Prisca 5.2.0.13

Date of report: 26/10/2024

JITM Diagnostics

Patient data				
Name	MRS. SANGAM			
Birthday	15/10/2001 S			2410048975/NOD
Age at sample date	23.0)	26/10/2024
Gestational age	13 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 73	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	Ultrasound data			
Parameter Value	lue Corr. MoM Gestational age		13 + 1	
PAPP-A 2.28 mIU/m	I 0.53	Method CRL Robinson		
fb-hCG 49.1 ng/ml	1.27	Scan date 25/10/2024		
Risks at sampling date		Crown rump length in mm		71.6
Age risk	1:1056	· · · · · · · · · · · · · · · · · · ·		0.73
Biochemical T21 risk	1:845			present
Combined trisomy 21 risk				DR. NEERJA CHOPRA
Trisomy 13/18 + NT	<1:10000	Qualification Trisomy 21	is in measuring NT	M.D
1:10 1:250 1:1000 1:10000 1:110000 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000	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 5131 women with the same data, there is one woman with a trisomy 21 pregnancy and 5130 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!			

Sign of Physician