## JITM Diagnostics

Patient data					
Name	MRS. RUCHI	Patient ID			
Birthday	23/03/1986	Sample ID		2505022752/NOD	
Age at sample date	39.1			12/05/2025	
Gestational age	13 + 0				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 58.8	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data	data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 12 + 2			
PAPP-A 4.16 mIU/m	0.83	Method CRL Robinson			
fb-hCG 96.8 ng/ml	2.26				
Risks at sampling date			Crown rump length in mm		
Age risk	1:101	Nuchal trans	slucency MoM	1.16	
Biochemical T21 risk	1:59	Nasal bone Sonographer DR.		present	
Combined trisomy 21 risk				DR. VINOD KUMAR	
Trisomy 13/18 + NT			s in measuring NT	M.D	
Risk 1:10		Trisomy 21	ited risk for Trisomy 2		
1:1000 1:10	translucency) is above the cut off, which indicates an increased risk.  After the result of the Trisomy 21 test (with NT) it is expected that among 152 women with the same data, there is one woman with a trisomy 21 pregnancy and 151 women with not affected pregnancies.  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