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				Date of Report PRISCA	29/12/2022 5.1.0.17
Patient Data					
Name		MRS. AARTI	Patient ID		012212280011
Birthday		25/6/1995	Sample ID		11468016
Age at term		27.11	Sample Date		28/12/2022
Gestational age		11+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	53 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			11+3
PAPP-A	2.65 mIU/ml	0.63	Method		CRL (<>Robinson)
fb-hCG	40.3 ng/ml	0.82	Scan date		27/12/2022
Risks at sampling date			Nuchal translucency 1.2		
Age Risk 1:799		1:799	Nuchal translucency MoM 0.9		
Biochemical T21 risk	ochemical T21 risk 1:2593		Nasal bone Presen		
Combined trisomy 21 risk <		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:10 1:250 1:10000 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 more than 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
	k Above Cut Off		Risk above Ag	e Risk	Risk below Age risk