

Patient data			
Name	Mrs. C E MOUNIKA	Patient ID	0012210020130
Birthday	26-04-1999	Sample ID	22472952
Age at sample date	23.4	Sample Date	02-10-2022
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.72 mIU/mL	0.56	
fb-hCG	45.65 ng/mL	1.22	
Risks at sampling date			
Age risk		1:1034	
Biochemical T21 risk		1:1037	
Combined trisomy 21 risk		1:6180	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 6
			Method CRL Robinson
			Scan date 01-10-2022
			Crown rump length in mm 68.41
			Nuchal translucency MoM 0.58
			Nasal bone unknown
			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 6180 women with the same data, there is one woman with a trisomy 21 pregnancy and 6179 women with not affected pregnancies. 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