

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

18-01-2024

Patient data			
Name		Mrs. PRIYA SINGH	
Patient ID		0382401160124	
Birthday		18-08-1994	
Sample ID		A0184050	
Age at sample date		29.4	
Sample Date		16-01-2024	
Gestational age		13 + 3	
Correction factors			
Fetuses	1	IVF	no
Weight	45.6	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	9.56 mIU/mL	1.25	
fb-hCG	38.99 ng/mL	1.07	
Gestational age		11 + 6	
Method		CRL Robinson	
Scan date		05-01-2024	
Risks at sampling date		Crown rump length in mm	
Age risk		1:714	
Biochemical T21 risk		1:6071	
Combined trisomy 21 risk		<1:10000	
Trisomy 13/18 + NT		<1:10000	
Nuchal translucency MoM		0.70	
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 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

