

Prisca 5.1.0.17
Date of report: 20/12/25

Patient data			
Name Mrs. SANDHYA SINGH W/O SUBHAM		Patient ID 0012512200252	
Birthday 15/07/02		Sample ID B4370079	
Age at sample date 23.4		Sample Date 17/12/25	
Gestational age 11 + 4			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 70	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 3
PAPP-A	1.31 mIU/mL	0.60	Method CRL Robinson
fb-hCG	48.03 ng/mL	0.98	Scan date 16/12/25
Risks at sampling date			Crown rump length in mm 49.65
Age risk 1:983			Nuchal translucency MoM 0.93
Biochemical T21 risk 1:1983			Nasal bone present
Combined trisomy 21 risk 1:9746			Sonographer NA
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT Sonographer
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 9746 women with the same data, there is one woman with a trisomy 21 pregnancy and 9745 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