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				Date of Report PRISCA	17/8/2022 5.1.0.17
Patient Data					
Name	MRS. NUP	UR PANDEY	Patient ID		012208160075
Birthday		15/9/1990	Sample ID		11619157
Age at term		32.03	Sample Date		16/8/2022
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	71 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	12+4
PAPP-A	3.15 mIU/ml	0.66	Method		CRL (<>Robinson)
fb-hCG	17.8 ng/ml	0.53	Scan date		15/8/2022
Risks at sampling date			Nuchal translucency 1.1		
Age Risk	e Risk 1:494		Nuchal translucency MoM 0.		
Biochemical T21 risk	emical T21 risk 1:4517		Nasal bone	one Presen	
Combined trisomy 21 ris	sk	<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000			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!		
Trisomy 13/18+NT The calculated risk for 7 <1:10000 , which indica	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		