Prisca 5.2.0.13

Date of report: 23/04/2025

Patient data				
Name	MRS. LAIBA	Patient ID		
Birthday	5/01/2003	Sample ID 2504045912/NOD		
Age at sample date	22.3	Sample Date	<b>;</b>	22/04/2025
Gestational age	12 + 4			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 70	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	Ultrasound data			
Parameter Value	Corr. MoM	Gestational age 12 + 3		
PAPP-A 1.6 mIU/m	0.47	Method		CRL Robinson
fb-hCG 31.9 ng/ml	0.75	Scan date 21/04/2025		
Risks at sampling date		Crown rump	Crown rump length in mm	
Age risk	1:1050	Nuchal translucency MoM 0.82		
Biochemical T21 risk	1:1950	Nasal bone present		
Combined trisomy 21 risk	,		r	
Trisomy 13/18 + NT	<1:10000	Qualification	is in measuring NT	
1:100 1:250 1:1000 1:11000 1:11000 1:1000 1:1000 1:10000 1	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 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!			

Sign of Physician