

Prisca 5.1.0.17
Date of report: 17-12-2024

N A

Patient data				
Name	Mrs. VIMAL MANOJ SHELKE		Patient ID	
Birthday	10-08-2004		Sample ID	
Age at sample date	20.3		Sample Date	
Gestational age	12 + 6			
Correction factors				
Fetuses	1	IVF	no	
Weight	52	diabetes	no	
Smoker	no	Origin	Asian	
		Previous trisomy 21 pregnancies	unknown	
Biochemical data		Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	
PAPP-A	12.52 mIU/mL	2.35	Method	
fb-hCG	37.07 ng/mL	0.92	Scan date	
Risks at sampling date			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!	
Age risk	1:1096			
Biochemical T21 risk	<1:10000			
Combined trisomy 21 risk	<1:10000			
Trisomy 13/18 + NT	<1:10000			
			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