KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.1.0.17

Date of report: 26-02-2020

Patient data

Name	MRS. MAMTA	Patient ID	2002220845/AMB
Birthday	23-06-1992	Sample ID	2002220845/AMB
Age at delivery	28.2	Sample Date	24-02-2020
Gestational age	12 + 1		

Correction factors

Fetuses	1	IVF	yes	Previous trisomy 21 pregnancies	no
Weight	59.8	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	1 mIU/ml	0.34
fb-hCG	110 ng/ml	2.91

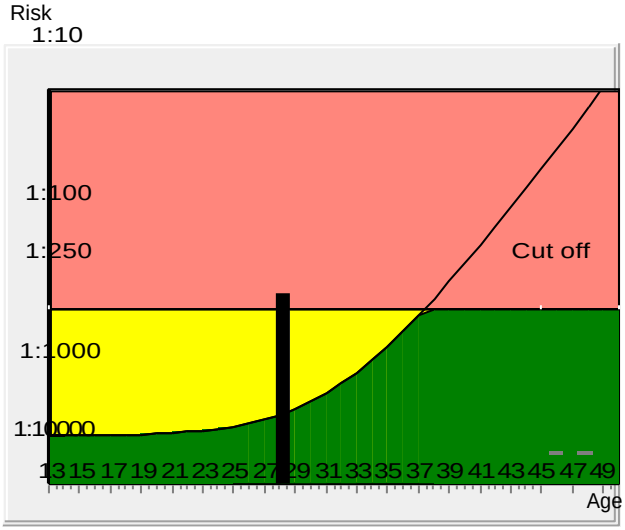
Risks at term

Age risk	1:1146
Biochemical T21 risk	>1:50
Combined trisomy 21 risk	1:195
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 0
Method	CRL Robinson
Scan date	23-02-2020
Crown rump length in mm	56.8
Nuchal translucency MoM	0.94
Nasal bone	present
Sonographer	.
Qualifications in measuring NT	MD

Risk



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 195 women with the same data, there is one woman with a trisomy 21 pregnancy and 194 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