KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.1.0.17

Date of report: 08-11-2020

Patient data

Name	MRS. RANDHIR	Patient ID	2011220140/AMB
Birthday	01-01-1977	Sample ID	2011220140/AMB
Age at sample date	43.8	Sample Date	05-11-2020
Gestational age	12 + 3		

Correction factors

Fetuses	1	IVF	yes	Previous trisomy 21 pregnancies	no
Weight	75.9	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

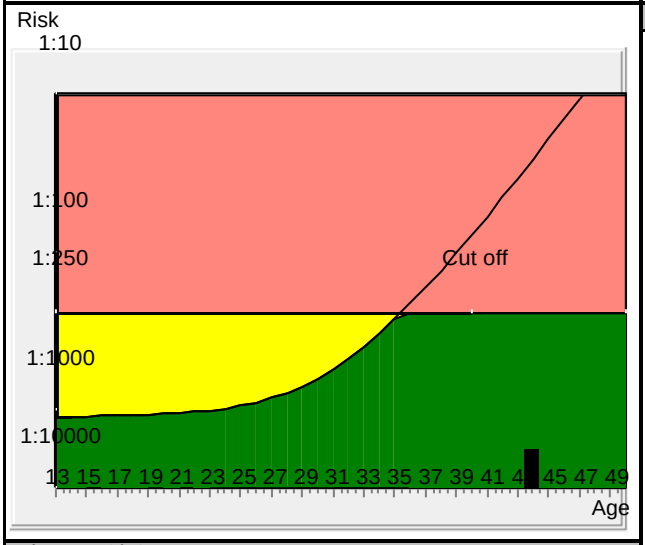
Parameter	Value	Corr. MoM
PAPP-A	1.64 mIU/ml	0.57
fb-hCG	13.3 ng/ml	0.31

Risks at sampling date

Age risk	1:27
Biochemical T21 risk	1:401
Combined trisomy 21 risk	1:1838
Trisomy 13/18 + NT	1:527

Ultrasound data

Gestational age	12 + 3
Method	CRL Robinson
Scan date	05-11-2020
Crown rump length in mm	62.2
Nuchal translucency MoM	0.93
Nasal bone	present
Sonographer	.
Qualifications in measuring NT	M.D



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:527, which represents a low risk.

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 1838 women with the same data, there is one woman with a trisomy 21 pregnancy and 1837 women with not affected pregnancies.

The free beta HCG 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!

Sign of Physician

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