

Prisca 5.1.0.17
Date of report: 01-10-2025

N A

Patient data			
Name Mrs. DIVYA. W/O. SRAVAN KUMAR		Patient ID 0312509290042	
Birthday 18-04-1994		Sample ID A0281033	
Age at sample date 31.4		Sample Date 29-09-2025	
Gestational age 12 + 4			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 52	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 4
PAPP-A	6.3 mIU/mL	1.32	Method CRL Robinson
fb-hCG	40.75 ng/mL	0.94	Scan date 29-09-2025
Risks at sampling date			Crown rump length in mm 64
Age risk 1:529			Nuchal translucency MoM 1.34
Biochemical T21 risk 1:6711			Nasal bone present
Combined trisomy 21 risk 1:9013			Sonographer N A
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT MD
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 9013 women with the same data, there is one woman with a trisomy 21 pregnancy and 9012 women with not affected pregnancies.</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