

Prisca 5.1.0.17
Date of report: 24/05/25

N A

Patient data			
Name Mrs. NAMRATA KALYANSHETTI		Patient ID 0662505230210	
Birthday 06/03/92		Sample ID b2793246	
Age at sample date 33.2		Sample Date 23/05/25	
Gestational age 12 + 1			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 67	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 1
PAPP-A	2.2 mIU/mL	0.74	Method CRL Robinson
fb-hCG	43.88 ng/mL	1.00	Scan date 23/05/25
Risks at sampling date			Crown rump length in mm 58
Age risk 1:385			Nuchal translucency MoM 0.99
Biochemical T21 risk 1:1239			Nasal bone present
Combined trisomy 21 risk 1:5244			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 5244 women with the same data, there is one woman with a trisomy 21 pregnancy and 5243 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