

Patient data			
Name		Mrs. PRANITHA TWIN A	
Patient ID		0012509180007	
Birthday		14/08/92	
Sample ID		B3616062 TWIN A	
Age at sample date		33.1	
Sample Date		17/09/25	
Gestational age		12 + 0	
Correction factors			
Fetuses	2	IVF	no
Weight	78	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.02 mIU/mL	1.63	Gestational age
fb-hCG	47.27 ng/mL	0.50	Method
Risks at sampling date			CRL Robinson
Age risk			1:392
Biochemical T21 risk			<1:10000
Combined trisomy 21 risk			<1:10000
Trisomy 13/18 + NT			<1:10000
Scan date			17/09/25
Crown rump length in mm			55.9
Nuchal translucency MoM			0.88
Nasal bone			present
Sonographer			NA
Qualifications in measuring NT			NA
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 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.			

 below cut off
 Below Cut Off, but above Age Risk
 above cut off