

Patient data			
Name		Mrs. SANDHYA	
Patient ID		0942511120001	
Birthday		12-11-2000	
Sample ID		B3690701	
Age at sample date		25.0	
Sample Date		12-11-2025	
Gestational age		12 + 2	
Correction factors			
Fetuses	1	IVF	no
Weight	49	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.32 mIU/mL	0.51	
fb-hCG	40.01 ng/mL	0.84	
Gestational age		12 + 2	
Method		CRL Robinson	
Scan date		12-11-2025	
Crown rump length in mm		60.1	
Nuchal translucency MoM		0.96	
Nasal bone		present	
Sonographer		N A	
Qualifications in measuring NT		MD	
Risks at sampling date		Trisomy 21	
Age risk	1:953	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 8192 women with the same data, there is one woman with a trisomy 21 pregnancy and 8191 women with not affected pregnancies. 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	1:1716		
Combined trisomy 21 risk	1:8192		
Trisomy 13/18 + NT	<1:10000		
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

 below cut off
 Below Cut Off, but above Age Risk
 above cut off