

Patient data					
Name	Mrs. SHAREEFA MAIMOONA YAMANI		Patient ID	0012203260315	
Birthday	10-06-1992		Sample ID	23264525	
Age at sample date	29.8		Sample Date	26-03-2022	
Gestational age	11 + 4				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	63	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	11 + 4	
PAPP-A	1.67 mIU/mL	0.68	Method	CRL Robinson	
fb-hCG	59.14 ng/mL	1.17	Scan date	26-03-2022	
Risks at sampling date			Crown rump length in mm	50.3	
Age risk	1:638		Nuchal translucency MoM	0.96	
Biochemical T21 risk	1:1155		Nasal bone	unknown	
Combined trisomy 21 risk	1:5322		Sonographer	DR NA	
Trisomy 13/18 + NT	<1:10000		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 5322 women with the same data, there is one woman with a trisomy 21 pregnancy and 5321 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