Date of report: 3/10/2024

JITM Diagnostics

Patient data			
Name	MRS. ANIQA ARISH	Patient ID	
Birthday	20/03/1988	Sample ID 2410004597/NOD	
Age at sample date	36.5	Sample Date	e 3/10/2024
Gestational age	11 + 6		
Correction factors			
Fetuses 1	IVF	no Previous trisomy 21 no	
Weight 89.8	diabetes	no pregnancies	
Smoker no	Origin	Asian	
Biochemical data		Ultrasound data	
Parameter Value	Corr. MoM	Gestational	age 11 + 5
PAPP-A 1.88 mIU/m	I 1.05	Method CRL Robinson	
fb-hCG 33.1 ng/ml	0.77	Scan date 2/10/2024	
Risks at sampling date		Crown rump length in mm 53	
Age risk	1:186	Nuchal translucency MoM 1.49	
Biochemical T21 risk	1:2307	Nasal bone present	
Combined trisomy 21 risk	1:1835	Sonographer	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	
KISK 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal
1:100 1:1000 1:1100		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 1835 women with the same data, there is one woman with a trisomy 21 pregnancy and 1834 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 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