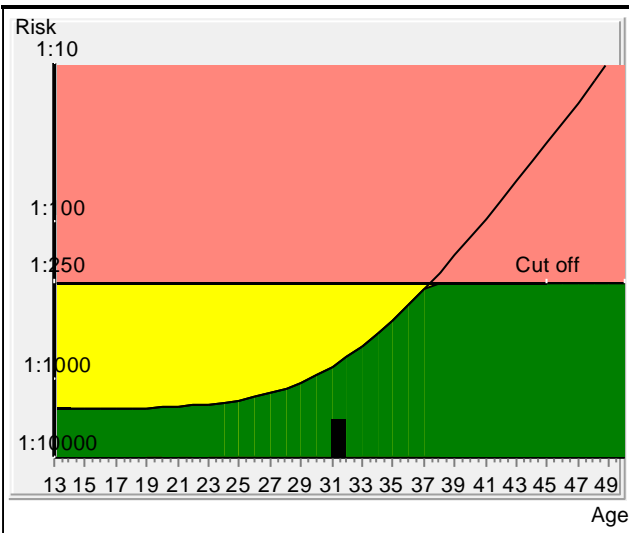


Patient data			
Name	MRS. NEHA BAJAJ	Patient ID	4053660
Birthday	01-02-1992	Sample ID	4053660
Age at delivery	31.4	Sample Date	29-12-2022
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	unknown
Weight	58.7	diabetes	unknown
		Previous trisomy 21 pregnancies	unknown

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 2
PAPP-A	3.9 mIU/ml	1.08	Method	CRL Robinson
fb-hCG	25 ng/ml	0.71	Scan date	27-12-2022
Risks at term			Crown rump length in mm	60
Age risk		1:804	Nuchal translucency MoM	0.77
Biochemical T21 risk		<1:10000	Nasal bone	present
Combined trisomy 21 risk		<1:10000	Sonographer	Dr. --
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	FMF



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

