

SAGEPATH LABS PVT LTD.

Prisca

5.1.0.17

Date of report:

03/07/24

Patient data			
Name		Mrs. JAMESHWARI	Patient ID
0352407010059			
Birthday		10/06/00	Sample ID
A0577245			
Age at sample date		24.1	Sample Date
01/07/24			
Gestational age		12 + 5	
Correction factors			
Fetuses	1	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.47 mIU/mL	0.72	12 + 5
fb-hCG	41.8 ng/mL	1.01	Method
Risks at sampling date			CRL Robinson
Age risk	1:1004		Scan date
Biochemical T21 risk	1:2883		01/07/24
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		66
			Nuchal translucency MoM
			1.01
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risk		Trisomy 21	
1:10		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
1:100		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.	
1:250		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
1:1000		Please note that risk calculations are statistical approaches and have no diagnostic value!	
1:10000		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Age			
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