

Prisca 5.1.0.17
Date of report: 04-12-2025

Patient data			
Name	Mrs. NAYAKAPU CHANDANA	Patient ID	0352512030035
Birthday	23-09-2002	Sample ID	A1904225
Age at sample date	23.2	Sample Date	03-12-2025
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	81	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.97 mIU/mL	1.34	
fb-hCG	39.4 ng/mL	1.08	
Risks at sampling date			
Age risk	1:1031	Gestational age	12 + 5
Biochemical T21 risk	1:9819	Method	CRL Robinson
Combined trisomy 21 risk	<1:10000	Scan date	03-12-2025
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	65.6
		Nuchal translucency MoM	0.96
		Nasal bone	present
		Sonographer	NA
		Qualifications in measuring NT	Sonographer
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician

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