

Prisca 5.1.0.17
Date of report: 25-09-2024

N A

Patient data			
Name Mrs. POOJA GAJANAN PAWADE		Patient ID 0372409230169	
Birthday 19-05-1995		Sample ID A0423644	
Age at sample date 29.3		Sample Date 23-09-2024	
Gestational age 13 + 2			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 51	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 0
PAPP-A	4.3 mIU/mL	0.67	Method CRL Robinson
fb-hCG	38.53 ng/mL	1.06	Scan date 21-09-2024
Risks at sampling date			Crown rump length in mm 70
Age risk 1:715			Nuchal translucency MoM 1.02
Biochemical T21 risk 1:1579			Nasal bone present
Combined trisomy 21 risk 1:6253			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 6253 women with the same data, there is one woman with a trisomy 21 pregnancy and 6252 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