Prisca   5.0.2.37

Date of report: 17-07-2018

Patient data

Name	MRS. SIMMI	Patient ID	
Birthday	24-12-1994	Sample ID	1807220398/AMB
Age at sample date	23.6	Sample Date	14-07-2018

Gestational age	11 + 5		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21	no

Weight	51	diabetes	no	pregancies	
Smoker	no	Origin		Asian	

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 3
PAPP-A	6.7 mIU/ml	2.24	Method	CRL Robinson
fb-hCG	48.1 ng/ml	1.12	Scan date	12-07-2018

Risks at sampling date

		Crown rump length in mm	49.2
Age risk	1:985	Nuchal translucency MoM	1.43
Biochemical T21 risk	<1:10000	Nasal bone	present
Combined trisomy 21 risk	<1:10000	Sonographer	.
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MD

Risk
1:10

1:100

1:250   Cut off

1:1000

1:1000

13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49
Age

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!

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

Trisomy 13/18 + NT

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

risk.

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Sign of Physician