Patient Data					
Name MRS. MUMTAZ		Patient ID		012212100270	
Birthday		1/1/1994	Sample ID		11498915
Age at term		29.03	Sample Date		10/12/2022
Gestational age		13+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	7.5 Diabetes		NO	Pregnancies	unknown
Smoker I	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter Va	lue	Corr Mom			13+2
PAPP-A	6.5 mIU/ml	1	Method		CRL (<>Robinson)
fb-hCG	21.4 ng/ml	0.77	Scan date		9/12/2022
Risks at sampling date			Nuchal translucency 1.4		
Age Risk		1:751	Nuchal translucency MoM 0.		0.8
Biochemical T21 risk		1:8109	Nasal bone Presen		
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age  Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			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		