

Patient data			
Name	MRS. ANSHIKA		Patient ID
Birthday	31/01/2001	Sample ID	2502032956/NOD
Age at sample date	24.0	Sample Date	17/02/2025
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.1 mIU/ml	1.43	Gestational age 13 + 2
fb-hCG	47.8 ng/ml	1.19	Method CRL Robinson
			Scan date 17/02/2025
Risks at sampling date			Crown rump length in mm 73
Age risk	1:1023		Nuchal translucency MoM 0.72
Biochemical T21 risk	1:8766		Nasal bone present
Combined trisomy 21 risk	<1:10000		Sonographer DR. AMIT RAI
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT MD
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 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