

Patient data			
Name	Mrs. NIDHI KUMARI	Patient ID	0662511120089
Birthday	02-02-1990	Sample ID	B3725941
Age at sample date	35.8	Sample Date	12-11-2025
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	51	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.4 mIU/mL	0.29	Gestational age 12 + 3
fb-hCG	39.76 ng/mL	0.91	Method CRL Robinson
Risks at sampling date			Scan date 11-11-2025
Age risk	1:229		Crown rump length in mm 61
Biochemical T21 risk	1:67		Nuchal translucency MoM 0.63
Combined trisomy 21 risk	1:457		Nasal bone unknown
Trisomy 13/18 + NT	1:2878		Sonographer N A
			Qualifications in measuring NT MD
Risk			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 457 women with the same data, there is one woman with a trisomy 21 pregnancy and 456 women with not affected pregnancies. The PAPP-A level is low. 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 1:2878, which represents a low risk.			

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