Prisca 5.1.0.17

Date of report: 26/03/22

SELF

Patient data

Name	Mrs. RUPINDER KAUR	Patient ID	01172203250015
Birthday	18/07/94	Sample ID	L3333524
Age at delivery	28.2	Sample Date	25/03/22
Gestational age	12 + 3		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	43	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

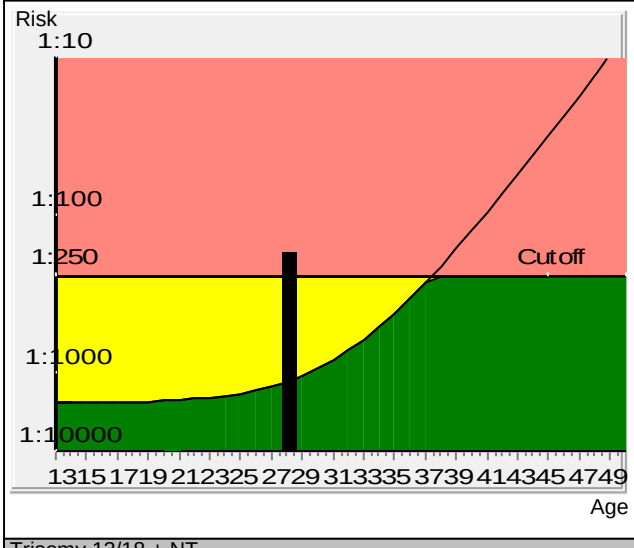
Parameter	Value	Corr. MoM
PAPP-A	2.15 mIU/mL	0.40
fb-hCG	223 ng/mL	4.92

Risks at term

Age risk	1:1146
Biochemical T21 risk	>1:50
Combined trisomy 21 risk	1:173
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	11 + 4
Method	CRL Robinson
Scan date	19/03/22
Crown rump length in mm	51.3
Nuchal translucency MoM	0.73
Nasal bone	present
Sonographer	DR.DEEPAK BANSAL.
Qualifications in measuring NT	MD



Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 173 women with the same data, there is one woman with a trisomy 21 pregnancy and 172 women with not affected pregnancies.
The free beta HCG level is high.
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 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