Prisca 5.2.0.13

Date of report: 9/08/2024

Patient data				
Name	MRS. SHEFALI			
Birthday	13/09/1996	Sample ID 2408015630/NOD		
Age at sample date	27.9	Sample Date 9/08/2024		
Gestational age	12 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 51	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data Ultrasound data		ata		
Parameter Value	Corr. MoM	// Gestational age 12 + 5		
PAPP-A 7.1 mIU/m	l 1.28	Method	CRL Robinson	
fb-hCG 73.1 ng/ml	1.60	Scan date	8/08/2024	
Risks at sampling date		Crown rump length in mm 66.6		
Age risk	1:811	Nuchal translucency MoM 0.77		
Biochemical T21 risk	1:2754	Nasal bone present		
Combined trisomy 21 risk	risk <1:10000		Sonographer	
Trisomy 13/18 + NT	<1:10000 G		Qualifications in measuring NT	
Risk	1	Trisomy 21		
1:1000 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:1000000 1:100000 1:100000 1:100000 1:100000 1:100000 1:1000000 1:1000000 1:10000000 1:100000000		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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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!		