Prisca 5.2.0.13

Date of report: 7/11/2024

JITM Diagnostics

Patient data				
Name	MRS. KIRTI			
Birthday	3/09/1992			2411009704/NOD
Age at sample date	32.2	Sample Date	9	6/11/2024
Gestational age	13 + 3			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 63.9	diabetes	no pregnancies		
Smoker no	Origin	Asian		
Biochemical data	data		Ultrasound data	
Parameter Value	Corr. MoM	M Gestational age 13 + 2		
PAPP-A 0.65 mIU/m	l 0.12	Method CRL Robinson		
fb-hCG 49.1 ng/ml	1.24	Scan date 5/11/2024		
Risks at sampling date			Crown rump length in mm 74.3	
Age risk	1:486		Nuchal translucency MoM 0.7	
Biochemical T21 risk	>1:50			present
Combined trisomy 21 risk	risk 1:177		er	DR. NEERJA CHOPRA
Trisomy 13/18 + NT			s in measuring NT	M.D
Risk 1:10		Trisomy 21		
1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 33 Trisomy 13/18 + NT The calculated risk for Trisomy 13 translucency) is 1:97, which represents.	The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 test (with NT) it is expected that among 177 women with the same data, there is one woman with a trisomy 21 pregnancy and 176 women with not affected pregnancies. The PAPP-A 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