

JITM Diagnostics

Patient data			
Name	MRS. ILMA TAHIR		Patient ID
Birthday	12/08/1992		Sample ID
Age at sample date	32.9		Sample Date
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	66	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.15 mIU/ml	0.72	
fb-hCG	24.5 ng/ml	0.59	
Risks at sampling date			
Age risk	1:425		
Biochemical T21 risk	1:3848		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
			Gestational age
			12 + 6
			Method
			CRL Robinson
			Scan date
			18/06/2025
			Crown rump length in mm
			68
			Nuchal translucency MoM
			0.81
			Nasal bone
			present
			Sonographer
			DR. SUNIL SUNEJA
			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 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