

Patient data			
Name Mrs. W/O L/NK RAVIKANT BHATT FETUA-A		Patient ID 0032105170182	
Birthday 15-10-1992		Sample ID K2448542	
Age at sample date 28.6		Sample Date 17-05-2021	
Gestational age 11 + 3			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 48.6	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 3
PAPP-A	2.49 mIU/mL	0.43	Method CRL Robinson
fb-hCG	68.62 ng/mL	0.56	Scan date 17-05-2021
Risks at sampling date			Crown rump length in mm 48.2
Age risk 1:723			Nuchal translucency MoM 0.99
Biochemical T21 risk 1:1929			Nasal bone present
Combined trisomy 21 risk 1:8706			Sonographer DR 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 8706 women with the same data, there is one woman with a trisomy 21 pregnancy and 8705 women with not affected pregnancies. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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