

SAGEPATH LABS PVT LTD.

Prisca

5.1.0.17

Date of report:

24-06-2024

Patient data					
Name		Mrs. VINITA GUPTA		Patient ID	0212406190028
Birthday		30-03-1998		Sample ID	24252914
Age at sample date		26.2		Sample Date	19-06-2024
Gestational age		12 + 4			
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	79	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age 12 + 4		
PAPP-A	3.28 mIU/mL	1.13	Method CRL Robinson		
fb-hCG	39.76 ng/mL	1.04	Scan date 19-06-2024		
Risks at sampling date			Crown rump length in mm 64		
Age risk		1:905	Nuchal translucency MoM 0.61		
Biochemical T21 risk		1:6670	Nasal bone unknown		
Combined trisomy 21 risk		<1:10000	Sonographer N A		
Trisomy 13/18 + NT		<1:10000	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