Date of report: 7/04/2025

JITM Diagnostics

Patient data				
Name	MRS. MANITA			
Birthday	11/01/1993	•		2504010817/NOD
Age at sample date	32.2	Sample Date		6/04/2025
Gestational age	13 + 3			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 74	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	nta	
Parameter Value	Corr. MoM	Gestational age		13 + 2
PAPP-A 3.28 mIU/m	I 0.74	Method		CRL Robinson
fb-hCG 40.1 ng/ml	1.06	Scan date		5/04/2025
Risks at sampling date		Crown rump length in mm		74.9
Age risk	1:481	Nuchal translucency MoM		0.70
Biochemical T21 risk	1:1320	Nasal bone		present
Combined trisomy 21 risk	1:7170	Sonographer		DR. NEERJA CHOPRA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT Trisomy 21		M.D
1:10 1:1000 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000	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 7170 women with the same data, there is one woman with a trisomy 21 pregnancy and 7169 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!			