

JITM Diagnostics

Patient data				
Name	MRS. RIMPY (A)		Patient ID	2502031234/NOD (A)
Birthday	29/08/1997		Sample ID	2502031234/NOD (A)
Age at sample date	27.5		Sample Date	16/02/2025
Gestational age	12 + 5			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	55	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	5.38 mIU/ml	0.60	Method	CRL Robinson
fb-hCG	98.9 ng/ml	1.01	Scan date	16/02/2025
Risks at sampling date			Crown rump length in mm	65
Age risk	1:836		Nuchal translucency MoM	1.02
Biochemical T21 risk	1:1551		Nasal bone	present
Combined trisomy 21 risk	1:6270		Sonographer	DR. GAGANDEEP KAUR
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	M.D
Risk			Trisomy 21	
			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 6270 women with the same data, there is one woman with a trisomy 21 pregnancy and 6269 women with not affected pregnancies. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
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

JITM Diagnostics

Patient data			
Name	MRS. RIMPY (B)	Patient ID	2502031234/NOD (B)
Birthday	29/08/1997	Sample ID	2502031234/NOD (B)
Age at sample date	27.5	Sample Date	16/02/2025
Gestational age	13 + 1		
Correction factors			
Fetuses	2	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.38 mIU/ml	0.51	13 + 1
fb-hCG	98.9 ng/ml	1.06	Method
			CRL Robinson
			Scan date
			16/02/2025
			Crown rump length in mm
			72
			Nuchal translucency MoM
			0.89
			Nasal bone
			present
			Sonographer
			DR. GAGANDEEP KAUR
			Qualifications in measuring NT
			M.D
Risks at sampling date		Trisomy 21	
Age risk	1:848	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 4943 women with the same data, there is one woman with a trisomy 21 pregnancy and 4942 women with not affected pregnancies. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Biochemical T21 risk	1:902		
Combined trisomy 21 risk	1:4943		
Trisomy 13/18 + NT	<1:10000		
Risk Age			
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