Date of report: 21-06-2023

Patient data							
Name		MRS	S. ANJANA	Patient ID			
Birthday	20-06-1992			Sample ID 2306220465/AN		2306220465/AMB	
Age at sample date	31.0			Sample Date		19-06-2023	
Gestational age	12 + 2						
Correction factors							
Fetuses	1	IVF		no	Previous trisomy 21	no	
Weight	53	diabetes		no	pregnancies		
Smoker	no	Origin		Asian			
Biochemical data				Ultrasound da	ata		
Parameter	Value		Corr. MoM	Gestational	age	12 + 2	
PAPP-A	0.98 mIU/m	ıl	0.24	Method		CRL Robinson	
fb-hCG	25.1 ng/ml		0.52	Scan date		19-06-2023	
Risks at sampling date				Crown rump length in mm 59.			
Age risk				Nuchal translucency MoM 0.5			
Biochemical T21 risk				· · ·		present	
Combined trisomy 21 risk 1:1787				Sonographer			
Trisomy 13/18 + NT	-				Qualifications in measuring NT MD		
Risk 1:10				Trisomy 21 The calculated risk for Trisomy 21 (with nuchal			
1: 00 1: 50 Cut off 1: 1000 1: 1000 1: 10000 1: 10000 1: 10000 1: 10000 1: 10000 1: 10000 1: 10000 1: 10000 1: 10000 1: 1000 1: 25 27 29 31 33 35 37 39 41 43 45 47 49 Age				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 1787 women with the same data, there is one woman with a trisomy 21 pregnancy and 1786 women with not affected pregnancies. 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!			
Trisomy 13/18 + NT The calculated risk for	or Trisomy 1	3/18 (with	nuchal				
translucency) is 1:10							

Sign of Physician