Date of report: 27/03/2025

## JITM Diagnostics

Patient data				
Name	MRS. NEHA			
Birthday	17/03/1990			2503052967/NOD
Age at sample date	35.0	Sample Date		26/03/2025
Gestational age	13 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 67	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	Ultrasound data			
Parameter Value	Corr. MoM	Gestational age		13 + 5
PAPP-A 3.65 mIU/m	I 0.62	Method		CRL Robinson
fb-hCG 53.2 ng/ml	1.43	Scan date		25/03/2025
Risks at sampling date		Crown rump length in mm		79.9
Age risk	1:284 Nuchal translucency MoM		0.68	
Biochemical T21 risk	1:257	Nasal bone		present
Combined trisomy 21 risk	1:1524	Sonographer		DR. NEERJA CHOPRA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.D
Risk 1:10		Trisomy 21	ated risk for Trisomy	
1:100 1:250 1:1000 1:11000 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.	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 1524 women with the same data, there is one woman with a trisomy 21 pregnancy and 1523 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!			