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				Date of Report PRISCA	4/2/2022 5.2.0.13
Patient Data					
Name	1	MRS. PREETI	Patient ID		012202020132
Birthday		16/08/1997	Sample ID		11431525
Age at delivery		24.5	Sample Date		2/2/2022
Gestational age		12+6			
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	Gestational age	2	12+5
PAPP-A	3.28 mIU/ml	0.56	Method		CRL (<>Robinson)
fb-hCG	46.3 ng/ml	0.99	Scan date		2/2/2022
Risks at sampling date			Nuchal translucency 1.7		
Age Risk		1:994	Nuchal translucency MoM 1.0		1.06
Biochemical T21 risk		1:1610	Nasal Bone		Present
Combined T21 risk		1:5947			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21 S	Screening)
Risk 1:10 1:250 Cut off 1:1000 1:250 Cut off 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000 1:10000 1:10000 1:			 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 5947 women with the same data, there is one woman with a trisomy 21 pregnancy and 5946 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 Age Risk 		