

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

01-11-2023

Patient data			
Name		Mrs. SAMAN MALIK	
Patient ID		0382310310140	
Birthday		10-10-1998	
Sample ID		24996713	
Age at sample date		25.1	
Sample Date		31-10-2023	
Gestational age		12 + 0	
Correction factors			
Fetuses	1	IVF	no
Weight	81.5	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.91 mIU/mL	1.33	Gestational age
fb-hCG	45.65 ng/mL	1.06	Method
Risks at sampling date			CRL Robinson
Age risk			30-10-2023
Biochemical T21 risk			Crown rump length in mm
Combined trisomy 21 risk			54.66
Trisomy 13/18 + NT			Nuchal translucency MoM
			0.97
			Nasal bone
			unknown
			Sonographer
			NA
			Qualifications in measuring NT
			MD
Risk		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

☐ below cut off

 Below Cut Off, but above Age Risk

above cut off