Prisca 5.2.0.13

Date of report: 27/04/2025

Patient data				
Name	MRS. MANJU			
Birthday	10/05/1995	Sample ID 2504		2504054807/NOD
Age at sample date	30.0	Sample Date	;	26/04/2025
Gestational age	13 + 0			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 65	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	Ultrasound data			
Parameter Value	Value Corr. MoM G		age	13 + 0
PAPP-A 4.56 mIU/m	I 1.03	Method CRL Robinson		
fb-hCG 26.8 ng/ml	0.65	Scan date 26/04/2025		
Risks at sampling date			Crown rump length in mm 70.	
Age risk	1:660	Nuchal translucency MoM		0.74
Biochemical T21 risk	<1:10000			present
Combined trisomy 21 risk	,		r	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		
1:100 1:250 1:1000 1:10000	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!			