

Patient data					
Name	Mrs. E. POOJITHA		Patient ID	0042306240001	
Birthday	01-10-2002		Sample ID	20030191	
Age at sample date	20.7		Sample Date	23-06-2023	
Gestational age	13 + 1				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	67	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 1	
PAPP-A	1.96 mIU/mL	0.44	Method	CRL Robinson	
fb-hCG	35.65 ng/mL	1.03	Scan date	23-06-2023	
Risks at sampling date			Crown rump length in mm	72	
Age risk	1:1100		Nuchal translucency MoM	0.84	
Biochemical T21 risk	1:879		Nasal bone	unknown	
Combined trisomy 21 risk	1:5290		Sonographer	NA	
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 5290 women with the same data, there is one woman with a trisomy 21 pregnancy and 5289 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