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

Prisca 5.0.2.37

Date of report: 22-08-2017

Birthday         01-02-1991         Sample ID         1708220571/AMB           Age at sample date         26.6         Sample Date         21-08-2017           Gestational age         12 + 6         Sample Date         21-08-2017           Correction factors           Fetuses         1 IVF         no Previous trisomy 21 no pregancies         no pregancies           Smoker         no Origin         Asian         Asian           Biochemical data         Ultrasound data           Parameter         Value         Corr. MoM Gestational age         12 + 6           PAPP-A         6.46 mIU/ml         1.74 Method         CRL Robinson           fb-hCG         35.6 ng/ml         1.09 Scan date         21-08-2017           Risks at sampling date         Crown rump length in mm         68.3           Age risk         1:896         Nuchal translucency MoM         0.47	Patient data						
Age at sample date  Gestational age  12 + 6  Correction factors  Fetuses  1 IVF  no Previous trisomy 21 no Weight 63 diabetes Smoker no Origin  Asian  Ultrasound data  Parameter  Value  Corr. MoM PAPP-A 6.46 mIU/ml 1.74  Method CRL Robinson fb-hCG 35.6 ng/ml 1.09 Risks at sampling date  Age risk 1:10000 Combined trisomy 21 risk 1:10000 Risk	Name	MRS. DEEPIKA SINGH			Patient ID		1708220571/AMB
Gestational age 12 + 6  Correction factors  Fetuses 1 IVF no Previous trisomy 21 no pregancies  Smoker no Origin Asian  Biochemical data  Parameter Value Corr. MoM Gestational age 12 + 6  PAPP-A 6.46 mIU/ml 1.74 Method CRL Robinson 5 can date 21-08-2017  Risks at sampling date Crown rump length in mm 68.3  Age risk 1:896 Biochemical T21 risk 1:10000  Combined trisomy 21 risk 1:10000  Risks 1:10000  Risks 1:10000  Risks 1:10000  Combined trisomy 21 risk 1:10000  Risk 1:10  Combined trisomy 21 risk 1:10000  Risk 1:10  Combined trisomy 21 risk 1:10000  Risk 1:10  Cut off 1:250  Cut off 1:10000  1:1000  1:1000  1:1000  1:11000  1	Birthday	01-02-1991		Sample ID		1708220571/AMB	
Fetuses 1 IVF no Previous trisomy 21 no pregancies  Smoker no Origin Asian  Biochemical data  Parameter Value Corr. MoM Gestational age 12 + 6 PAPP-A 6.46 mIU/ml 1.74 Biochemical T21 risk 21:0000 Combined trisomy 21 risk 1:10000 Risk 1:100	Age at sample date	26.6		Sample Date	Э	21-08-2017	
Fetuses 1 IVF no Previous trisomy 21 no Pregancies  Smoker no Origin Asian  Biochemical data  Parameter Value Corr. MoM Gestational age 12 + 6  PAPP-A 6.46 mIU/ml 1.74 Method CRL Robinson fb-hCG 35.6 ng/ml 1.09  Risks at sampling date Crown rump length in mm 68.3  Age risk 1:896  Combined trisomy 21 risk 1:10000  Combined trisomy 21 risk 1:10000  Risks 1:10000  1:1000  1:1000  1:1000  1:1000  1:1000  1:1000  1:110	Gestational age	12 + 6					
Weight 63 diabetes no Origin Asian  Biochemical data  Parameter Value Corr. MoM Gestational age 12 + 6  PAPP-A 6.46 mIU/ml 1.74 Method CRL Robinson to-hCG 35.6 ng/ml 1.09  Risks at sampling date Crown rump length in mm 68.3  Age risk 1:896 Combined trisomy 21 risk 1:10000  Combined trisomy 21 risk 4:1:10000  Risk 1:10  Risk 1:10  Cut off 1:10  Cut of	Correction factors						
Smoker no Origin Asian  Biochemical data  Parameter Value Corr. MoM Gestational age 12 + 6  PAPP-A 6.46 mIU/ml 1.74  Method CRL Robinson fb-hCG 35.6 ng/ml 1.09  Scan date 21-08-2017  Scan date 21-08-2017  Crown rump length in mm 68.3  Age risk 1:896  Biochemical T21 risk <1:10000  Combined trisomy 21 risk <1:10000  Risk 1:10000  Risk 1:10000  Risk 1:10000  1:1000  1:1000  1:1000  1:1000  1:1000  1:1000  1:1000  1:1000  1:1	Fetuses	1	IVF		no		no
Biochemical data  Parameter Value Corr. MoM PAPP-A 6.46 mIU/ml 1.74 Ib-nCG 35.6 ng/ml 1.09 Risks at sampling date  Age risk 1:896 Combined trisomy 21 risk 4:10000 Combined trisomy 21 risk 4:10000 Combined trisomy 13/18 + NT 4:10000 Risks 1:10000 Risks 1:	Weight	63	diabetes		no	pregancies	
