Prisca 5.2.0.13

Date of report: 10/10/2024

JITM Diagnostics

Patient data					
Name MRS.	me MRS. SHIVANGI SHARMA				
Birthday	21/02/1991			2410018387/NOD	
Age at sample date	33.6)	10/10/2024	
Gestational age	estational age 12 + 4				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 82	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data	U		Ultrasound data		
Parameter Value	Value Corr. MoM		age	12 + 3	
PAPP-A 1.38 mIU/m	0.49	Method CRL Robinson			
fb-hCG 50.1 ng/ml	1.23	Scan date 9/10/2024			
		Crown rump length in mm 61.1			
Age risk	1:361	Nuchal translucency MoM		0.44	
Biochemical T21 risk	1:258	Nasal bone Sonographer DR.		present	
Combined trisomy 21 risk				DR. P.K. ANAND MD	
Trisomy 13/18 + NT	<1:10000	1	s in measuring NT	MD	
Risk 1:10		Trisomy 21	ated risk for Trisomy 2		
1:1000 1:10000 1:110000 1:100000 1:100000 1:10000 1:100000 1:100000 1:100000 1:100000 1:100000 1:10000		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 1588 women with the same data, there is one woman with a trisomy 21 pregnancy and 1587 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