Prisca 5.2.0.13

Date of report: 30/09/2024

JITM Diagnostics

Patient data				
Name	MRS. MONIKA	Patient ID		
Birthday	21/11/1992			2409063347/NOD
Age at sample date	31.9	Sample Date)	29/09/2024
Gestational age	11 + 3			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 2	1 no
Weight 53.2	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Value Corr. MoM Gestational ag		age	11 + 3
PAPP-A 1.65 mIU/m	I 0.61	Method CRL Robinson		
fb-hCG 77.1 ng/ml	1.46	Scan date 29/09/2024		
Risks at sampling date	Crown rump length in mm		49	
Age risk	1:475	Nuchal translucency MoM		1.05
Biochemical T21 risk	1:389	Nasal bone		present
Combined trisomy 21 risk	risk 1:1455		r	DR. GAGANDEEP KAUR
Trisomy 13/18 + NT	<1:10000		s in measuring NT	M.D
Risk 1:10		Trisomy 21		
1:1000 1:10000 1:10000 1:110000 1:10000 1:10000 1:10000 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low 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 1455 women with the same data, there is one woman with a trisomy 21 pregnancy and 1454 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