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				Date of Report PRISCA	10/10/2022 5.1.0.17
Patient Data					
Name		MRS. SEEMA	Patient ID		012210090084
Birthday		5/7/1994	Sample ID		11590835
Age at term		28.09	Sample Date		9/10/2022
Gestational age		12+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	68 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	12+1
PAPP-A	5.68 mIU/ml	1.33	Method		CRL (<>Robinson)
fb-hCG	72.4 ng/ml	1.89	Scan date		8/10/2022
Risks at sampling date			Nuchal translucency 1.3		
Age Risk	ge Risk 1:770		Nuchal translucency MoM 0.89		
Biochemical T21 risk	risk 1:1874		Nasal bone Preser		
Combined trisomy 21 risl	ĸ	1:8793			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:250 1:250 1:1000 1:10			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 with NT test it is expected that among 8793 women with the same data, there is one woman with a trisomy 21 pregnancy and 8792 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		
<1:10000 , which indicat		1 IN I ) IS	on the risk ass values Risk above Ag		is have no diagnostic