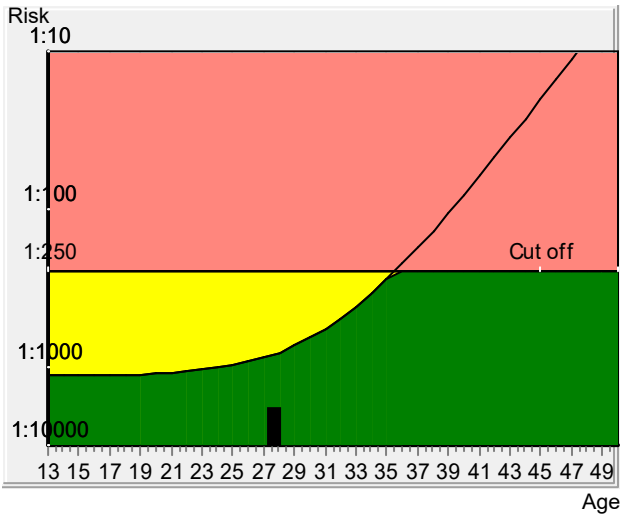


Prisca 5.1.0.17
Date of report: 02-08-2025

N A

Patient data			
Name		Mrs. VAISHNAVI GAURAV UKINKAR	
Patient ID		0662508010018	
Birthday		17-11-1997	
Sample ID		B3296997	
Age at sample date		27.7	
Sample Date		01-08-2025	
Gestational age		13 + 1	
Correction factors			
Fetuses	1	IVF	no
Weight	48	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.77 mIU/mL	0.58	13 + 1
fb-hCG	45.28 ng/mL	1.17	Method
			CRL Robinson
Risks at sampling date			Scan date
			01-08-2025
Age risk			Crown rump length in mm
1:833			72
Biochemical T21 risk			Nuchal translucency MoM
1:1009			0.67
Combined trisomy 21 risk			Nasal bone
1:5918			present
Trisomy 13/18 + NT			Sonographer
<1:10000			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 5918 women with the same data, there is one woman with a trisomy 21 pregnancy and 5917 women with not affected pregnancies. 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