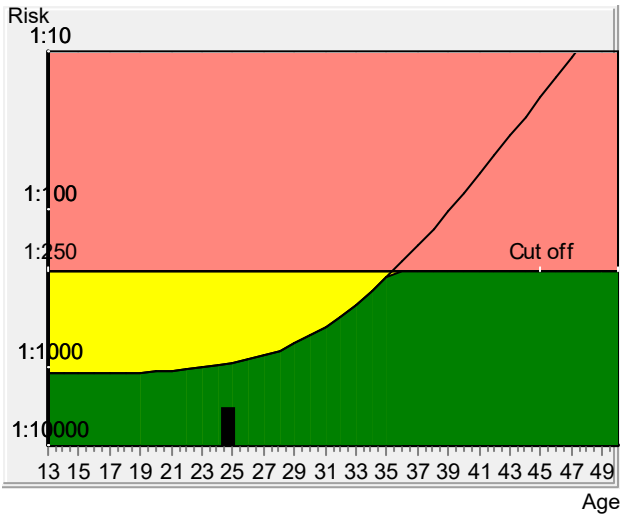


Patient data		
Name	Mrs. SUPRATIKSHA PARAB	Patient ID 0662402010180
Birthday	19-05-1999	Sample ID a0128392
Age at sample date	24.7	Sample Date 01-02-2024
Gestational age	12 + 2	
Correction factors		
Fetuses	1	IVF no
Weight	71	diabetes no
Smoker	no	Origin Asian
Previous trisomy 21 pregnancies		unknown
Biochemical data		Ultrasound data
Parameter	Value	Corr. MoM
PAPP-A	2.91 mIU/mL	0.99
fb-hCG	42.5 ng/mL	1.01
Risks at sampling date		Gestational age 12 + 2
Age risk	1:965	Method CRL Robinson
Biochemical T21 risk	1:5742	Scan date 01-02-2024
Combined trisomy 21 risk	<1:10000	Crown rump length in mm 59
Trisomy 13/18 + NT	<1:10000	Nuchal translucency MoM 1.04
		Nasal bone unknown
		Sonographer N A
		Qualifications in measuring NT MD
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