

Prisca 5.1.0.17
Date of report: 13-01-2025

NA

Patient data			
Name	Mrs. M.RAJASREE	Patient ID	0312501120024
Birthday	29-01-1986	Sample ID	a2081088
Age at sample date	39.0	Sample Date	12-01-2025
Gestational age	13 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.12 mIU/mL	0.88	Gestational age 13 + 6
fb-hCG	13.88 ng/mL	0.49	Method CRL Robinson
			Scan date 12-01-2025
Risks at sampling date			Trisomy 21
Age risk	1:109		Crown rump length in mm 82.4
Biochemical T21 risk	1:2274		Nuchal translucency MoM 0.87
Combined trisomy 21 risk	<1:10000		Nasal bone present
Trisomy 13/18 + NT	<1:10000		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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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