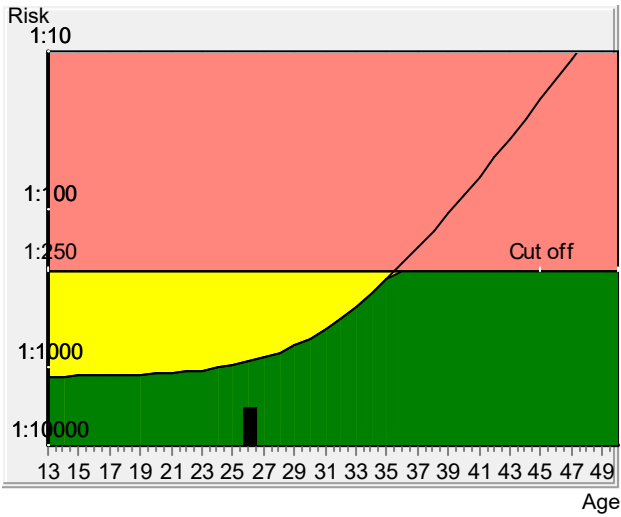


SAGEPATH LABS PVT

Prisca 5.1.0.17
Date of report: 10-05-2024

Patient data		
Name	Mrs. HANIKA	Patient ID 0382405060071
Birthday	06-04-1998	Sample ID A0687892
Age at sample date	26.1	Sample Date 06-05-2024
Gestational age	13 + 2	
Correction factors		
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown
Weight 89	diabetes no	
Smoker no	Origin Asian	
Biochemical data		Ultrasound data
Parameter	Value	Corr. MoM
PAPP-A	7.38 mIU/mL	2.27
fb-hCG	45.98 ng/mL	1.49
Risks at sampling date		
Age risk	1:934	
Biochemical T21 risk	1:8297	
Combined trisomy 21 risk	<1:10000	
Trisomy 13/18 + NT	<1:10000	
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