

Patient data			
Name	Mrs. MAYA	Fetus A	Patient ID 0012111290522
Birthday		04/02/97	Sample ID 22655558
Age at sample date		24.8	Sample Date 29/11/21
Gestational age		13 + 0	
Correction factors			
Fetuses	2	IVF	yes
Weight	40	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 4
PAPP-A	11.04 mIU/mL	0.78	Method CRL Robinson
fb-hCG	95.33 ng/mL	1.02	Scan date 26/11/21
Risks at sampling date			Crown rump length in mm 64.7
Age risk		1:985	Nuchal translucency MoM 1.03
Biochemical T21 risk		1:3406	Nasal bone present
Combined trisomy 21 risk		<1:10000	Sonographer Dr Rajani Peethala MBBS., DGO.,
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT MBBS., DGO.,
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.			

Sign of Physician

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