Prisca 5.2.0.13

Date of report: 15/02/2025

Patient data				
Name MF	MRS. MOHINI TRIPATHI			
Birthday	2/04/1995		Sample ID 2502026	
Age at sample date	29.9		Sample Date 14/02/2025	
Gestational age	12 + 4			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 56	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ıta	
Parameter Value	Corr. MoM	Gestational age 12 + 3		
PAPP-A 3.46 mIU/n	nl 0.78	Method CRL Robinson		
fb-hCG 67.8 ng/ml	1.49	Scan date 13/02/2025		
Risks at sampling date		Crown rump length in mm 61		
Age risk	1:657	Nuchal translucency MoM 0.95		
Biochemical T21 risk	1:942	Nasal bone present		
Combined trisomy 21 risk	1:4370	Sonographer		
Trisomy 13/18 + NT	<1:10000			
1:100 1:250 1:1000 1:1000 1:100000 1:10000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000000	Trisomy 21 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 4370 women with the same data, there is one woman with a trisomy 21 pregnancy and 4369 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!			