JITM Diagnostics

Patient data				
Name	MRS. KOMAL			
Birthday	12/07/2000	Sample ID 2503061532		2503061532/NOD
Age at sample date	24.7	Sample Date		30/03/2025
Gestational age	13 + 1			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 53	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Value Corr. MoM G		Gestational age 13 + 0	
PAPP-A 4.61 mIU/m	l 0.77	Method CRL Robinson		
fb-hCG 375 ng/ml	8.60	Scan date 29/03/2025		
Risks at sampling date		Crown rump length in mm		70
Age risk	1:993		Nuchal translucency MoM	
Biochemical T21 risk	1:106	Nasal bone		present
Combined trisomy 21 risk 1:423		Sonographe	r	DR. RUCHI JAIN
Trisomy 13/18 + NT	<1:10000	Qualification	s in measuring NT	MD
Risk 1:10 1:10 1:250 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000, which risk.	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 423 women with the same data, there is one woman with a trisomy 21 pregnancy and 422 women with not affected pregnancies. The free beta HCG level is high. 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