

Prisca 5.1.0.17
Date of report: 18-03-2025

N A

Patient data			
Name Mrs. KEHKASHA ZAID ANSARI		Patient ID 0662503170098	
Birthday 17-12-1994		Sample ID B2564954	
Age at sample date 30.2		Sample Date 17-03-2025	
Gestational age 12 + 3			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 83	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 1
PAPP-A	1.99 mIU/mL	0.78	Method CRL Robinson
fb-hCG	74.23 ng/mL	1.91	Scan date 15-03-2025
Risks at sampling date			Crown rump length in mm 58
Age risk 1:624			Nuchal translucency MoM 0.86
Biochemical T21 risk 1:477			Nasal bone present
Combined trisomy 21 risk 1:2632			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 2632 women with the same data, there is one woman with a trisomy 21 pregnancy and 2631 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