

SAGEPATH LABS PVT LTD.

Prisca

5.1.0.17

Date of report:

26-05-2024

Patient data			
Name	Mrs. PRATIKSHA SHINGARE		Patient ID
Birthday	28-03-2001	Sample ID	0662405250104
Age at sample date	23.2	Sample Date	A0721670
Gestational age	12 + 0		25-05-2024
Correction factors			
Fetuses	1	IVF	no
Weight	75	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.66 mIU/mL	1.51	11 + 6
fb-hCG	24.73 ng/mL	0.56	Method
			CRL Robinson
			Scan date
			24-05-2024
Risks at sampling date		Crown rump length in mm	
Age risk	1:1007	55	
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000	0.69	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		N A	
		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