Sagepath Labs Pvt Ltd

Prisca 5.0.2.37

Date of report: 23/07/20

Patient data

Name	MRS. MANI	Patient ID	20008439
Birthday	03/05/90	Sample ID	20008439
Age at delivery	30.8	Sample Date	20/07/2020
Gestational age	11 + 5		

Correction factors

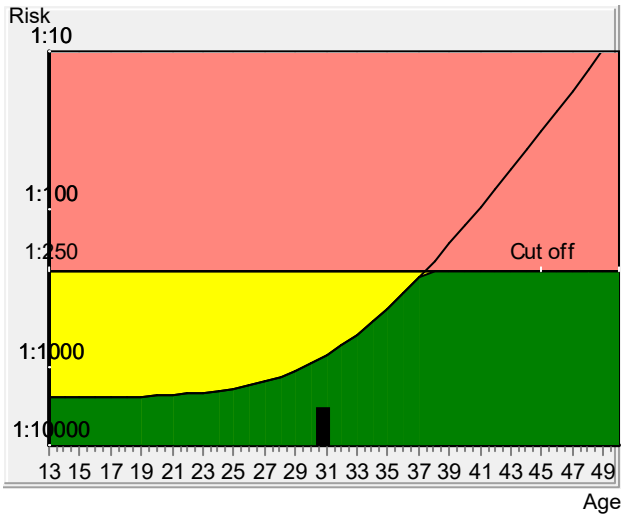
Fetuses	1	IVF	unknown	Previous trisomy 21 pregancies	unknown
Weight	100	diabetes	unknown		
Smoker	unknown	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	0.722 mIU/ml	0.55
fb-hCG	52.1 ng/ml	1.33

Risks at term

Age risk	1:880
Biochemical T21 risk	1:711
Combined trisomy 21 risk	1:2826
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	11 + 4
Method	CRL Robinson
Scan date	20/07/20
Crown rump length in mm	50
Nuchal translucency MoM	1.04
Nasal bone	present
Sonographer	DR. PHANINDRANATH
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 2826 women with the same data, there is one woman with a trisomy 21 pregnancy and 2825 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