

Patient data			
Name		Mrs. SAYLI PATIL	
Patient ID		0662409140106	
Birthday		02/05/98	
Sample ID		A0990321	
Age at sample date		26.4	
Sample Date		14/09/24	
Gestational age		13 + 1	
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.09 mIU/mL	1.19	
fb-hCG	54.39 ng/mL	1.45	
Gestational age		12 + 6	
Method		CRL Robinson	
Scan date		12/09/24	
Crown rump length in mm		68	
Nuchal translucency MoM		0.87	
Nasal bone		present	
Sonographer		N A	
Qualifications in measuring NT		MD	
Risks at sampling date		Trisomy 21	
Age risk	1:914	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
Biochemical T21 risk	1:3480	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.	
Combined trisomy 21 risk	<1:10000	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!	
Trisomy 13/18 + NT	<1:10000	The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
Qualifications in measuring NT		MD	
Risk		The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
1:10			
1:100			
1:250			
1:1000			
1:10000			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
Age			
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