

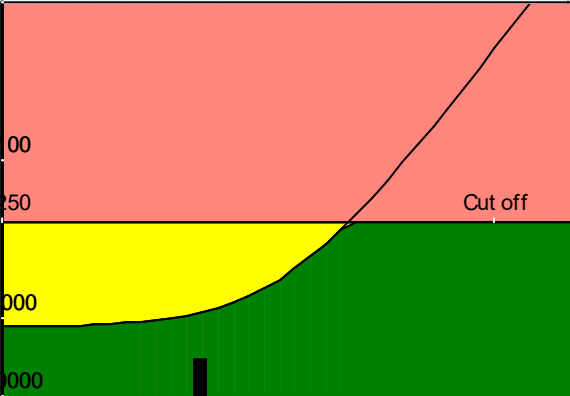
SAGEPATH LABS PVT LTD.

Prisca

5.1.0.17

Date of report:

06-03-2024

Patient data						
Name		Mrs. PURNIMA PRADHAN		Patient ID	0662403040186	
Birthday		04-05-1998		Sample ID	A0129806	
Age at sample date		25.8		Sample Date	04-03-2024	
Gestational age		12 + 6				
Correction factors						
Fetuses	1	IVF	unknown	Previous trisomy 21 pregnancies	unknown	
Weight	52.5	diabetes	unknown			
Smoker	unknown	Origin	Caucasian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age			12 + 4
PAPP-A	7.51 mIU/mL	1.66	Method			CRL Robinson
fb-hCG	36.03 ng/mL	0.93	Scan date			02-03-2024
Risks at sampling date			Crown rump length in mm			64.2
Age risk		1:934	Nuchal translucency MoM			0.49
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 1:10 1:100 1:250 1:1000 1:10000  Age			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