

Prisca 5.1.0.17
Date of report: 20-03-2024

Patient data				
Name	Mrs. ROSHNI SHELAR	Patient ID	0662403190118	
Birthday	31-05-1996	Sample ID	A0107884	
Age at sample date	27.8	Sample Date	19-03-2024	
Gestational age	12 + 1			
Correction factors				
Fetuses	1	IVF	no	
Weight	74	diabetes	no	
Smoker	no	Origin	Asian	
		Previous trisomy 21 pregnancies	unknown	
Biochemical data		Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 1
PAPP-A	6.09 mIU/mL	2.32	Method	CRL Robinson
fb-hCG	56.01 ng/mL	1.31	Scan date	19-03-2024
Risks at sampling date			Crown rump length in mm	57.4
Age risk	1:798		Nuchal translucency MoM	0.46
Biochemical T21 risk	1:9780		Nasal bone	present
Combined trisomy 21 risk	<1:10000		Sonographer	NA
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MD
RISK 1:10 1:100 1:250 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 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