

Prisca 5.1.0.17
Date of report: 05-10-2024

NA

Patient data			
Name	Mrs. A.SANDHYA	Patient ID	0352410010040
Birthday	06-07-1999	Sample ID	23204288
Age at sample date	25.2	Sample Date	01-10-2024
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.98 mIU/mL	0.73	13 + 4
fb-hCG	47.41 ng/mL	1.51	Method
			CRL Robinson
			Scan date
			30-09-2024
Risks at sampling date			Trisomy 21
Age risk	1:989		Crown rump length in mm
Biochemical T21 risk	1:1165		79
Combined trisomy 21 risk	1:6451		Nuchal translucency MoM
Trisomy 13/18 + NT	<1:10000		0.84
			Nasal bone
			unknown
			Sonographer
			NA
			Qualifications in measuring NT
			MD
			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 6451 women with the same data, there is one woman with a trisomy 21 pregnancy and 6450 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