

JITM Diagnostics

Patient data			
Name	MRS. MANISHA		Patient ID
Birthday	1/01/1983		Sample ID
Age at sample date	42.4		Sample Date
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	88	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.49 mIU/ml	1.11	12 + 1
fb-hCG	18.1 ng/ml	0.44	Method
Risks at sampling date			CRL Robinson
Age risk	1:40		Scan date
Biochemical T21 risk	1:1730		2/06/2025
Combined trisomy 21 risk	1:7487		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		58.3
			Nuchal translucency MoM
			0.79
			Nasal bone
			present
			Sonographer
			DR. SANJAY SHERKAR
			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 7487 women with the same data, there is one woman with a trisomy 21 pregnancy and 7486 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