Typolog Software Evaluation version
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Not allowed to use on real patients

Prisca 5.1.0.17

Date of report: 26-08-2025

NA

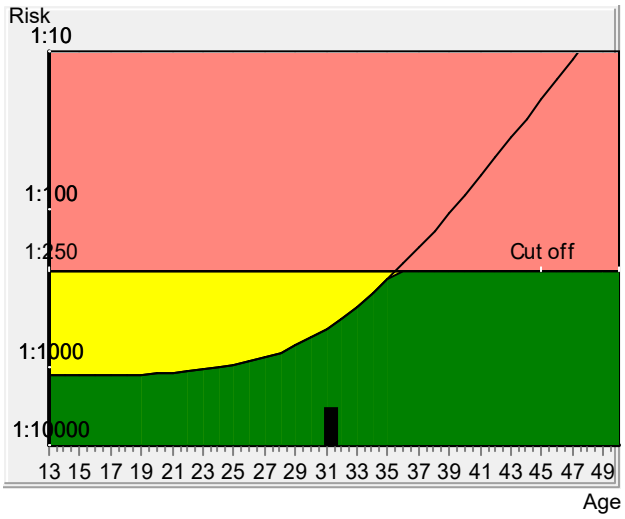
Patient data			
Name	Mrs. KAJAL SRIVASTAVA	Patient ID	0482508250129
Birthday	24-04-1994	Sample ID	B3671422
Age at sample date	31.3	Sample Date	25-08-2025
Gestational age	13 + 1		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	76	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data		
Parameter	Value	Corr. MoM
PAPP-A	2.76 mIU/mL	0.63
fb-hCG	33.02 ng/ml	1.03

Risks at sampling date	
Age risk	1:550
Biochemical T21 risk	1:1109
Combined trisomy 21 risk	1:6213
Trisomy 13/18 + NT	<1:10000

Ultrasound data	
Gestational age	12 + 6
Method	CRL Robinson
Scan date	23-08-2025
Crown rump length in mm	68.4
Nuchal translucency MoM	0.81
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	MD



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 6213 women with the same data, there is one woman with a trisomy 21 pregnancy and 6212 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