Prisca 5.2.0.13

Date of report: 30/11/2024

JITM Diagnostics

Patient data				
Name	MRS. NIVA			
Birthday	30/09/1991	Sample ID 2411053928/NOD		
Age at sample date	33.2		;	29/11/2024
Gestational age	12 + 0			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 49	diabetes	no pregnancies		
Smoker no	Origin	Asian		
ochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 11 + 6		
PAPP-A 2.3 mIU/m	I 0.58	Method CRL Robinson		
fb-hCG 19.2 ng/ml	0.38	Scan date 28/11/2024		
Risks at sampling date		Crown rump length in mm 55		
Age risk	1:387	Nuchal translucency MoM		1.10
Biochemical T21 risk	1:4776	Nasal bone		present
Combined trisomy 21 risk	<1:10000	Sonographe	r	DR. URVASHI JAIN
Trisomy 13/18 + NT	1:7336	Qualifications in measuring NT		M.D
KISK 1:10		Trisomy 21	ated risk for Trisomy 2	
1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 Trisomy 13/18 + NT The calculated risk for Trisomy 13 translucency) is 1:7336, which rep	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 free beta HCG level is low. 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