Prisca 5.2.0.13

Date of report: 13/09/2024

Patient data						
Name	MRS. NANCY			Patient ID		
Birthday	3/05/1999		Sample ID		2409026851/NOD	
Age at sample date	25.4			Sample Date 12/09/202		12/09/2024
Gestational age	11 + 5					
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	66	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound da	ata	
Parameter	Value Corr. MoM		Gestational age 11 + 4			
PAPP-A	1.27 mIU/m	าไ	0.52	Method		CRL Robinson
fb-hCG	84.1 ng/ml		1.76	Scan date		11/09/2024
Risks at sampling date	, 0			Crown rump length in mm 51.1		
Age risk	1:918			Nuchal translucency MoM 0.7		0.73
Biochemical T21 risk	1:319		Nasal bone		present	
Combined trisomy 21 r				Sonographer		
Trisomy 13/18 + NT	13/18 + NT <1:10000			Qualifications in measuring NT		
Risk			Trisomy 21			
1:10 1:250 1:1000 1:10000 1:110000 1:110000	Cut off  5 27 29 31 33 35 37 39 41 43 45 47 49  Age			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 2037 women with the same data, there is one woman with a trisomy 21 pregnancy and 2036 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!		
Trisomy 13/18 + NT The calculated risk for translucency) is < 1: risk.						