

Prisca 5.1.0.17
Date of report: 31/12/25

Patient data			
Name	Mrs. G UMA	Patient ID	0182512310026
Birthday	14/08/07	Sample ID	B3247664
Age at sample date	18.4	Sample Date	31/12/25
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	48	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.38 mIU/mL	0.49	13 + 1
fb-hCG	35.04 ng/mL	0.94	Method
			CRL Robinson
			Scan date
			30/12/25
Risks at sampling date		Trisomy 21	
Age risk	1:1133	Crown rump length in mm	71
Biochemical T21 risk	1:1466	Nuchal translucency MoM	0.79
Combined trisomy 21 risk	1:8700	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	NA
		Qualifications in measuring NT	Sonographer
		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 8700 women with the same data, there is one woman with a trisomy 21 pregnancy and 8699 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off