Prisca 5.2.0.13

Date of report: 12/10/2024

JITM Diagnostics

Patient data				
Name MRS. VARSHA SHARMA		Patient ID		
Birthday	29/09/1992		Sample ID 2410022374/NC	
Age at sample date	32.0		Sample Date 12/10/2024	
Gestational age 13 + 1				
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 74	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 13 + 0		
PAPP-A 1.51 mIU/m	ıl 0.38	Method CRL Robinson		
fb-hCG 62.8 ng/ml	1.60	Scan date 11/10/2024		
Risks at sampling date		Crown rump length in mm 6		69.9
Age risk	ge risk 1:492		Nuchal translucency MoM 0.	
Biochemical T21 risk	1:90	Nasal bone		present
Combined trisomy 21 risk	1:616	Sonographe	r DI	R. ASHISH GARG
Trisomy 13/18 + NT	<1:10000		s in measuring NT	M.D
Risk Trisomy 2				
1:10 1:250 1:10000 1:10000 1:110000 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 616 women with the same data, there is one woman with a trisomy 21 pregnancy and 615 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