Date of report: 26/03/2025

JITM Diagnostics

Patient data				
Name	MRS. PRIYANKA			
Birthday	8/03/2000	Sample ID		2503051330/NOD
Age at sample date	25.0	Sample Date		25/03/2025
Gestational age	13 + 5			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 67.5	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	ta Ultrasound data			
Parameter Value	Corr. MoM	Gestational age 13 + 4		
PAPP-A 2.38 mIU/m	I 0.43	Method CRL Robinsor		
fb-hCG 23.8 ng/ml	0.63	Scan date 24/03/2025		
Risks at sampling date		Crown rump length in mm 78.5		
Age risk	1:997		Nuchal translucency MoM	
Biochemical T21 risk	1:2055	Nasal bone		present
Combined trisomy 21 risk	<1:10000	Sonographer		DR. VINOD KUMAR
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.D
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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!			