

Patient data					
Name	Mrs. G.MOUNIKA FETUS-A		Patient ID	0352204100060	
Birthday	10-05-1998		Sample ID	22664721	
Age at sample date	23.9		Sample Date	10-04-2022	
Gestational age	11 + 5				
Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	51	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	11 + 2	
PAPP-A	2 mIU/mL	0.32	Method	CRL Robinson	
fb-hCG	51.22 ng/mL	0.45	Scan date	07-04-2022	
Risks at sampling date			Crown rump length in mm	46.82	
Age risk	1:973		Nuchal translucency MoM	1.33	
Biochemical T21 risk	1:1632		Nasal bone	present	
Combined trisomy 21 risk	1:2745		Sonographer	DR NA	
Trisomy 13/18 + NT	1:973		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 2745 women with the same data, there is one woman with a trisomy 21 pregnancy and 2744 women with not affected pregnancies. The PAPP-A level is low. 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 1:973, which represents a low risk.					

Sign of Physician