

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	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>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.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician