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	e	12/09/2024
Gestational age	nal 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	nl	0.52	Method CRL I		CRL Robinson
fb-hCG	84.1 ng/ml		1.76	Scan date		11/09/2024
Risks at sampling date				Crown rump length in mm 51.1		
Age risk	1:918			Nuchal translucency MoM 0.7		
Biochemical T21 risk	1:319			Nasal bone		present
Combined trisomy 21 risk 1:2037			Sonographer .			
Trisomy 13/18 + NT <1:10000				Qualifications in measuring NT		
Risk 1:10				Trisomy 21		
1:: 000 1::250 Cut off 1::10000 1::10000 13:15:17:19:21:23:25:27:29:31:33:35:37:39:41:43:45:47:49 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 trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.						