVI (F2)	Patient ID		012203100027	
Birthday		10/7/1992	Sample ID		11257843	
Age at term 30.01		Sample Date		10/3/2022		
Gestational age						
Correction factors						
Fetuses	2 IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	60 Diabetes		NO	Pregnancies	unknown	
Smoker	NO Origin		Asian			
Biochemical Data			Ultrasound Data			
Parameter	Value	Corr Mom	Gestational age	e	13+4	
PAPP-A	5.28 mIU/ml	0.42	Method		CRL (<>Robinson)	
fb-hCG	27.2 ng/ml	0.31	Scan date		9/3/2022	
Risks at sampling date			Nuchal translucency 2			
Age Risk		1:700	Nuchal translucency MoM		1.08	
Biochemical T21 risk		1:4646	Nasal Bone		Presesnt	
Combined trisomy 21 risk <1:10		<1:10000				
Trisomy 13/18+NT		1:2914				
Risk			Down's Syndrome Risk (Trisomy 21 Screening)			
Risk 1:10 1:100 1:250 1:1000 1:1000 1:10000 1:10000 1:3 15 17 19 21 23 25 27 Trisomy 13/18+NT The calculated risk for T which indicates a low risk	risomy 13/18 with	Age	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 it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy 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 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						