

Prisca 5.1.0.17
Date of report: 08/11/25

N A

Patient data			
Name Mrs. PRAJAKTA DONGAREB3763746		Patient ID 0672511070051	
Birthday 09/09/98		Sample ID B3763746	
Age at sample date 27.2		Sample Date 07/11/25	
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 + 0
PAPP-A	1.71 mIU/mL	0.48	Method CRL Robinson
fb-hCG	40.6 ng/mL	0.92	Scan date 05/11/25
Risks at sampling date			Crown rump length in mm 56.5
Age risk 1:842			Nuchal translucency MoM 0.87
Biochemical T21 risk 1:1045			Nasal bone present
Combined trisomy 21 risk 1:5872			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 5872 women with the same data, there is one woman with a trisomy 21 pregnancy and 5871 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