

Patient data			
Name	Mrs. MAYA Fetus B	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	
PAPP-A	11.04 mIU/mL	0.78	
fb-hCG	95.33 ng/mL	1.02	
Risks at sampling date			
Age risk		1:985	
Biochemical T21 risk		1:3406	
Combined trisomy 21 risk		<1:10000	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 4 Method CRL Robinson Scan date 26/11/21 Crown rump length in mm 63.5 Nuchal translucency MoM 0.92 Nasal bone present Sonographer Dr Rajani Peethala MBBS., DGO., Qualifications in measuring NT MBBS., DGO.,
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