

Patient data					
Name	Mrs. SANDHYA		Patient ID	0012206140435	
Birthday	24/01/85		Sample ID	23271800	
Age at sample date	37.4		Sample Date	14/06/22	
Gestational age	12 + 3				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	unknown
Weight	57	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 3	
PAPP-A	5.7 mIU/mL	1.41	Method	CRL Robinson	
fb-hCG	39.55 ng/mL	0.91	Scan date	14/06/22	
Risks at sampling date			Crown rump length in mm	62	
Age risk	1:155		Nuchal translucency MoM	0.81	
Biochemical T21 risk	1:2384		Nasal bone	unknown	
Combined trisomy 21 risk	<1:10000		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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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