Prisca 5.1.0.17

Date of report: 09/08/20

Patient data

Name	MRS.SRAVANTHI	Patient ID	20022259
Birthday	21/12/87	Sample ID	20022259
Age at delivery	33.2	Sample Date	05/08/20
Gestational age	12 + 1		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	52.2	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

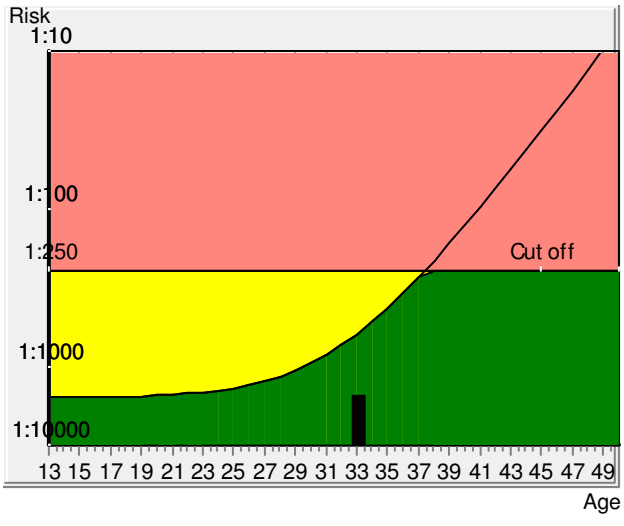
Parameter	Value	Corr. MoM
PAPP-A	1.72 mIU/ml	0.44
fb-hCG	41 ng/ml	0.94

Risks at term

Age risk	1:610
Biochemical T21 risk	1:582
Combined trisomy 21 risk	1:1516
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 1
Method	CRL Robinson
Scan date	05/08/20
Crown rump length in mm	58
Nuchal translucency MoM	1.18
Nasal bone	unknown
Sonographer	NA
Qualifications in measuring NT	NA



Trisomy 13/18 + NT

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

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 1516 women with the same data, there is one woman with a trisomy 21 pregnancy and 1515 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