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

Prisca 5.0.2.37

Date of report: 19-07-2017

Biochemical T21 risk Combined trisomy 21 risk Trisomy 13/18 + NT <1:1000 Risk 1:10 1:100 1:100 1:1000 1:200 Cut off Trisomy 21 risk 1:2418 Nasal bone Sonographer Qualifications in measuring NT MD 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)).	Patient data					
Age at sample date 29.7 Gestational age 11 + 5 Correction factors Fetuses 1 IVF no diabetes no Origin Asian Biochemical data Parameter Value Corr. MoM Gestational age 11 + 4 PAPP-A 1.9 mIU/ml 1.08 Method CRL Robinson b-hCG 50.1 ng/ml 1.34 Risks at sampling date Crown rump length in mm 51 Age risk 1:2418 Combined trisomy 21 risk 1:2418 Combined trisomy 21 risk 1:10000 Risk 1:10 Cut off 1.1000 1.1000 1.1000 1.1000 1.1100	Name	М	RS. BALJEET KAUR	Patient ID 170		1707220520/AMB
Gestational age Tetuses IVF no Gright T9.5 Glabetes Smoker No Origin Asian Biochemical data Parameter Value Corr. MoM Gestational age 11 + 4 PAPP-A 1.9 mIU/ml 1.08 Method CRL Robinson fb-hCG 50.1 ng/ml 1.34 Scan date 17-07-2017 Risks at sampling date Crown rump length in mm 51 Nuchal translucency MoM 0.95 Biochemical T21 risk 1:2418 Nasal bone Combined trisomy 21 risk 1:10000 Risk 1:40 Trisomy 13/18 + NT Combined trisomy 21 risk Combined trinsomy 21 risk Combined trisomy 21 risk Com	Birthday	19-10-1987		Sample ID		1707220520/AMB
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Weight 79.5 diabetes no Origin Asian Pregancies	Correction factors					
Smoker no Origin Asian Biochemical data Parameter Value Corr. MoM Gestational age 11+ 4 PAPP-A 1.9 mIU/ml 1.08 Method CRL Robinson 50.1 ng/ml 1.34 Scan date 17-07-2017 Risks at sampling date 51-2418 Sinchemical T21 risk 1:2418 Combined trisomy 21 risk 1:10000 Combined risk 1:10000	Fetuses	1	IVF	no	Previous trisomy 21	no
Biochemical data Parameter Value Corr. MoM Gestational age 11+ 4 PAPP-A 1.9 mIU/ml 1.08 Method CRL Robinson fb-hCG 50.1 ng/ml 1.34 Scan date 17-07-2017 Risks at sampling date Age risk 1:2418 Combined trisomy 21 risk 1:10000 Combined trisomy 21 risk 1:10000 Trisomy 13/18 + NT 2.110000 Combined trisomy 21 risk 1:10000 Combined risk for Trisomy 21 risk 1:10000 Combined risk	Weight	79.5	diabetes	no	pregancies	
Parameter Value Corr. MoM Gestational age 11+ 4 PAPP-A 1.9 mIU/ml 1.08 Method CRL Robinson fb-hCG 50.1 ng/ml 1.34 Risks at sampling date Age risk 1:646 Biochemical T21 risk 1:2418 Combined trisomy 21 risk 1:2418 Combined trisomy 21 risk 1:2418 Combined trisomy 21 risk 1:10000 Risk 1:10000 Trisomy 13/18 + NT 1:0000 Cut off Trisomy 13/18 + NT 1.0000 Cut off 1:10000 Cut off 1:100	Smoker	no	Origin	Asian		
PAPP-A 1.9 mIU/ml 1.08 fb-hCG 50.1 ng/ml 1.34 Scan date 17-07-2017 Risks at sampling date Age risk 1:646 Nuchal translucency MoM 0.95 Biochemical T21 risk 1:2418 Combined trisomy 21 risk 1:2418 Combined trisomy 21 risk 1:10000 Risk 1:10 1	Biochemical data			Ultrasound data		
fb-hCG 50.1 ng/ml 1.34 Risks at sampling date Age risk 1:646 Biochemical T21 risk 1:2418 Combined trisomy 21 risk 1:10000 Risk 1:10 Risk 1:10 Gt off Trisomy 13/18 + NT Combined trisomy 21 risk 1:2418 Combined trisomy 21 ris	Parameter	Value	Corr. MoN	1 Gestational	age	11 + 4
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Age risk Biochemical T21 risk 1:2418 Combined trisomy 21 risk 1:2418 Combined trisomy 21 risk 41:10000 Risk 1:10 Risk 1:10 1	fb-hCG	50.1 ng/ml	1.34			
Biochemical T21 risk Combined trisomy 21 risk 1:2418 Combined trisomy 21 risk 4:10000 Risk 1:10 1:1000 1:10	Risks at sampling date			Crown rump	length in mm	51
Combined trisomy 21 risk Trisomy 13/18 + NT Sonographer Qualifications in measuring NT 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	Age risk		1:646	Nuchal trans	slucency MoM	0.95
Trisomy 13/18 + NT Qualifications in measuring NT MD Trisomy 21 The ealeulated 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	Biochemical T21 risk		1:2418	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!	Combined trisomy 21 r	isk	<1:10000	Sonographe	er	•
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translucency) is < 1:10000, which represents a low risk.	1:1000 1:1000 13 15 17 19 21 23 25 Trisomy 13/18 + NT The calculated risk for translucency) is < 1:		5 37 39 41 43 45 47 49 Age /18 (with nuchal	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		
				<u> </u>	Sign of Physician	

Sign of Physician

above cut off

below cut off Below Cut Off, but above Age Risk