

Prisca 5.1.0.17
Date of report: 21-12-2024

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Patient data			
Name	Mrs. AKHILA		Patient ID
Birthday	26-01-2000	Sample ID	
Age at sample date	24.9	Sample Date	
Gestational age	11 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	48	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.84 mIU/mL	0.63	Method
fb-hCG	90.35 ng/mL	1.53	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 4310 women with the same data, there is one woman with a trisomy 21 pregnancy and 4309 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!
Age risk	1:921		
Biochemical T21 risk	1:722		
Combined trisomy 21 risk	1:4310		
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