KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.1.0.17

Date of report: 12-09-2019

Patient data

Name	MRS. PARWINDER KAUR	Patient ID	1909220722/AMB
Birthday	25-08-1996	Sample ID	1909220722/AMB
Age at delivery	23.6	Sample Date	11-09-2019
Gestational age	12 + 6		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	53.4	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	2.65 mIU/ml	0.59
fb-hCG	79.6 ng/ml	2.30

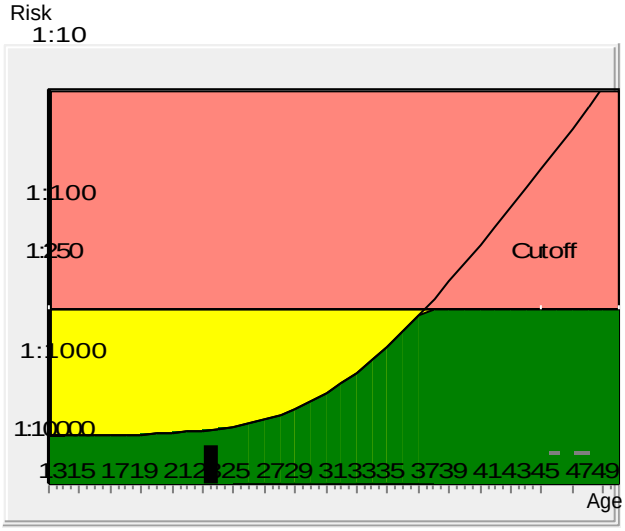
Risks at term

Age risk	1:1444
Biochemical T21 risk	1:357
Combined trisomy 21 risk	1:2287
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 5
Method	CRL Robinson
Scan date	10-09-2019
Crown rump length in mm	66
Nuchal translucency MoM	0.77
Nasal bone	present
Sonographer	.
Qualifications in measuring NT	MD

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 2287 women with the same data, there is one woman with a trisomy 21 pregnancy and 2286 women with not affected pregnancies.

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