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

Prisca 5.0.2.37

Date of report: 01/03/19

Patient data				
Name	NAVNEET KAUF	Patient ID		211902280001
Birthday	19/01/91	Sample ID		211902280001
Age at sample date	28.0	Sample Date		28/02/19
Gestational age	12 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 77	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data	Ultrasound data			
Parameter Value	Corr. MoM Gestational age 12 + 0		12 + 6	
PAPP-A 2.0 mIU/m	nl 0.64	0.64 Method CRL Robinson		
fb-hCG 59.8 ng/ml	1.78	1.78 Scan date		26/02/19
Risks at sampling date	, <u> </u>		Crown rump length in mm	
Age risk	1:970 Nuchal trans		slucency MoM	0.69
Biochemical T21 risk			Nasal bone pr	
Combined trisomy 21 risk 1:3320		Sonographer		
Trisomy 13/18 + NT <1:10000		Qualifications in measuring NT MD		
1:100 1:250 1:1000 1:1000 1:1500 1:1000 1:1500 1:1000 1:1500 1:1000 1:1500 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 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 3312 women with the same data, there is one woman with a trisomy 21 pregnancy and 3311 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!			

Sign of Physician