

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

25-01-2024

Patient data			
Name		Mrs. DIVYA	
Patient ID		0012401250110	
Birthday		11-08-1999	
Sample ID		A0035149	
Age at sample date		24.5	
Sample Date		25-01-2024	
Gestational age		12 + 2	
Correction factors			
Fetuses	1	IVF	no
Weight	56	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	0.75	
fb-hCG	42.5 ng/mL	0.94	
Risks at sampling date			
Age risk		1:975	
Biochemical T21 risk		1:3615	
Combined trisomy 21 risk		1:917	
Trisomy 13/18 + NT		<1:10000	
Gestational age			11 + 4
Method			CRL Robinson
Scan date			20-01-2024
Crown rump length in mm			51
Nuchal translucency MoM			1.75
Nasal bone			present
Sonographer			NA
Qualifications in measuring NT			NA
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 917 women with the same data, there is one woman with a trisomy 21 pregnancy and 916 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!
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