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

5.0.2.37

Date of report: 10-11-2018

Prisca

Patient data			
Name	MRS. SHYAMWATI	Patient ID	1811220015/AMB
Birthday	01-01-1989	Sample ID	1811220015/AMB
Age at sample date	29.8	Sample Date	e 02-11-2018
Gestational age	12 + 2		
Correction factors			
Fetuses 1	IVF		Previous trisomy 21 no
Weight 53	diabetes	no	pregancies
Smoker no	Origin	Asian	
Biochemical data		Ultrasound da	lata
Parameter Value	Corr. MoM	Gestational a	age 12 + 2
PAPP-A 0.95 mIU/n	nl 0.26	Method	CRL Robinson
fb-hCG 45.1 ng/ml	1.17	Scan date	02-11-2018
Risks at sampling date		Crown rump	p length in mm 59
ge <u>risk 1:653</u>		Nuchal translucency MoM 0.53	
Biochemical T21 risk	1:80	Nasal bone	present
Combined trisomy 21 risk 1:587		Sonographe	er DR. (MRS.) POONAM LOOMBA
Trisomy 13/18 + NT	1:7039	Qualificatior Trisomy 21	ns in measuring NT MD
Risk 1 10 1:100 1:1000 1:1000 1:10000 1:10000 1:10000	5 37 39 41 43 45 47 49 Age	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 587 women with the same data, there is one woman with a trisomy 21 pregnancy and 586 women with not affected pregnancies. The PAPP-A level is low. 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

The calculated H5k 10		
below <b>translucency) is 1:7039</b>	, which we presents a low risk Risk	above cut off

Sign of Physician