SagePath Labs Pvt Ltd

Prisca 5.0.0.10

Date of report: 14/09/20

Patient data

Name	Mrs. P.RANI	Patient ID	0162009110001
Birthday	15/04/87	Sample ID	20027634
Age at sample date	33.4	Sample Date	11/09/20

Correction factors

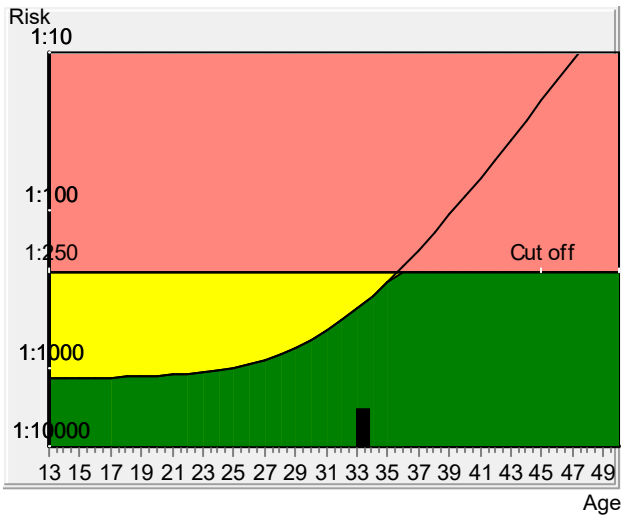
Fetuses	1	IVF	no	Previous trisomy 21 pregancies	unknown
Weight	47	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	3.84 mIU/mL	0.39
fb-hCG	6.45 ng/ml	0.20

Risks at sampling date

Age risk	1:393
Biochemical T21 risk	1:2037
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	1:576



Ultrasound data

Gestational age	13 + 2
Method	CRL Robinson
Scan date	08/09/20
Crown rump length in mm	74.56
Nuchal translucency MoM	0.82
Absent nasal bone	no
Sonographer	NA
Qualifications in measuring NT	NA

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 free beta HCG level is low.
The PAPP-A level is low.
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 18

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:576, which represents a low risk.

Sign of Physician

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