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

Prisca 5.0.2.37

Date of report: 22-07-2017

Weight 80 diabetes no Origin Asian  Biochemical data  Parameter Value Corr. MoM Gestational age 11 + 4  PAPP-A 3.1 mIU/ml 1.90 Method CRL Robinsor fb-hCG 20.4 ng/ml 0.53  Risks at sampling date Crown rump length in mm 49.8  Age risk 1:23 Nuchal translucency MoM 0.67  Biochemical T21 risk 1:1913 Nasal bone presen Combined trisomy 21 risk 1:7443 Sonographer	Patient data					
Age at sample date  44.2 Correction factors  Fetuses 1   VF   no pregancies  Smoker no Origin Asian  Biochemical data  PAPP-A 3.1 mIU/ml 1.90 Method CRL Robinsor fb-hCG 20.4 ng/ml 0.53  Biochemical T21 risk 1:1913  Bioc	Name		MRS. ANTAKSHRI	Patient ID	170722	0588/AMB
Gestational age  11 + 4  Correction factors  Fetuses 1   IVF   no pregancies  Smoker no Origin   Asian  Biochemical data  Parameter   Value   Corr. MoM   Gestational age   11 + 4  PAPP-A   3.1 mIU/ml   1.90   Method   CRL Robinsor   Crown rump length in mm   49.8    Age risk   1:23   Biochemical T21 risk   1:7443   Sara	Birthday		15-05-1973	Sample ID	170722	0588/AMB
Correction factors  Fetuses 1   IVF   no pregancies  Smoker   no Origin   Asian  Biochemical data  Parameter   Value   Corr. MoM   Gestational age   11 + 2   PAPP-A   3.1 mIU/ml   1.90   Method   CRL Robinsor   Crown rump length in mm   4.92   Age risk   1.23   Biochemical T21 risk   1.1943   Combined trisomy 21 risk   1.1943   Combined trisomy 21 risk   1.1943   Combined trisomy 21 risk   1.1940   Risk   1.190   Asian   Value   Correct   Crown rump length in mm   4.92   Nuchal translucency MoM   0.67   Nasal bone   presen   Sonographer   Crown rump length in mm   4.92   Value   Crown rump length in mm   4.92   Value   Crown rump length in mm   4.93   Value   Trisomy 13/18 + NT   Value   Value   Value   Trisomy 13/18 + NT   Value   Value   Value   Value   Value   Value   Corr. MoM   Value   Valu	Age at sample date	44.2		Sample Date	Sample Date 21-07-20	
Fetuses 1   IVF   no diabetes   no origin   Nasian   Previous trisomy 21   no pregancies   no	Gestational age		11 + 4			
Weight 80 diabetes no Origin Asian    Parameter   Value   Corr. MoM   Gestational age   11 + 2	Correction factors					
Smoker no Origin Asian  Biochemical data  Parameter Value Corr. MoM Gestational age 11+4  PAPP-A 3.1 mIU/ml 1.90  Method CRL Robinsor Crown rump length in mm 49.8  Age risk 1:23  Biochemical T21 risk 1:1913  Combined trisomy 21 risk 1:7443  Trisomy 13/18 + NT 4:1000  1:1000  1:1000  1:1000  1:1000  1:1000  1:1000  Trisomy 13/18 + NT 4:5 4744  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal approaches and have no diagnostic value!  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal approaches and have no diagnostic value!  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal approaches and have no diagnostic value!  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal approaches and have no diagnostic value!  The calculated risk for trisomy 13/18 (with nuchal approaches and have no diagnostic value!  The calculated risk for trisomy 13/18 (with nuchal approaches and have no diagnostic value!  The calculated risk for trisomy 13/18 (with nuchal approaches and have no diagnostic value!	Fetuses	1	IVF	no	Previous trisomy 21	no
