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		20/12/2024
Gestational age		12 + 1			
Correction factors					
Fetuses 1	IVF		no	Previous trisomy 21	no
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	I	1.27	Method		CRL Robinson
fb-hCG 54.1 ng/ml		1.22			
			Crown rump length in mm 55		
Age risk					1.51
Biochemical T21 risk			·		present
-			Sonographe		
Trisomy 13/18 + NT		<1:10000	Qualification Trisomy 21	s in measuring NT	
1:10 1:250 Cut off 1:10000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			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