

JITM Diagnostics

Patient data			
Name	MRS. MANSI	Patient ID	
Birth day	25/12/2000	Sample ID	2502029001/NOD
Age at sample date	24.1	Sample Date	15/02/2025
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	43.5	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.1 mIU/ml	0.81	Gestational age 12 + 1
fb-hCG	98.6 ng/ml	2.00	Method CRL Robinson
Risks at sampling date			Scan date 11/02/2025
Age risk		1:1001	Crown rump length in mm 58
Biochemical T21 risk		1:755	Nuchal translucency MoM 0.99
Combined trisomy 21 risk		1:3235	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. URVASHI JAIN
			Qualifications in measuring NT M.D
Risk		Trisomy 21	
		<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 3235 women with the same data, there is one woman with a trisomy 21 pregnancy and 3234 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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician