

Prisca 5.1.0.17
Date of report: 16/05/25

N A

Patient data				
Name	Mrs. C NAGA TEJASWINI		Patient ID	0042505160001
Birthday	24/08/93		Sample ID	23168776
Age at sample date	31.7		Sample Date	15/05/25
Gestational age	12 + 1			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	49	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 1
PAPP-A	1.36 mIU/mL	0.32	Method	CRL Robinson
fb-hCG	44.52 ng/mL	0.91	Scan date	15/05/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:499		58	
Biochemical T21 risk	1:200		Nuchal translucency MoM	
Combined trisomy 21 risk	>1:50		1.84	
Trisomy 13/18 + NT	1:125		Nasal bone	
			present	
			Sonographer	
			N A	
			Qualifications in measuring NT	
			MD	
Trisomy 21				
The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 pregnancy. 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 1:125, which represents a low risk.				

Sign of Physician