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				Date of Report PRISCA	9/8/2022 5.1.0.17
Patient Data					
Name	MR	S. BHAWNA	Patient ID		012208080177
Birthday		25/01/1990	Sample ID		11566983
Age at term		32.11	Sample Date		8/8/2022
Gestational age		12+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	97 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+1
PAPP-A	2.94 mIU/ml	1.02	Method		CRL (<>Robinson)
fb-hCG	47.9 ng/ml	1.43	Scan date		6/8/2022
Risks at sampling date			Nuchal translucency 0.59		
Age Risk 1		1:441	Nuchal translucency MoM		0.41
Biochemical T21 risk	ochemical T21 risk 1:		Nasal bone		Present
Combined trisomy 21 ri	isk	1:6552			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:250 1:100 1:250 Cut off			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 6552 women with the same data, there is one woman with a trisomy 21 pregnancy and 6551 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		
Trisomy 13/18+NT The calculated risk for <1:10000 , which indica		Age	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 Risk below Age risk		