

Prisca 5.1.0.17
Date of report: 14-05-2025

N A

Patient data			
Name Mrs. ARCHANA KUMARI BANIK		Patient ID 0482505130306	
Birthday 25-06-1996		Sample ID B2881897	
Age at sample date 28.9		Sample Date 13-05-2025	
Gestational age 12 + 2			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 60	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 1
PAPP-A	1.82 mIU/mL	0.51	Method CRL Robinson
fb-hCG	42.5 ng/mL	0.96	Scan date 12-05-2025
Risks at sampling date			Crown rump length in mm 57.5
Age risk 1:726			Nuchal translucency MoM 1.13
Biochemical T21 risk 1:963			Nasal bone present
Combined trisomy 21 risk 1:2976			Sonographer N A
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT MD
Trisomy 21			
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 2976 women with the same data, there is one woman with a trisomy 21 pregnancy and 2975 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
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