

JITM Diagnostics

Patient data			
Name	MRS. KAMINI	Patient ID	
Birthday	2/10/1989	Sample ID	2504059585/NOD
Age at sample date	35.6	Sample Date	29/04/2025
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	74.6	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.42 mIU/ml	1.12	
fb-hCG	46.1 ng/ml	1.18	
Risks at sampling date			
Age risk		1:245	
Biochemical T21 risk		1:1317	
Combined trisomy 21 risk		1:6532	
Trisomy 13/18 + NT		<1:10000	
		Gestational age 13 + 0	
		Method CRL Robinson	
		Scan date 28/04/2025	
		Crown rump length in mm 69	
		Nuchal translucency MoM 0.75	
		Nasal bone present	
		Sonographer DR. NEERJA CHOPRA	
		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 6532 women with the same data, there is one woman with a trisomy 21 pregnancy and 6531 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