

SAGEPATH LABS PVT LTD.

Prisca

5.1.0.17

Date of report:

06-05-2024

Patient data			
Name	Mrs. NADAPRIYA		Patient ID
0012405040409			
Birthday	04-07-1998		Sample ID
A0373510			
Age at sample date	25.8		Sample Date
04-05-2024			
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	74	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	10.67 mIU/mL	2.35	13 + 4
fb-hCG	54.1 ng/mL	1.82	Method
Risks at sampling date			CRL Robinson
Age risk	1:956		Scan date
Biochemical T21 risk	1:5223		04-05-2024
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		79
Risk			Nuchal translucency MoM
1:10			0.78
1:100			Nasal bone
1:250			present
1:1000			Sonographer
1:10000			N A
Age			Qualifications in measuring NT
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			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