Date of report: 10/06/2025

## J.I.T.M. DIAGNOSTICS

Patient data				
Name	MRS. MANISHA	Patient ID		
Birthday	5/01/1988	Sample ID	2506014603/NOD	
Age at sample date	37.4	Sample Date	e 9/06/2025	
Gestational age	13 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 77	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	L		Ultrasound data	
Parameter Value	Corr. MoM	Gestational	age 13 + 6	
PAPP-A 4.6 mIU/m	0.93	Method	CRL Robinson	
fb-hCG 35.5 ng/ml	0.99	Scan date	9/06/2025	
Risks at sampling date		Crown rump	o length in mm 82	
Age risk	1:162	Nuchal translucency MoM 0.72		
Biochemical T21 risk	1:872	Nasal bone present		
Combined trisomy 21 risk	1:4418	Sonographer DR. AMIT RAI		
Trisomy 13/18 + NT	<1:10000	1 3		
1:1000 1:250		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 4418 women with the same data, there is one woman with a trisomy 21 pregnancy and 4417 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!		