SAGEPATH LABS PVT LTD.

Prisca   5.1.0.17

Date of report: **05/08/24**

Patient data			
Name	Mrs. CH.BHAVANI	Patient ID	0352408020063
Birthday	06/06/87	Sample ID	23422718
Age at sample date	37.2	Sample Date	02/08/24
Gestational age	12 + 3		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	61	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data				Ultrasound data	
Parameter	Value	Corr. MoM		Gestational age	12 + 3
PAPP-A	6.42 mIU/mL	1.72		Method	CRL Robinson
fb-hCG	41.12 ng/mL	0.97		Scan date	02/08/24
Risks at sampling date				Crown rump length in mm	61
Age risk		1:164		Nuchal translucency MoM	0.76
Biochemical T21 risk		1:3152		Nasal bone	unknown
Combined trisomy 21 risk		<1:10000		Sonographer	N A
Trisomy 13/18 + NT		<1:10000		Qualifications in measuring NT	MD

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