

Patient data			
Name	Mrs. MANISHA PATIL		Patient ID
Birthday	09-10-1984		Sample ID
Age at sample date	38.8		Sample Date
Gestational age	12 + 4		
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	Gestational age
PAPP-A	4.25 mIU/mL	0.89	12 + 3
fb-hCG	45.66 ng/mL	1.05	Method
			CRL Robinson
			Scan date
			13-07-2023
Risks at sampling date			Crown rump length in mm
Age risk			61.6
Biochemical T21 risk			Nuchal translucency MoM
1:474			0.82
Combined trisomy 21 risk			Nasal bone
1:2444			unknown
Trisomy 13/18 + NT			Sonographer
<1:10000			NA
			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 2444 women with the same data, there is one woman with a trisomy 21 pregnancy and 2443 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 held 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.			

Sign of Physician