Parameter Value Corr. MoM Gestational age 12 + 6 PAPP-A 6.46 mIU/ml 1.74 fb-hCG 35.6 ng/ml 1.09 Risks at sampling date  Age risk 1:896 Combined trisomy 21 risk <1:10000 Combined trisomy 21 risk <1:10000 Risk Trisomy 13/18 + NT	Smoker	no	Origin		Asian		
PAPP-A 6.46 mIU/ml 1.74 fb-hCG 35.6 ng/ml 1.09 Scan date 21-08-2017 Crown rump length in mm 68.3 Age risk 1:896 Biochemical T21 risk <1:10000 Combined trisomy 21 risk <1:10000 Tisomy 13/18 + NT	Biochemical data				Ultrasound da	ata	
Risks at sampling date  Age risk Biochemical T21 risk Crown rump length in mm Combined trisomy 21 risk C1:10000 Risk 1:10 Risk Risk Risk Risk Risk Risk Risk Risk	Parameter	Value		Corr. MoM	Gestational	age	12 + 6
Risks at sampling date  Age risk  1:896 Biochemical T21 risk  Combined trisomy 21 risk  1:10000 Risk  1:10  Cut off  1:10000  1:10000  1:10000  1:10000  1:10000  1:10000  1:10000  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal-translucency) is < 1:10000, which represents a low  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal-translucency) is < 1:10000, which represents a low  Crown rump length in mm  68.3  Nuchal translucency MoM  0.47  Nasal bone  PR. POONAM LOOMBA  Qualifications in measuring NT  MD  Trisomy 21  The ealculated 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!	PAPP-A	6.46 mIU/m	l	1.74	Method		CRL Robinson
Age risk Biochemical T21 risk Combined trisomy 21 risk C1:10000 Risk Trisomy 13/18 + NT C1:10000 Risk  1:10 Cut off  Cut off  Cut off  Cut off  Trisomy 13/18 + NT Tre calculated risk for trisomy 13/18 (with nuchal-translucency) is < 1:10000, which represents a low  Nuchal translucency MoM O.47 Nasal bone Present Sonographer DR. POONAM LOOMBA Qualifications in measuring NT MD Trisomy 21 Trisomy 21 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!	fb-hCG	35.6 ng/ml		1.09	Scan date		21-08-2017
Biochemical T21 risk	Risks at sampling date				Crown rump	length in mm	68.3
Combined trisomy 21 risk  Trisomy 13/18 + NT  Cut off  Cut off  1:1000	Age risk			1:896	Nuchal trans	slucency MoM	0.47
Trisomy 13/18 + NT	Biochemical T21 risk			<1:10000	Nasal bone		present
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!  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low	Combined trisomy 21 ris	sk		<1:10000	Sonographe	er	DR. POONAM LOOMBA
Trisomy 13/18 + NT  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!  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal-translucency) is < 1:10000, which represents a low	Trisomy 13/18 + NT			<1:10000	Qualification	ns in measuring NT	MD
transtucency) 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!  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal-translucency) is < 1:10000, which represents a low					Trisomy 21		
	1:250  1:1000  1:10000  13 15 17 19 21 23 25  Trisomy 13/18 + NT  The calculated risk for translucency) is < 1:10	r trisomy 13	√18 (with r	3 45 47 49 ''''''''''''''''''''''''''''''''''''	translucen low risk.  After the re expected th same data, pregnancy. The calcula of the inforr Please note approaches The patient was done a 18: 511-52; The laborat on the risk	sult of the Trisomy 2: nat among more than there is one woman atted risk by PRISCA of mation provided by the that risk calculation is and have no diagnot combined risk presunccording to accepted 3 (1998)). The sassessment ! Calculation is calculation of the combined risk presunccording to accepted 3 (1998).	1 test (with NT) it is 10000 women with the with a trisomy 21 depends on the accuracy he referring physician. s are statistical ostic value! heres the NT measurement d guidelines (Prenat Diagn responsible for their impact

Sign of Physician

below cut off Below Cut Off, but above Age Risk above cut off