SAGEPATH LABS PVT LTD.

Prisca   5.1.0.17

Date of report: **14-01-2024**

Patient data			
Name	Mrs.RAJASHRI AJAY THAKUR	Patient ID	0662401130271
Birthday	06-10-2000	Sample ID	24973332
Age at sample date	23.3	Sample Date	13-01-2024
Gestational age	11 + 3		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	48.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 3
PAPP-A	4.44 mIU/mL	1.43	Method	CRL Robinson
fb-hCG	50.93 ng/mL	0.89	Scan date	13-01-2024

Risks at sampling date		Ultrasound data (cont.)	
		Crown rump length in mm	48.5
Age risk	1:982	Nuchal translucency MoM	0.76
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

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