

Prisca 5.1.0.17
Date of report: 15-11-2024

NA

Patient data			
Name	Mrs. KRITESHWARI	Patient ID	0622411120094
Birthday	13-07-1996	Sample ID	A1635126
Age at sample date	28.3	Sample Date	12-11-2024
Gestational age	12 + 5		
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	4.89 mIU/mL	0.95	
fb-hCG	83.15 ng/mL	1.97	
Risks at sampling date		Gestational age	
Age risk		12 + 5	
Biochemical T21 risk		Method	
1:777		CRL Robinson	
Combined trisomy 21 risk		Scan date	
1:858		12-11-2024	
Trisomy 13/18 + NT		Crown rump length in mm	
1:3265		65	
<1:10000		Nuchal translucency MoM	
		1.02	
		Nasal bone	
		present	
		Sonographer	
		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 3265 women with the same data, there is one woman with a trisomy 21 pregnancy and 3264 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 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