Prisca 5.2.0.13

Date of report: 24/09/2024

Patient data				
Name	MRS. AASHI JAIN	Patient ID		
Birthday	15/09/1995	Sample ID 2409050739/NOD		
Age at sample date	29.0	Sample Date	24/09/2024	
Gestational age	12 + 4			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 59.1	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data U		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 12 + 3		
PAPP-A 3.16 mIU/m	l 0.76	·I		
fb-hCG 52.1 ng/ml	1.16	Scan date 23/09/2024		
Risks at sampling date		Crown rump length in mm 62.2		
Age risk	1:723 N		Nuchal translucency MoM 0.81	
Biochemical T21 risk	1:1727	Nasal bone present		
Combined trisomy 21 risk 1:9390		Sonographer		
Trisomy 13/18 + NT <1:10000		Qualifications in measuring NT		
Risk		Trisomy 21		
1:10 1:250 Cut off 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 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 9390 women with the same data, there is one woman with a trisomy 21 pregnancy and 9389 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