

Prisca 5.1.0.17
Date of report: 30-06-2025

NA

Patient data			
Name	Mrs. SRAVANI	Patient ID	0312506300026
Birthday	02-04-1988	Sample ID	b2920205
Age at sample date	37.2	Sample Date	30-06-2025
Gestational age	11 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.56 mIU/mL	1.31	11 + 0
fb-hCG	54.28 ng/mL	0.94	Method
			CRL Robinson
			Scan date
			30-06-2025
Risks at sampling date			Trisomy 21
Age risk	1:151		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:1854		After the result of the Trisomy 21 test (with NT) it is expected that among 7967 women with the same data, there is one woman with a trisomy 21 pregnancy and 7966 women with not affected pregnancies.
Combined trisomy 21 risk	1:7967		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!
Trisomy 13/18 + NT	<1:10000		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