Prisca 5.2.0.13

Date of report: 29/10/2024

JITM Diagnostics

Patient data				
Name	MRS. NIKITA	Patient ID		
Birthday	19/05/2003	Sample ID	2410053727/NOD	
Age at sample date	21.4	Sample Date	29/10/2024	
Gestational age	13 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 37	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data			Ultrasound data	
Parameter Value	Corr. MoM	Gestational	age 13 + 5	
PAPP-A 8.79 mIU/m	0.76	Method	CRL Robinson	
fb-hCG 26.4 ng/ml	0.57	Scan date	28/10/2024	
Risks at sampling date			length in mm 80.5	
Age risk		Nuchal translucency MoM 1.01		
Biochemical T21 risk	<pre><1:10000 Nasal bone</pre>			
Combined trisomy 21 risk	 			
Trisomy 13/18 + NT	<1:10000	Qualification Trisomy 21	s in measuring NT M.D	
1:10 1:250 Cut off 1:1000 1:1000 1:1000 1:11000 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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