

Prisca 5.1.0.17
Date of report: 10/04/25

DR
S MANJULA MBBS

Patient data			
Name Mrs. SHUBHANGI MOUNDEKAR		Patient ID 0372504080118	
Birthday 27/06/00		Sample ID A1558661	
Age at sample date 24.8		Sample Date 08/04/25	
Gestational age 11 + 1			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 52	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 1
PAPP-A	1.66 mIU/mL	0.67	Method CRL Robinson
fb-hCG	55.41 ng/mL	0.94	Scan date 08/04/25
Risks at sampling date			Crown rump length in mm 45
Age risk 1:921			Nuchal translucency MoM 0.81
Biochemical T21 risk 1:2573			Nasal bone present
Combined trisomy 21 risk <1:10000			Sonographer DR S MANJULA MBBS
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT DMCH
Risk			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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