Date of Report 26-03-19 PRISCA 5.0.2.37

Birthday 10-02-91 Sample ID 102066. Age at delivery 28.1 Sample Date 23/03/20 Gestational age by 13+1 Correction factors Fetuses 1 IVF unknown Previous trisomy 21 unknow Pregnancies Smoker no Origin Asian Biochemical Data Ultrasound Data Parameter Value Corr Mont Gestational age PAPP-A 1.28 mIU/ml 0.42 Method CRI. (⇔Hadloe Ib-hCG 16.85 ng/ml 0.45 Scan Date 22-03-CRL measurements 64 Risks at sampling date Age Risk 1:804 Nasal bone Prese DR.Ruby RAH Combined Trisomy 21 Risk 1:2958 Sonographer DR.Ruby RAH Combined Trisomy 21 Risk 4:10000 Trisomy 13/18 < 1:10000 1:100 1:100 Cot off Trisomy 13/18 Cot off Trisomy 13/18 Cot off Trisomy 13/18 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 for 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 for 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 for 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 approach and have note that risk calculations are statistical approach and have note that risk calculations are statistical approach and have note that risk calculations are statistical approach and have note that risk calculations are statistical approach and have note that risk calculations are statistical approach and have note that risk calculations are statistical approach and have note that risk calculations are statistical approach and have note that risk calculations are statistical approach and have note that risk calculations are statistical approach an						PRISCA	5.0.2.37
Birthday Age at delivery Gestational age by 13+1 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) 13/18 (with nuchal translucency) 13/18 (with nuchal translucency of the information provided by the referring physician. Please note that risk calculations are statistical approach and have n diagnostic values! The calculated risk for trisomy 13/18 (with nuchal translucency was done according to accepted guidelines Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency was done according to accepted guidelines Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency was done according to accepted guidelines Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency was done according to accepted guidelines	Patient Data						
Age at delivery 28.1 Gestational age by 13+1 Correction factors Fetuses 1 IVF unknown Previous trisomy 21 unknown Pregnancies Smoker no Origin Asian Biochemical Data Parameter Value Corr Mon Gestational age 12- PAPP-A 1.28 mIU/ml 0.42 Method CRL (~Hadloe 16-hCG 16.85 ng/ml 0.45 Scan Date 22-03- CRL measurments 64- Risks at sampling date Age Risk 1:804 Nasal bone Prese Sonographer DR.RUBY RAH Combined Trisomy 21 Risk 1:2958 Combined Trisomy 21 Risk 4:10000 Risk 8: 1:10000 Risk 8: 1:10000 Risk 9: 1:10000 Risk 1:10000 Ri	Name	ľ	Mrs Bhav	ini Sharma	Patient ID		011903230245
Gestational age by 13+1 Correction factors Fetuses 1 IVF unknown Weight in kg 86 Diabetes no Origin Asian Biochemical Data Parameter Value Corr Mon PAPP-A 1.28 mIU/ml 0.42 fb-hCG 16.85 ng/ml 0.45 Risks at sampling date Age Risk 1:804 Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk 1:2958 Combined Trisomy 21 Risk 4:110000 Risk Risk Resk 1:00 Risk Resk 1:1000 Risk Risk (Irisomy 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 pregnancies Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal) Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal) Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal) Risk Risk (Irisomy 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 pregnancies	Birthday			10-02-91	Sample ID		10206658
Fetuses 1 IVF unknown Previous trisomy 21 unknown Pregnancies Smoker no Origin Asian Biochemical Data Parameter Value Corr Mon Corr Mo	Age at delivery			28.1	Sample Date		23/03/2019
Fetuses I I IVF unknown Weight in kg 86 Diabetes no Pregnancies March Previous trisomy 21 Unknown Previous trisomy 21 Unknown Previous trisomy 21 Unknown Pregnancies March Parameter Value	Gestational age by			13+1			
Weight in kg 86 Diabetes no Origin Asian Corr Mon Corr Mon Corr Mon	Correction factors						
Biochemical Data Parameter Value Corr Mom PAPP-A 1.28 mIU/ml 0.42 fb-hCG 16.85 ng/ml 0.45 Risks at sampling date Age Risk 1:804 Biochemical T21 Risk 1:2958 Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk 4:1:10000 Risk Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk 4:1:10000 Risk Down's Syndrome Risk (Trisomy 21 Screening) 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. Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal) Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal) Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal) Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal)	Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Parameter Value Corr Mom PAPP-A 1.28 mIU/ml 0.42 fb-hCG 16.85 ng/ml 0.45 Risks at sampling date Age Risk 1:804 Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk 1:2958 Combined Trisomy 21 Risk 4:1:10000 Risk Risk Down's Syndrome Risk (Trisomy 21 Screening) 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 (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. Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal) Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal)	Weight in kg	86	Diabetes		no	Pregnancies	
