

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

07-03-2024

Patient data			
Name		Mrs. MANISHA	
Patient ID		0012403060563	
Birthday		10-07-2003	
Sample ID		A0229816	
Age at sample date		20.7	
Sample Date		06-03-2024	
Gestational age		11 + 6	
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	
PAPP-A	4.13 mIU/mL	1.79	
fb-hCG	49.11 ng/mL	1.08	
Gestational age		11 + 6	
Method		CRL Robinson	
Scan date		06-03-2024	
Crown rump length in mm		54.7	
Nuchal translucency MoM		0.28	
Nasal bone		unknown	
Sonographer		NA	
Qualifications in measuring NT		MD	
Risks at sampling date		Trisomy 21	
Age risk	1:1054	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!	
Biochemical T21 risk	<1:10000		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
Risk Age			
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