

Patient data		Patient ID	
Name	Mrs. SANGHAMITRA ROUTRAY	Patient ID	0652405240192
Birthday	06-10-1990	Sample ID	A0711867
Age at sample date	33.6	Sample Date	24-05-2024
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	83	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.143 mIU/mL	1.89	Gestational age
fb-hCG	56.98 ng/mL	1.51	Method
Risks at sampling date		Scan date	
Age risk	1:362	Crown rump length in mm	
Biochemical T21 risk	1:2844	Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000	Nasal bone	
Trisomy 13/18 + NT	<1:10000	Sonographer	
Trisomy 21		Qualifications in measuring NT	
RISK 		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 held 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

