Prisca 5.1.0.17

Date of report: **31-08-2025**

NA

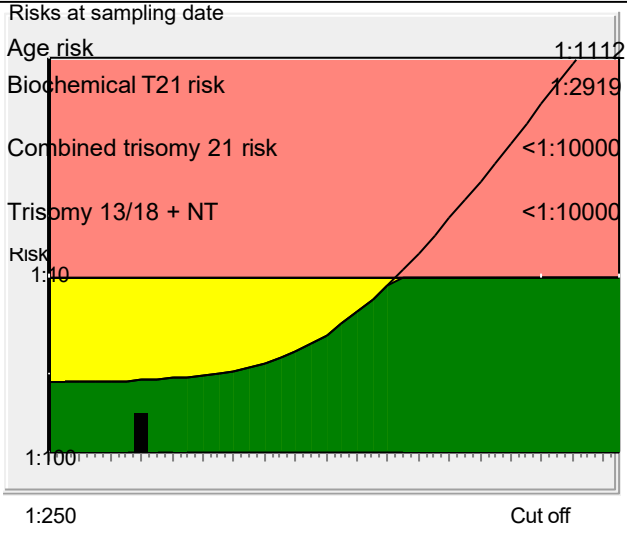
Patient data			
Name	Mrs. A RITHWIK	Patient ID	0182508300040
Birthday	30-08-2006	Sample ID	A1894264
Age at sample date	19.0	Sample Date	30-08-2025
Gestational age	12 + 6		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	55	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data		
Parameter	Value	Corr. MoM
PAPP-A	3.81 mIU/mL	0.67
fb-hCG	37.01 ng/ml	0.98

Ultrasound data	
Gestational age	12 + 5
Method	CRL Robinson
Scan date	29-08-2025
Crown rump length in mm	65
Nuchal translucency MoM	0.72
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	MD

Risks at sampling date	
Age risk	1:1112
Biochemical T21 risk	1:2919
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000



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