

Patient data			
Name	MRS. POOJA (A)	Patient ID	2506006805/NOD(A)
Birthday	12/08/1998	Sample ID	2506006805/NOD(A)
Age at sample date	26.8	Sample Date	4/06/2025
Gestational age	12 + 3		
Correction factors			
Fetuses	2	IVF	no
Weight	74	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.31 mIU/ml	0.96	
fb-hCG	42.5 ng/ml	0.46	
Risks at sampling date		Gestational age	
Age risk	1:868	12 + 2	
Biochemical T21 risk	<1:10000	Method	
Combined trisomy 21 risk	<1:10000	CRL Robinson	
Trisomy 13/18 + NT	<1:10000	Scan date	
		3/06/2025	
		Crown rump length in mm	
		59.1	
		Nuchal translucency MoM	
		0.78	
		Nasal bone	
		present	
		Sonographer	
		..	
		Qualifications in measuring NT	
		..	
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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