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

5.0.2.37

Date of report: 16-12-2016

Prisca

Patient data				
Name	MRS. JYOTI			1612220184/AMB
Birthday	09-05-1988	Sample ID		1612220184/AMB
Age at sample date	28.6	Sample Date		15-12-2016
Gestational age	12 + 2			
Correction factors				
Fetuses 1	IVF	yes	Previous trisomy 21	no
Weight 77	diabetes	no pregancies		
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age		12 + 2
PAPP-A 2.84 mIU/m	l 1.22	Method		CRL Robinson
fb-hCG 25.9 ng/ml	0.76			15-12-2016
Risks at sampling date			length in mm	59.8
Age risk	1:746	Nuchal translucency MoM		0.77
Biochemical T21 risk		Nasal bone		present
Combined trisomy 21 risk				DR. POONAM LOOMBA
Trisomy 13/18 + NT Risk	<1:10000		s in measuring NT	MD
110 1:10 1:100 1:250 Cut off 1:000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, 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 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