Prisca 5.2.0.13

Date of report: 11/10/2024

JITM Diagnostics

Patient data			
Name MRS	S. PRATIMA KUMARI	Patient ID	
Birthday	10/02/1998	Sample ID	2410019194/NOD
Age at sample date	26.7	Sample Date	10/10/2024
Gestational age	12 + 5		
Correction factors			
Fetuses 1	IVF	no	Previous trisomy 21 no
Weight 59.8	diabetes	no	pregnancies
Smoker no	Origin	Asian	
Biochemical data	Ultrasound data		
Parameter Value	Corr. MoM	Gestational	age 12 + 4
PAPP-A 3.51 mIU/ml	I 0.81	Method	CRL Robinson
fb-hCG 78.1 ng/ml	1.78	Scan date	9/10/2024
Risks at sampling date		Crown rump length in mm 63	
Age risk	1:885	,	
Biochemical T21 risk	1:879	Nasal bone	present
Combined trisomy 21 risk	·		·
Trisomy 13/18 + NT	<1:10000		s in measuring NT MD
Risk 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal
1:1000 1:10000 1:10000 1:110000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:10000000 1:100000000		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 5015 women with the same data, there is one woman with a trisomy 21 pregnancy and 5014 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