KOS DIAGNOSTIC LAB 6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 25-10-2018

Patient data				
Name	MRS. DEEPIKA Patient ID			1810220442/AMB
Birthday	19-09-1986	Sample ID		1810220442/AMB
Age at sample date	32.1	Sample Date		24-10-2018
Gestational age	12 + 4			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 63	diabetes	no pregancies		
Smoker no	Origin	Asian		
Biochemical data	Ultrasound data			
Parameter Value	Corr. MoM	Gestational age 12 + 2		12 + 2
PAPP-A 2.69 mIU/m	l 0.81	Method CRL Rob		CRL Robinson
fb-hCG 19.8 ng/ml	0.57	Scan date		22-10-2018
Risks at sampling date		Crown rump length in mm		60.1
Age risk	1:478	Nuchal translucency MoM		1.22
Biochemical T21 risk	1:6022	Nasal bone		present
Combined trisomy 21 risk	<1:10000	Sonographer		DR. VIKAS KAUSHAL
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		MD
Risk 1:10		Trisomy 21	ated risk for Trisomy	
1:1000 1:1000 1:110	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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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