

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

17/01/24

Patient data			
Name		Mrs. KIRAN GHULE	Patient ID
Birthday		11/11/95	0662401150190
Age at sample date		28.2	Sample ID
Gestational age		12 + 4	A0121367
			Sample Date
			15/01/24
Correction factors			
Fetuses	1	IVF	no
Weight	64.9	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.55 mIU/mL	2.05	11 + 6
fb-hCG	41.21 ng/mL	1.02	Method
Risks at sampling date			CRL Robinson
Age risk	1:784		Scan date
Biochemical T21 risk	<1:10000		10/01/24
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		53.8
			Nuchal translucency MoM
			0.77
			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

 below cut off
 Below Cut Off, but above Age Risk
 above cut off