Prisca 5.2.0.13

Date of report: 14/10/2024

JITM Diagnostics

Patient data				
Name	MRS. MINAL			
Birthday	12/06/1987			2410025049/NOD
Age at sample date	37.3)	14/10/2024
Gestational age	12 + 4			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	unknown
Weight 103	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoN	. MoM Gestational age		12 + 3
PAPP-A 0.85 mIU/m	I 0.41	Method CRL Robin		CRL Robinson
fb-hCG 41.3 ng/ml	1.07	Scan date 13/10/2024		
Risks at sampling date			length in mm	62.7
Age risk		1:157 Nuchal translucency MoM 1:92 Nasal bone		0.74
Biochemical T21 risk				unknown
Combined trisomy 21 risk			r 	DR. NEERJA CHOPRA
Trisomy 13/18 + NT	<1:10000	Qualification Trisomy 21	ns in measuring NT	M.D
1:100 1:250 1:1000 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 < 1:10000, which risk.	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 586 women with the same data, there is one woman with a trisomy 21 pregnancy and 585 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