Prisca 5.2.0.13

Date of report: 1/11/2024

JITM Diagnostics

Patient data				
Name	MRS. VIBHUTI	Patient ID		
Birthday	23/10/1994		2411000197/NOD	
Age at sample date	30.0	Sample Date	e 1/11/2024	
Gestational age	13 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 48.6	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational	age 13 + 0	
PAPP-A 10.1 mIU/m	l 1.45	Method	CRL Robinson	
fb-hCG 81.1 ng/ml	1.83	Scan date	30/10/2024	
Risks at sampling date		Crown rump length in mm 70		
Age risk	1:661	Nuchal translucency MoM 0.74		
Biochemical T21 risk	1:2047	Nasal bone present		
-		Sonographer		
Trisomy 13/18 + NT			Qualifications in measuring NT	
7 1.0.1			Trisomy 21	
1:100 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 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