Patient data					
Name	М	RS. SIMRAN VERMA	Patient ID		
Birthday			Sample ID	2411003391/NOD	
Age at sample date		36.1	Sample Date	e 2/11/2024	
Gestational age		12 + 2	2		
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 no	
Weight	71.4	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound da	Ultrasound data	
Parameter	Value Corr. MoM		Gestational age 12 + 2		
PAPP-A	1.55 mIU/m	I 0.53	Method CRL Robinson		
fb-hCG	14.2 ng/ml	0.33	Scan date 2/11/2024		
Risks at sampling date				Crown rump length in mm 59.19	
Age risk 1:209			Nuchal translucency MoM 1.42		
Biochemical T21 risk 1:2610			Nasal bone present		
Combined trisomy 21 risk 1:3065			Sonographer		
Trisomy 13/18 + NT 1:558			Qualification	Qualifications in measuring NT	
Risk 1:10			Trisomy 21	Trisomy 21 The calculated risk for Trisomy 21 (with nuchal	
1::00 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:558, which represents a low risk.			translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 3065 women with the same data, there is one woman with a trisomy 21 pregnancy and 3064 women with not affected pregnancies. The free beta HCG level is low. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!		
translucency) is 1:558	8, which rep	resents a low risk.			

Sign of Physician