KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

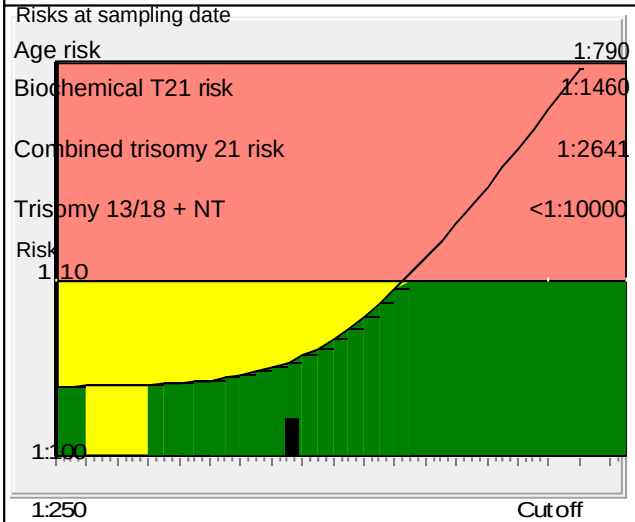
Date of report: 02-05-2018

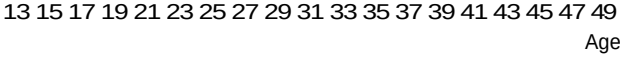
Patient data			
Name	MRS. PARMINDER KAUR	Patient ID	
Birthday	14-12-1989	Sample ID	1804220693/AMB
Age at sample date	28.4	Sample Date	28-04-2018
Gestational age	13 + 2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight	66	diabetes	unknown	pregancies	
Smoker	unknown	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 2
PAPP-A	2.8 mIU/ml	0.69	Method	CRL Robinson
fb-hCG	35.1 ng/ml	1.17	Scan date	28-04-2018
			Crown rump length in mm	73
			Nuchal translucency MoM	1.27
			Nasal bone	unknown
			Sonographer	.
			Qualifications in measuring NT	MD

Risks at sampling date	
Age risk	1:790
Biochemical T21 risk	1:1460
Combined trisomy 21 risk	1:2641
Trisomy 13/18 + NT	<1:10000





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 2641 women with the same data, there is one woman with a trisomy 21 pregnancy and 2640 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