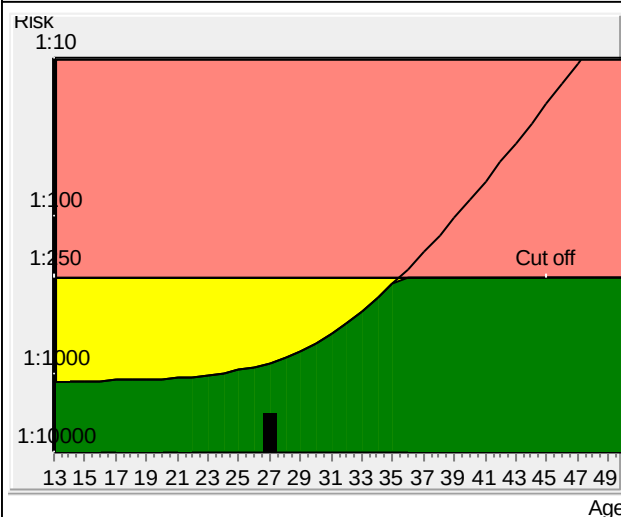


Prisca 5.1.0.17
Date of report: 24-03-2024

Patient data		
Name	Mrs. MANISHA NAGARE	Patient ID 0372403230157
Birthday	04-04-1997	Sample ID 25142224
Age at sample date	27.0	Sample Date 23-03-2024
Gestational age	12 + 4	
Correction factors		
Fetuses	1	IVF no
Weight	58	diabetes no
Smoker	no	Origin Asian
		Previous trisomy 21 pregnancies unknown
Biochemical data		Ultrasound data
Parameter	Value	Corr. MoM
PAPP-A	7.74 mIU/mL	1.84
fb-hCG	48.66 ng/mL	1.16
Risks at sampling date		
Age risk	1:863	
Biochemical T21 risk	<1:10000	
Combined trisomy 21 risk	<1:10000	
Trisomy 13/18 + NT	<1:10000	
		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