

Patient data					
Name	Mrs. V.SANDHYA W/O. KRTISHNA		Patient ID	0312405060028	
Birthday	02-06-1990		Sample ID	A0015616	
Age at sample date	33.9		Sample Date	06-05-2024	
Gestational age	12 + 2				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	61	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 2	
PAPP-A	5.01 mIU/mL	1.42	Method	CRL Robinson	
fb-hCG	60.14 ng/mL	1.37	Scan date	06-05-2024	
Risks at sampling date			Crown rump length in mm	59	
Age risk	1:337		Nuchal translucency MoM	1.17	
Biochemical T21 risk	1:2059		Nasal bone	present	
Combined trisomy 21 risk	1:4837		Sonographer	N A	
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 4837 women with the same data, there is one woman with a trisomy 21 pregnancy and 4836 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 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

below cut off

Below Cut Off, but above Age Risk

above cut off