

J.I.T.M. DIAGNOSTICS

Patient data			
Name	MRS. ANITA	Patient ID	
Birthday	14/07/1998	Sample ID	2504037671/NOD
Age at sample date	26.8	Sample Date	18/04/2025
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	72.9	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.47 mIU/ml	0.55	
fb-hCG	45.1 ng/ml	1.02	
Risks at sampling date			
Age risk	1:861		
Biochemical T21 risk	1:1245		
Combined trisomy 21 risk	1:7258		
Trisomy 13/18 + NT	<1:10000		
			Gestational age 12 + 0
			Method CRL Robinson
			Scan date 17/04/2025
			Crown rump length in mm 56.93
			Nuchal translucency MoM 0.67
			Nasal bone present
			Sonographer DR. GAURAV KUMAR
			Qualifications in measuring NT M.D
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 7258 women with the same data, there is one woman with a trisomy 21 pregnancy and 7257 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!			
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician