Date of report: 20/05/2025

## JITM Diagnostics

Patient data				
Name	MRS. NEHA YADAV	Patient ID		
Birthday	10/10/1991	Sample ID 2505037904/NOD		
Age at sample date	33.6	Sample Date 19/05/202		19/05/2025
Gestational age	12 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 56	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data			Ultrasound data	
Parameter Value	Corr. MoM	Gestational	age	12 + 6
PAPP-A 3.85 mIU/m	I 0.77	Method CRL Robinson		
fb-hCG 25.3 ng/ml	0.57	Scan date 19/05/2025		
Risks at sampling date		Crown rump	length in mm	67
Age risk	1:367	Nuchal translucency MoM 0.65		
Biochemical T21 risk	1:4171	Nasal bone present		
Combined trisomy 21 risk	ined trisomy 21 risk <1:10000		Sonographer DR. AMIT F	
Trisomy 13/18 + NT	<1:10000	3		
1:100 1:250 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 3 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 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 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!			