

JITM Diagnostics

Patient data			
Name	MRS. NEHA SHARMA		Patient ID
Birthday	11/07/1992		Sample ID
Age at sample date	32.9		Sample Date
Gestational age	13 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	92.5	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.84 mIU/ml	0.98	13 + 4
fb-hCG	27.1 ng/ml	0.80	Method
Risks at sampling date			CRL Robinson
Age risk	1:435		Scan date
Biochemical T21 risk	1:4307		29/05/2025
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		78.7
			Nuchal translucency MoM
			0.79
			Nasal bone
			present
			Sonographer
			DR. VINOD KUMAR
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.	
1:250		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!	
1:1000		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
1:10000		The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Cut off			
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