

## JITM Diagnostics

Patient data			
Name	MRS. MANSI		Patient ID
Birthday	25/12/2000		Sample ID
Age at sample date	24.1		Sample Date
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	Gestational age
PAPP-A	5.1 mIU/ml	0.81	12 + 1
fb-hCG	98.6 ng/ml	2.00	Method
Risks at sampling date			CRL Robinson
Age risk	1:1001		Scan date
Biochemical T21 risk	1:755		11/02/2025
Combined trisomy 21 risk	1:3235		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		58
			Nuchal translucency MoM
			0.99
			Nasal bone
			present
			Sonographer
			DR. URVASHI JAIN
			Qualifications in measuring NT
			M.D
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