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Patient data

Name	SAMTA	Patient ID	2M036599
Birthday	30/08/94	Sample ID	2M036599
Age at sample date	31.1	Sample Date	05/10/25
Gestational age	13 + 4		

Correction factors

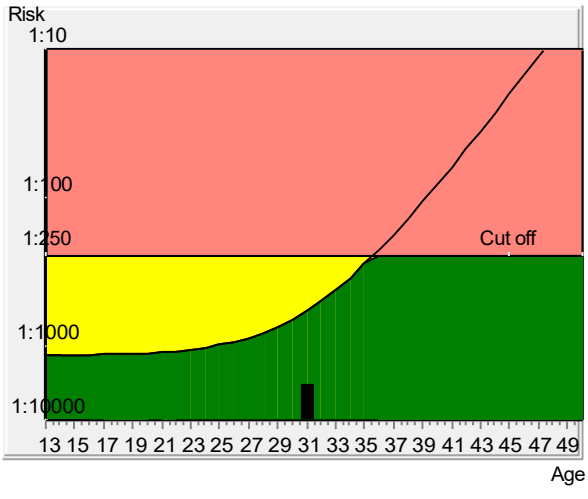
Fetuses	1	IVF	unknown	Previous trisomy 21 pregnancies	unknown
Weight	52	diabetes	unknown		
Smoker	unknown	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	6 mIU/mL	0.75
fb-hCG	49 ng/mL	1.77

Risks at sampling date

Age risk	1:578
Biochemical T21 risk	1:494
Combined trisomy 21 risk	1:2850
Trisomy 13/18 + NT	<1:10000



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Ultrasound data

Gestational age	12 + 1
Method	CRL Robinson
Scan date	25/09/25
Crown rump length in mm	57
Nuchal translucency MoM	0.80
Nasal bone	unknown
Sonographer	DR SELF
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 2850 women with the same data, there is one woman with a trisomy 21 pregnancy and 2849 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!

Sign of Physician

below cut off | Below Cut Off, but above Age Risk | above cut off