

JITM Diagnostics

Patient data			
Name	MRS. MAMTA	Patient ID	
Birthday	10/07/1995	Sample ID	2507010784/NOD
Age at sample date	30.0	Sample Date	7/07/2025
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	43	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.2 mIU/ml	1.46	11 + 5
fb-hCG	27.6 ng/ml	0.52	Method
			CRL Robinson
			Scan date
			4/07/2025
Risks at sampling date			Crown rump length in mm
			52
Age risk		1:637	Nuchal translucency MoM
Biochemical T21 risk		<1:10000	1.08
Combined trisomy 21 risk		<1:10000	Nasal bone
Trisomy 13/18 + NT		<1:10000	present
			Sonographer
			DR. P.S. UPPAL
			Qualifications in measuring NT
			M.D
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!
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