

ANANTKAWALAS

Patient data			
Name			Patient ID 0702305090002
Birthday	21-03-1993	Sample ID	24262243
Age at sample date	30.1	Sample Date	08-05-2023
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	90	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 6
PAPP-A	3.67 mIU/mL	1.34	Method CRL Robinson
fb-hCG	37.07 ng/mL	1.08	Scan date 08-05-2023
Risks at sampling date			Crown rump length in mm 68.3
Age risk	1:643		Nuchal translucency MoM 0.70
Biochemical T21 risk	1:6112		Nasal bone present
Combined trisomy 21 risk	<1:10000		Sonographer ANANTKAWALAS
Trisomy 13/18 + NT	<1:10000		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 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.			

Sign of Physician

■ below cut off

■ Below Cut Off, but above Age Risk

■ above cut off