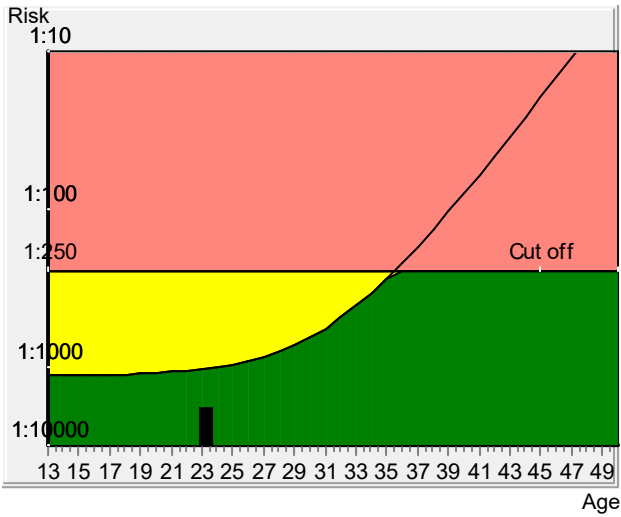


Patient data				
Name	Mrs.SWAPNALI SATISH SHELAR	Patient ID	0662403010268	
Birthday	25-11-2000	Sample ID	A0129716	
Age at sample date	23.3	Sample Date	01-03-2024	
Gestational age	12 + 6			
Correction factors				
Fetuses	1	IVF	no	
Weight	42	diabetes	no	
Smoker	no	Origin	Asian	
		Previous trisomy 21 pregnancies	unknown	
Biochemical data		Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	23.78 mIU/mL	3.50	Method	CRL Robinson
fb-hCG	44.42 ng/mL	1.01	Scan date	29-02-2024
Risks at sampling date			Crown rump length in mm	66.2
Age risk	1:1034		Nuchal translucency MoM	0.77
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			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 PAPP-A level is high. 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