

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

26/11/21

Patient data			
Name		Mrs. ANUSHA	
Patient ID		0252111250002	
Birthday		21/09/99	
Sample ID		22682324	
Age at sample date		22.2	
Sample Date		25/11/21	
Gestational age		12 + 6	
Correction factors			
Fetuses	1	IVF	no
Weight	40	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.02 mIU/mL	0.56	
fb-hCG	38.96 ng/mL	0.87	
Risks at sampling date			
Age risk		1:1063	
Biochemical T21 risk		1:2283	
Combined trisomy 21 risk		<1:10000	
Trisomy 13/18 + NT		<1:10000	
Gestational age			12 + 6
Method			CRL Robinson
Scan date			25/11/21
Crown rump length in mm			68.3
Nuchal translucency MoM			0.87
Nasal bone			present
Sonographer			Dr 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