

Prisca 5.1.0.17
Date of report: 13-09-2024

N A

Patient data			
Name		Mrs. PRIYANKA SAROJ	
Patient ID		0662409120191	
Birthday		03-04-1998	
Sample ID		A0997664	
Age at sample date		26.4	
Sample Date		12-09-2024	
Gestational age		13 + 1	
Correction factors			
Fetuses	1	IVF	no
Weight	44	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.86 mIU/mL	0.54	13 + 0
fb-hCG	54.39 ng/mL	1.36	Method
Risks at sampling date			CRL Robinson
Age risk			11-09-2024
Biochemical T21 risk			1:910
Combined trisomy 21 risk			1:643
Trisomy 13/18 + NT			1:3915
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			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 3915 women with the same data, there is one woman with a trisomy 21 pregnancy and 3914 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!
1:100			
1:250			
Cut off			
1:1000			
1:10000			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
Age			
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

below cut off
 Below Cut Off, but above Age Risk
 above cut off