Prisca 5.2.0.13

Date of report: 28/01/2025

JITM Diagnostics

Patient data					
Name	MRS. SHALIKA				
Birthday	22/01/1996	Sample ID	2	501043180/NOD	
Age at sample date	29.0			27/01/2025	
Gestational age	11 + 6				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 75	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data		Ultrasound data			
Parameter Value	Corr. MoM	Gestational age 11 + 3			
PAPP-A 4.48 mIU/m	l 1.99	Method CRL Robinson			
fb-hCG 19.8 ng/ml	0.44				
				49	
Age risk			Nuchal translucency MoM 1.5		
Biochemical T21 risk	<1:10000	Nasal bone Sonographe		present	
Combined trisomy 21 risk	-				
Trisomy 13/18 + NT			Qualifications in measuring NT		
Risk 1:10		Trisomy 21	ited risk for Trisomy 21 (wit		
1: 100 1:250 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:	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