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				Date of Report PRISCA	18/6/2022 5.1.0.17
Patient Data					
Name	MRS. PREETI V	V/O ANIL(F2)	Patient ID		012206170157
Birthday		9/7/1992	Sample ID		11607989
Age at term		30.03	Sample Date		17/6/2022
Gestational age		13+0			
Correction factors					
Fetuses	2 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	12+5
PAPP-A	6.45 mIU/ml	0.45	Method		CRL(<>Robinson)
fb-hCG	41.5 ng/ml	0.56	Scan date		15/6/2022
Risks at sampling date			Nuchal Translucency 1		
Age Risk	1:662		Nuchal Translucency MoM 0.6		
Biochemical T21 risk		1:2011	Nasal Bone	Present	
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 1:250 Cut off 1:1000			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 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!		
13 15 17 19 21 23 25 2 Trisomy 13/18+NT The calculated risk for 7 <1:10000, which indicat Risk	Trisomy 13/18 (with	Age	The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values Risk above Age Risk		