

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

02-03-2024

Patient data			
Name	Mrs. SHRUTI TELURE		Patient ID
Birthday	24-01-1994	Sample ID	0662402270104
Age at sample date	30.1	Sample Date	A0100707
Gestational age	12 + 4		27-02-2024
Correction factors			
Fetuses	1	IVF	no
Weight	68.9	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	9.98 mIU/mL	2.92	12 + 3
fb-hCG	49.1 ng/mL	1.24	Method
			CRL Robinson
			Scan date
			26-02-2024
Risks at sampling date			Crown rump length in mm
Age risk	1:640		61.6
Biochemical T21 risk	1:8916		Nuchal translucency MoM
Combined trisomy 21 risk	<1:10000		0.88
Trisomy 13/18 + NT	<1:10000		Nasal bone
			present
			Sonographer
			NA
			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 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