Prisca 5.2.0.13

Date of report: 3/06/2024

Patient data			
Name	MRS. JOBA MANJHI		
Birthday	25/06/1993	Sample ID	10218722/NOD
Age at sample date	30.9	Sample Date	e 2/06/2024
Gestational age	13 + 0		
Correction factors			
Fetuses 1	IVF	no	Previous trisomy 21 no
Weight 48	diabetes	no	pregnancies
Smoker no	Origin	Asian	
Biochemical data		Ultrasound data	
Parameter Value	Corr. MoM	Gestational	age 11 + 1
PAPP-A 6.23 mIU/m	nl 0.99	Method	CRL Robinson
fb-hCG 30.1 ng/ml	0.65	Scan date	20/05/2024
Risks at sampling date		Crown rump length in mm 46.13	
Age risk	1:580	Nuchal translucency MoM 1.88	
Biochemical T21 risk	1:8754	Nasal bone present	
Combined trisomy 21 risk	1:1377	Sonographer	
Trisomy 13/18 + NT	1:6374	Qualifications in measuring NT	
Risk 1:10		Trisomy 21	
1:100 1:1000 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 1:6374, 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 1377 women with the same data, there is one woman with a trisomy 21 pregnancy and 1376 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!	