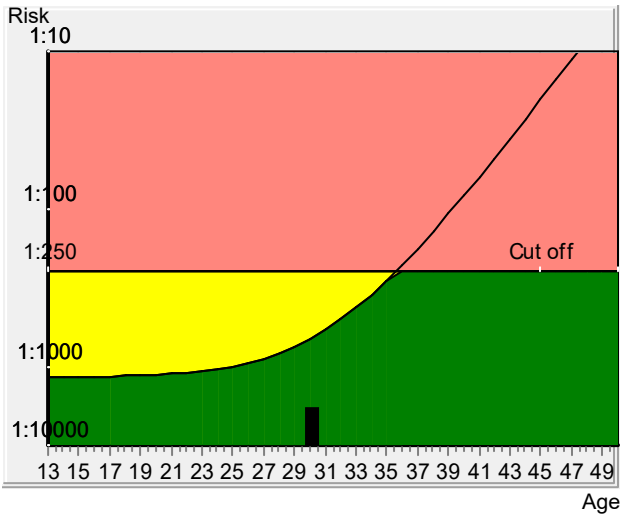


Prisca 5.1.0.17
Date of report: 05-04-2025

N A

Patient data						
Name		Mrs. SAVITA MEHRA	Patient ID	0822503300003.		
Birthday		11-02-1995	Sample ID	A2034341		
Age at sample date		30.1	Sample Date	30-03-2025		
Gestational age		13 + 5				
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown	
Weight	45	diabetes	no			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age			13 + 4
PAPP-A	5 mIU/mL	0.58	Method			CRL Robinson
fb-hCG	29.22 ng/mL	0.87	Scan date			29-03-2025
Risks at sampling date			Crown rump length in mm			78
Age risk		1:662	Nuchal translucency MoM			0.90
Biochemical T21 risk		1:1599	Nasal bone			present
Combined trisomy 21 risk		1:8285	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 8285 women with the same data, there is one woman with a trisomy 21 pregnancy and 8284 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