Prisca 5.2.0.13

Date of report: 20/12/2024

Patient data				
Name	MRS. MARIYAM	Patient ID		
Birthday	14/11/1996	Sample ID	2412033602/NOD	
Age at sample date	28.1	Sample Date	e 20/12/2024	
Gestational age	12 + 1			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	
Weight 71.2	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational	age 11 + 6	
PAPP-A 3.5 mIU/m		Method CRL Robinson		
fb-hCG 54.1 ng/ml	1.22	Scan date	18/12/2024	
Risks at sampling date		Crown rump length in mm 55		
Age risk	1:778	Nuchal translucency MoM 1.51		
Biochemical T21 risk	1:5070	Nasal bone present		
Combined trisomy 21 risk	21 risk 1:3403		Sonographer	
Trisomy 13/18 + NT	<1:10000	Qualification	Qualifications in measuring NT	
KISK 1:10		Trisomy 21		
1:1000 1:1000 1:1000 1:11000 13 15 17 19 21 23 25 27 29 31 33 35 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	Age s/18 (with nuchal	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 3403 women with the same data, there is one woman with a trisomy 21 pregnancy and 3402 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