SagePath Labs Pvt Ltd.

Prisca 5.1.0.17

Date of report: 22/09/20

Patient data

Name	Mrs. R.SIRISHA	Patient ID	0212009140004
Birthday	03/01/92	Sample ID	20022417
Age at sample date	28.7	Sample Date	14/09/20
Gestational age	13 + 3		

Correction factors

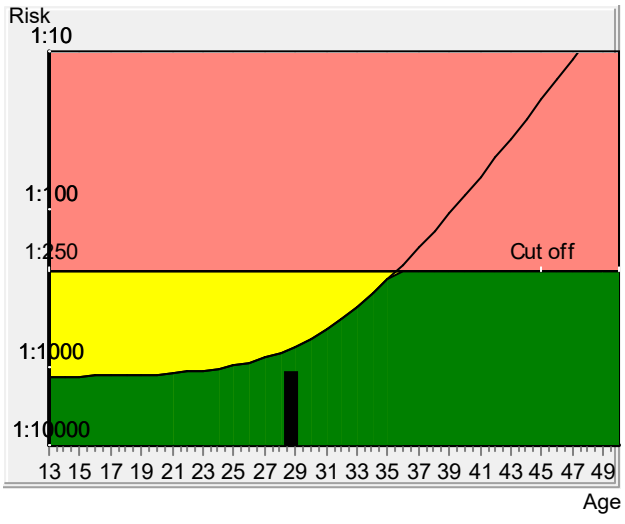
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	67	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	3.39 mIU/mL	0.59
fb-hCG	63.3 ng/ml	2.03

Risks at sampling date

Age risk	1:769
Biochemical T21 risk	1:258
Combined trisomy 21 risk	1:1088
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	13 + 3
Method	CRL Robinson
Scan date	14/09/20
Crown rump length in mm	76.5
Nuchal translucency MoM	1.02
Nasal bone	present
Sonographer	T
Qualifications in measuring NT	KEERTHI

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 1088 women with the same data, there is one woman with a trisomy 21 pregnancy and 1087 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