

JITM Diagnostics

Patient data			
Name		MRS. POOJA SHARMA	
Patient ID			
Birthday		18/09/1994	
Sample ID		2504048181/NOD	
Age at sample date		30.6	
Sample Date		23/04/2025	
Gestational age		13 + 2	
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.48 mIU/ml	0.63	Gestational age
fb-hCG	25.6 ng/ml	0.62	Method
Risks at sampling date			CRL Robinson
Age risk			20/04/2025
Biochemical T21 risk			Crown rump length in mm
Combined trisomy 21 risk			67.7
Trisomy 13/18 + NT			Nuchal translucency MoM
			1.28
			Nasal bone
			present
			Sonographer
			DR. ATEESH SINGHAL
			Qualifications in measuring NT
			M.D
Risk		Trisomy 21	
1:10		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
1:100		After the result of the Trisomy 21 test (with NT) it is expected that among 6461 women with the same data, there is one woman with a trisomy 21 pregnancy and 6460 women with not affected pregnancies.	
1:250		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
1:1000		Please note that risk calculations are statistical approaches and have no diagnostic value!	
1:10000		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
Cut off		The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Age			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
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