

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

02-04-2024

Patient data						
Name		Mrs. ARSHIYA		Patient ID	0012404010327	
Birthday		19-08-1999		Sample ID	22622801	
Age at sample date		24.6		Sample Date	01-04-2024	
Gestational age		13 + 4				
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown	
Weight	47	diabetes	no			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age			13 + 3
PAPP-A	23.75 mIU/mL	3.06	Method			CRL Robinson
fb-hCG	30.51 ng/mL	0.88	Scan date			31-03-2024
Risks at sampling date			Crown rump length in mm			76
Age risk		1:1011	Nuchal translucency MoM			0.81
Biochemical T21 risk		<1:10000	Nasal bone			present
Combined trisomy 21 risk		<1:10000	Sonographer			N A
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT			MD
Risk 1:10 1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			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 PAPP-A level is high. 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

