

Prisca 5.1.0.17
Date of report: 27-09-2024

N A

Patient data			
Name Mrs. SHRADDHA PRITESH MANGTANI		Patient ID 0662409260190	
Birthday 25-01-1999		Sample ID a0506859	
Age at sample date 25.7		Sample Date 26-09-2024	
Gestational age 13 + 3			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 70	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 3
PAPP-A	6.39 mIU/mL	1.38	Method CRL Robinson
fb-hCG	42.1 ng/mL	1.34	Scan date 26-09-2024
Risks at sampling date			Crown rump length in mm 76
Age risk 1:960			Nuchal translucency MoM 0.81
Biochemical T21 risk 1:5871			Nasal bone present
Combined trisomy 21 risk <1:10000			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 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