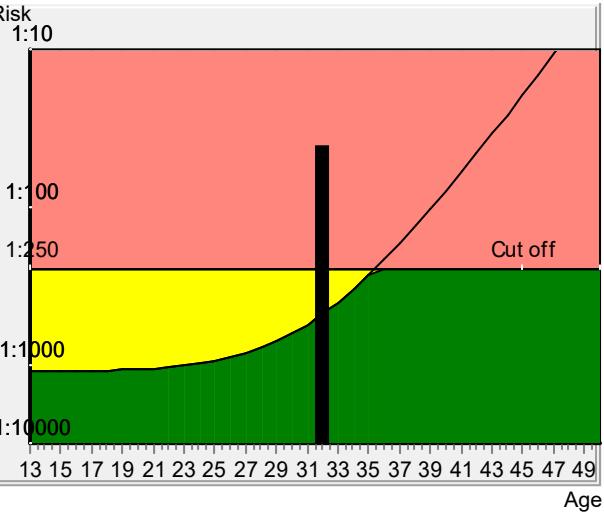


N A

Patient data			
Name	Mrs. KARISHMA BABBAR	Patient ID	0622507260070
Birthday	31-08-1993	Sample ID	A1820175
Age at sample date	31.9	Sample Date	26-07-2025
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	2.63 mIU/mL	0.78	Gestational age 12 + 0
fb-hCG	42.84 ng/mL	0.94	Method CRL Robinson
Risks at sampling date			
Age risk	1:485		Scan date 25-07-2025
Biochemical T21 risk	1:1984		Crown rump length in mm 55.7
Combined trisomy 21 risk	>1:50		Nuchal translucency MoM 2.31
Trisomy 13/18 + NT	1:427		Nasal bone present
Sonographer N A			
Qualifications in measuring NT MD			
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.			
After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one 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 held 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 1:427, which represents a low risk.			

Sign of Physician