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

Prisca 5.0.2.37

Date of report: 11-08-2017

fb-hCG 32.1 ng/ml 1.09 Risks at sampling date Age risk 1:990 Biochemical T21 risk 4:10000 Combined trisomy 21 risk 4:10000 Risk 1:10 Risk 1:10 Age risk 5:10000 Combined trisomy 21 risk 5:10000 Risk 1:10 Age risk 6:10000 Risk 6:10000 Risk 7:10000 Risk 7:10000 Risk 1:10 Age risk 6:10000 Risk 6:10000 Risk 6:10000 Risk 7:10000 Risk 7:10000 Risk 7:10000 Risk 7:10000 Risk 7:10000 Risk 8:10000 Risk 8:10000 Risk 8:10000 Risk 8:10000 Risk 8:100000 Risk 8:100000 Risk 8:1000000000000000000000000000000000000	Patient data					
Age at sample date 25.2 Sample Date 08-08-2017 Gestational age 13 + 5 Correction factors	Name		MRS. ANJANA	Patient ID		
Gestational age 13 + 5 Correction factors Fetuses 1 IVF no pregancies Smoker no Origin Asian Biochemical data Parameter Value Corr. MoM Gestational age 13 + 5 PAPP-A 9.8 mIIU/ml 1.68 Method CRL Robinson 5th-hCG 32.1 ng/ml 1.09 Risks at sampling date Crown rump length in mm 8.0.3. Age risk 1.990 Biochemical T21 risk 1.10000 Combined trisomy 21 risk 1.10000 Risks 1.10000 Risks 1.10000 Risks 1.10000 1.1000 1.1000 1.1000 1.1000 1.1000 1.11000	Birthday	25-05-1992		Sample ID		1708220193/AMB
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Smoker no Origin Asian Biochemical data Parameter Value Corr. MoM Gestational age 13 + 5 PAPP-A 9.8 mIU/ml 1.68 Method CRL Robinson fb-hCG 32.1 ng/ml 1.09 Scan date 08-08-2017 Risks at sampling date 1.990 Biochemical T21 risk 1.990 Corbined trisomy 21 risk 1.10000 Crown rump length in mm 80.3 Nuchal translucency MoM 0.48 Nasal bone present Sonographer DR. POONAM LOOMBA 1.10000 Risk 1.10000 Risk 1.10000 1.1000 1.1000 1.1000 1.1000 1.1100	Fetuses	1	IVF	no		no
Biochemical data Parameter Value Corr. MoM Gestational age 13 + 5 PAPP-A 9.8 mIU/ml 1.68 108-08-2017 Risks at sampling date 1.990 Biochemical T21 risk 1.990 Combined trisomy 21 risk 1.1000 1.100 1.100 1.100 1.100 1.100 1.100 1.100 1.1000 1.100 1.10000	Weight	55	diabetes	no	pregancies	
Parameter Value Corr. MoM Gestational age 13 + 5 PAPP-A 9.8 mIU/ml 1.68 fb-hCG 32.1 ng/ml 1.09 Risks at sampling date Age risk 1.990 Combined trisomy 21 risk <1:10000 Combined trisomy 21 risk <1:10000 Trisomy 13/18 + NT	Smoker	no	Origin	Asian		
PAPP-A 9.8 mIU/ml 1.68 fb-hCG 32.1 ng/ml 1.09 Scan date 08-08-2017 Crown rump length in mm 80.3 Nuchal translucency MoM 0.48 Biochemical T21 risk 1:10000 Scan date 1:10000 Scan date 08-08-2017 Crown rump length in mm 80.3 Nuchal translucency MoM 0.48 Nasal bone present Scangrapher DR. POONAM LOOMBA Callifications in measuring NT MD 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The calculated risk py 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	Biochemical data			Ultrasound data		
fb-hCG 32.1 ng/ml 1.09 Risks at sampling date Age risk 1:990 Biochemical T21 risk <1:10000 Combined trisomy 21 risk <1:10000 Trisomy 13/18 + NT	Parameter	Value Corr. MoM		Gestational age 13 + 5		
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Age risk 1:990 Biochemical T21 risk < 1:10000 Combined trisomy 21 risk	fb-hCG	32.1 ng/ml	1.09	Scan date 08-08-2017		
Biochemical T21 risk	Risks at sampling date			Crown rump length in mm 80.3		
Combined trisomy 21 risk Trisomy 13/18 + NT Cut off 1:10 Cut off 1:10 Cut off 1:10 Cut off 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 Sonographer DR. POONAM LOOMBA 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)). The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!	Age risk		1:990	Nuchal translucency MoM		0.48
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 r	mbined trisomy 21 risk <1:10000		Sonographer DR. POONAM LOOMBA		
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1:1000 1:250 Out off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:110000 1:1				Trisomy 21		
	1:250 Out off 1:10000 13 15 17 19 21 23 2 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low			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		

Sign of Physician

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