

Prisca 5.1.0.17
Date of report: 09-01-2026

Patient data			
Name	Mrs. K.MANGAMMA	Patient ID	0842601080030
Birthday	10-09-1994	Sample ID	B4141345
Age at sample date	31.3	Sample Date	08-01-2026
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	77	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.06 mIU/mL	0.38	12 + 3
fb-hCG	43.1 ng/mL	1.09	Method
			CRL Robinson
			Scan date
			08-01-2026
			Crown rump length in mm
			62
			Nuchal translucency MoM
			0.62
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			Sonographer
Risks at sampling date			
Age risk	1:536		
Biochemical T21 risk	1:238		
Combined trisomy 21 risk	1:1551		
Trisomy 13/18 + NT	<1:10000		
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 1551 women with the same data, there is one woman with a trisomy 21 pregnancy and 1550 women with not affected pregnancies. The PAPP-A level is low. 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