Patient data		-		
Name	MRS. BEANT KAUR		Patient ID	
Birthday	26-10-1993	Sample ID 2303220		2303220077/AMB
Age at sample date	29.3	Sample Date		02-03-2023
Gestational age	11 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 68	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 11 + 1		
PAPP-A 0.95 mIU/m	I 0.38	Method CRL Robinson		
fb-hCG 11.6 ng/ml	0.25	Scan date 25-02-2023		
Risks at sampling date	late		Crown rump length in mm 45	
Age risk	1:680	Nuchal translucency MoM		0.48
Biochemical T21 risk	1:3207			unknown
Combined trisomy 21 risk <1:10000		Sonographer		
Trisomy 13/18 + NT	1:1629	Qualifications in measuring NT MD   Trisomy 21		
1:10 1: 00 1: 250 1:1000 1:100000 1:100000 1:100000 1:100000 1:1000	The calculated risk for Trisomy 21 (with nuchal 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The free beta HCG level is low. The PAPP-A 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!			

Sign of Physician