

Prisca 5.1.0.17
Date of report: 31/12/25

Patient data			
Name	Mrs. SANDHYA	Patient ID	0772512300051
Birthday	01/07/89	Sample ID	B3668937
Age at sample date	36.5	Sample Date	30/12/25
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	51	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.33 mIU/mL	0.64	
fb-hCG	32.96 ng/mL	0.94	
Risks at sampling date			
Age risk		1:200	
Biochemical T21 risk		1:512	
Combined trisomy 21 risk		1:1321	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 4
			Method CRL Robinson
			Scan date 24/12/25
			Crown rump length in mm 62.9
			Nuchal translucency MoM 1.17
			Nasal bone present
			Sonographer NA
			Qualifications in measuring NT Sonographer
Risk		Trisomy 21	
The graph shows the risk of Trisomy 21 as a function of maternal age. The y-axis represents risk from 1:10 to 1:10000. The x-axis represents age from 13 to 49. A red shaded area represents the risk above the cut-off, a yellow shaded area represents the risk below the cut-off but above the age risk, and a green shaded area represents the risk below the cut-off. A black vertical line indicates the patient's age risk at 36.5 years.		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 1321 women with the same data, there is one woman with a trisomy 21 pregnancy and 1320 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off