Prisca 5.2.0.13

Date of report: 18/04/2025

JITM Diagnostics

Patient data					
Name	MRS. MEENU	Patient ID			
Birthday	8/07/1997	Sample ID 2504036690/NOD			
Age at sample date	27.8 S			18/04/2025	
Gestational age	12 + 6				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 72	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data			Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 12 + 5			
PAPP-A 4.9 mIU/m	l 1.33	Method CRL Robinson			
fb-hCG 53.1 ng/ml	1.30				
Risks at sampling date				65.4	
Age risk			Nuchal translucency MoM 1.0 Nasal bone prese		
Biochemical T21 risk				present	
Combined trisomy 21 risk	5		r		
Trisomy 13/18 + NT	Qualifications in measuring NT				
Risk 1:10		Trisomy 21	ted risk for Trisomy 21 (w	ith nuchal	
1:100 1:250 Cut off 1:1000			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