Biochemical data  Parameter  Value  Corr. MoM  Gestational age  11 + 4  PAPP-A  3.1 mIU/ml  1.90  Method  CRL Robinsor  Scan date  21-07-2017  Crown rump length in mm  49.8  Nuchal translucency MoM  Nuchal translucency MoM  Nuchal translucency MoM  Ocorr risomy 13/18 + NT  1:1000  1:1000  1:1000  Trisomy 13/18 + NT  Age  Trisomy 13/18 + NT  1:1000  1:1000  1:1000  Trisomy 13/18 + NT  Trisomy 13/18 + NT  1:1000  1:1000  1:1000  Trisomy 13/18 + NT  Trisomy 13/18 trisomy 13/18 (with nuchal according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).  Trisomy 13/18 + NT  Trisomy 13/18 + NT  Trisomy 13/18 + NT  Trisomy 13/18 + NT  Trisomy 13/18 trisomy 13/18 (with nuchal according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).  Trisomy 13/18 + NT  Trisomy 13/18 + NT  Trisomy 13/18 trisomy 13/18 (with nuchal according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).  Trisomy 13/18 + NT	Weight	80	diabetes	no	pregancies	
Parameter Value Corr. MoM  PAPP-A 3.1 mIU/ml 1.90  tb-hCG 20.4 ng/ml 0.53  Risks at sampling date  Age risk 1:23  Biochemical T21 risk 1:1913  Combined trisomy 21 risk 1:1900  Risk 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 < 1:1000  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:1000, which represents a low  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:1000, which represents a low	Smoker	no	Origin	Asian		
PAPP-A 3.1 mIU/ml 1.90 fb-hCG 20.4 ng/ml 0.53 Risks at sampling date  Age risk 1:23 Biochemical T21 risk 1:1913 Combined trisomy 21 risk 1:7443 Risk 1:1000  1	Biochemical data			Ultrasound da	ata	
Risks at sampling date  Age risk Age risk Biochemical T21 risk 1:1913 Combined trisomy 21 risk 1:7443 Trisomy 13/18 + NT 1:100  1:1000 1:1000 1:1000 1:11000  Age risk 1:23 Biochemical T21 risk 1:1913 Combined trisomy 21 risk 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:11	Parameter	Value	Corr. MoN	Gestational	age	11 + 4
Age risk  Age risk  1:23 Biochemical T21 risk  Combined trisomy 21 risk  1:7443 Trisomy 13/18 + NT  1:10000  1:10000	PAPP-A	3.1 mIU/ml	1.90	Method	CRI	Robinson
Age risk Biochemical T21 risk 1:1913 Combined trisomy 21 risk 1:7443 Trisomy 13/18 + NT Risk 1:10  1:100  1:1000 1:1000 1:1000 1:11000	fb-hCG	20.4 ng/ml	0.53	Scan date	2	21-07-2017
Biochemical T21 risk Combined trisomy 21 risk 1:7443 Trisomy 13/18 + NT Risk 1:10  1	Risks at sampling date			Crown rump	length in mm	49.8
Combined trisomy 21 risk  Trisomy 13/18 + NT  Altifo  Risk  1:10  Trisomy 21  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 pregnancy and 7442 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!  Trisomy 13/18 + NT  The calculated risk for trisomy 13/18 (with nuchal-translucency) is <1:1000, which represents a low	Age risk		1:23	Nuchal trans	slucency MoM	0.67
Trisomy 13/18 + NT	Biochemical T21 risk		1:1913	Nasal bone		present
Trisomy 21  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 7443 women with the same data, there is one woman with a trisomy 21 pregnancy and 7442 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 calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low	Combined trisomy 21 r	isk	1:7443	Sonographe	er	
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 7443 women with the same data, there is one woman with a trisomy 21 pregnancy and 7442 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)).  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
1:100 1:200 1:200 1:200 1:200 1:200 1:200 1:200 1:200 1:200 1:2100 1:2100 1:2100 1:220 1:2				Trisomy 21		
	it is it is ie data, y and 7442 accuracy sician. assurement enat Diagn heir impact					

Sign of Physician

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