

Patient data			
Name	Mrs. B.KRISHNAVENI	Patient ID	0352211240015
Birthday	01-01-2000	Sample ID	23263348
Age at sample date	22.9	Sample Date	23-11-2022
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no Previous trisomy 21 unknown
Weight	45	diabetes	no pregnancies
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.53 mIU/mL	0.31	12 + 2
fb-hCG	45.65 ng/mL	0.93	Method CRL Robinson
Risks at sampling date		Scan date	
Age risk	1:1025	Crown rump length in mm	59.06
Biochemical T21 risk	1:344	Nuchal translucency MoM	0.78
Combined trisomy 21 risk	1:2324	Nasal bone	unknown
Trisomy 13/18 + NT	<1:10000	Sonographer	NA
Risk		Qualifications in measuring NT	
		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 2324 women with the same data, there is one woman with a trisomy 21 pregnancy and 2323 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 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

 below cut off

 Below Cut Off, but above Age Risk

 above cut off