

Prisca 5.1.0.17
Date of report: 05/05/25

N A

Patient data			
Name Mrs. MAHITHA W/O PREM DAS		Patient ID 0012505010315	
Birthday 01/05/98		Sample ID B2750268	
Age at sample date 27.0		Sample Date 01/05/25	
Gestational age 12 + 4			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 48	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 4
PAPP-A	0.44 mIU/mL	0.08	Method CRL Robinson
fb-hCG	41.3 ng/mL	0.92	Scan date 01/05/25
Risks at sampling date			Crown rump length in mm 64
Age risk 1:861			Nuchal translucency MoM 0.61
Biochemical T21 risk 1:76			Nasal bone present
Combined trisomy 21 risk 1:574			Sonographer N A
Trisomy 13/18 + NT >1:50			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 574 women with the same data, there is one woman with a trisomy 21 pregnancy and 573 women with not affected pregnancies. The PAPP-A level is low. 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 test (with nuchal translucency) is >1:50, which indicates an increased risk.			

Sign of Physician