SagePath Labs Private Limited

Prisca 5.0.2.37

Date of report: 12-08-2020

Patient data

Name	MRS. SAHITHI	Patient ID 20015637	20015637
Birthday	01-06-1996	Sample ID	200156377
Age at delivery	24.7	Sample Date	10-08-2020
Gestational age	13 + 4		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21 pregancies	unknown
Weight	56	diabetes	unknown		
Smoker	unknown	Origin	Asian		

Biochemical data

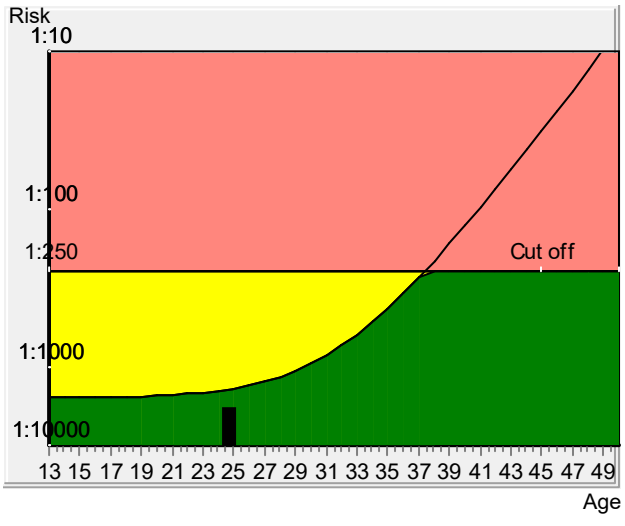
Parameter	Value	Corr. MoM
PAPP-A	7.62 mIU/ml	1.79
fb-hCG	31.2 ng/ml	0.87

Risks at term

Age risk	1:1394
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 4
Method	CRL Robinson
Scan date	06-08-2020
Crown rump length in mm	66.8
Nuchal translucency MoM	1.24
Nasal bone	present
Sonographer	DR. DEEPSHIKHA KHANNA
Qualifications in measuring NT	DMRD



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!

Trisomy 13/18 + NT

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

Sign of Physician

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