

Patient data			
Name	MRS. RUPINDER KAUR	Patient ID	
Birth day	18-10-1996	Sample ID	2301220420/AMB
Age at sample date	26.3	Sample Date	20-01-2023
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	59.7	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.68 mIU/ml	1.63	12 + 1
fb-hCG	175 ng/ml	3.92	Method
Risks at sampling date			CRL Robinson
Age risk		1:903	Scan date
Biochemical T21 risk		1:450	17-01-2023
Combined trisomy 21 risk		1:2386	Crown rump length in mm
Trisomy 13/18 + NT		<1:10000	58
			Nuchal translucency MoM
			0.66
			Nasal bone
			present
			Sonographer
			..
			Qualifications in measuring NT
			MD
Risk		Trisomy 21	
1:10		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
1:100		After the result of the Trisomy 21 test (with NT) it is expected that among 2386 women with the same data, there is one woman with a trisomy 21 pregnancy and 2385 women with not affected pregnancies.	
1:250		The free beta HCG level is high.	
1:1000		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
1:10000		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!	
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.			

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off