

Patient data			
Name		Mrs. KANAK SHARMA	
Patient ID		0522505020015	
Birthday		15/06/00	
Sample ID		b2781597	
Age at sample date		24.9	
Sample Date		02/05/25	
Gestational age		13 + 1	
Correction factors			
Fetuses	1	IVF	no
Weight	71	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.54 mIU/mL	0.62	
fb-hCG	45.36 ng/mL	1.33	
Gestational age		13 + 0	
Method		CRL Robinson	
Scan date		01/05/25	
Crown rump length in mm		69	
Nuchal translucency MoM		0.72	
Nasal bone		present	
Sonographer		N A	
Qualifications in measuring NT		MD	
Risks at sampling date		Trisomy 21	
Age risk	1:987	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 6106 women with the same data, there is one woman with a trisomy 21 pregnancy and 6105 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	1:1040		
Combined trisomy 21 risk	1:6106		
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