

Prisca 5.1.0.17
Date of report: 24-10-2025

N A

Patient data			
Name	Mrs. ROSE MERY		Patient ID 0182510230036
Birthday	04-11-1992		Sample ID A1253477
Age at sample date	33.0		Sample Date 23-10-2025
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	61	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 5
PAPP-A	2.89 mIU/mL	0.65	Method CRL Robinson
fb-hCG	39.62 ng/mL	1.03	Scan date 22-10-2025
Risks at sampling date			Crown rump length in mm 65.9
Age risk	1:414		Nuchal translucency MoM 0.83
Biochemical T21 risk	1:896		Nasal bone present
Combined trisomy 21 risk	1:4924		Sonographer N A
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT MD
Risk			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 4924 women with the same data, there is one woman with a trisomy 21 pregnancy and 4923 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