

Patient data			
Name	Mrs. PRAJAKTA VEER	Patient ID	0662509210055
Birthday	28/03/97	Sample ID	B3407189
Age at sample date	28.5	Sample Date	21/09/25
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	40.5	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.51 mIU/mL	0.70	Gestational age 12 + 0
fb-hCG	42 ng/mL	0.77	Method CRL Robinson
Risks at sampling date			Scan date 21/09/25
Age risk	1:747		Crown rump length in mm 56.5
Biochemical T21 risk	1:3699		Nuchal translucency MoM 0.94
Combined trisomy 21 risk	<1:10000		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer NA
			Qualifications in measuring NT NA
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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 < 1:10000, which represents a low risk.			

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