

Prisca 5.1.0.17
Date of report: 25-04-2025

NA

Patient data			
Name	W/O NITIN MOHAN	Patient ID	0692504240096
Birthday	10-04-1993	Sample ID	B2766080
Age at sample date	32.0	Sample Date	24-04-2025
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	0.64 mIU/mL	0.19	12 + 4
fb-hCG	40.76 ng/mL	1.03	Method
			CRL Robinson
			Scan date
			24-04-2025
			Crown rump length in mm
			63.46
			Nuchal translucency MoM
			0.64
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			NA
Risks at sampling date			Trisomy 21
Age risk		1:482	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 256 women with the same data, there is one woman with a trisomy 21 pregnancy and 255 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!
Biochemical T21 risk		>1:50	
Combined trisomy 21 risk		1:256	
Trisomy 13/18 + NT		1:913	
Risk 1:10 1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			
Trisomy 13/18 + NT			
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:913, which represents a low risk.			

Sign of Physician

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