Parameter Value Corr Mon PAPP-A 1.28 mlU/ml 0.42 fb-hCG 16.85 ng/ml 0.45 Risks at sampling date Age Risk 1:804 Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk 1:2958 Combined Trisomy 21 Risk 4:10000 Risk Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for 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 approach and have n diagnostic values! Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal	Smoker	no	Origin		Asian		
PAPP-A 1.28 mIU/ml 0.42 fb-hCG 16.85 ng/ml 0.45 Risks at sampling date Age Risk 1:804 Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk 1:2958 Combined Trisomy 21 Risk 4:10000 Risk Risk Risk Risk Risk Risk Risk Ris	Biochemical Data				Ultrasound Data		
fb-hCG 16.85 ng/ml 0.45 Risks at sampling date Age Risk 1:804 Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk 4:10000 Trisomy 13/18	Parameter	Value		Corr Mom	Gestational ag	e	12+6
Risk 1:804 Age Risk 1:2958 Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk 4:10000 Trisomy 13/18 Combined Trisomy 21 Risk 4:10000 Risk 2 Down's Syndrome Risk (Trisomy 21 Screening) 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 approach and have note that risk of the risk pr	PAPP-A	1.28	mIU/ml	0.42	Method		CRL (<>Hadlock)
Risks at sampling date Age Risk Biochemical T21 Risk Combined Trisomy 21 Risk Combined Trisomy 13/18 Risk Risk Risk 1:10000 Risk 1:10000	fb-hCG	16.85	ng/ml	0.45	Scan Date		22-03-19
Age Risk Biochemical T21 Risk 1:2958 Combined Trisomy 21 Risk Combined Trisomy 21 Risk 71:10000 Risk Risk Down's Syndrome Risk (Trisomy 21 Screening) 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 approach and have n diagnostic values! Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal					CRL measurm	ents	64.6
Biochemical T21 Risk Combined Trisomy 21 Risk 1:2958 Sonographer Qualifications in measuring NT CON. RADIOLOGI Trisomy 13/18 Council Trisomy 21 Risk 1:1000 Risk Risk 1:10 Down's Syndrome Risk (Trisomy 21 Screening) 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 approach and have n diagnostic values! Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal	Risks at sampling date				Nuchal translucency MoM 0.91		
Combined Trisomy 21 Risk Trisomy 13/18 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 approach and have n diagnostic values! Trisomy 13/18 + NT The calculated risk for 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 approach and have n diagnostic values! Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal) Trisomy 13/18 + NT The calculated risk for 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 patient combined risk presumes the NT measurement was done according to accepted guidelines. Trisomy 13/18 + NT The calculated risk for trisomy 21 (with nuchal) The calculated risk for 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 approach and have n diagnostic values! The patient combined risk presumes the NT measurement was done according to accepted guidelines. Trisomy 13/18 + NT The calculated risk for trisomy 21 (with nuchal) The calculated risk for 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 for trisomy 21 test (with NT) it is expected that among more than 10000 women with the same dat	Age Risk			1:804	Nasal bone		Present
Risk Risk 1:10 1:100 Cut off 1:1000 Cut off 1:1000 1:1000 1:1000 1:1000 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 approach and have note that risk c	Biochemical T21 Risk			1:2958	Sonographer		DR.RUBY RAHUL
Risk 1:10 1:200 1:250 Cut off 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:1100000 1:110000 1:1100000 1:1100000000	Combined Trisomy 21 Risk			<1:10000	Qualifications	in measuring NT	CON. RADIOLOGIST
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 approach and have n diagnostic values! Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal	Trisomy 13/18			<1:10000			
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 approach and have n diagnostic values! The patient combined risk presumes the NT measurement was done according to accepted guidelines Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal	Risk				Down's Syndi	rome Risk (Trisomy 21	1 Screening)
two nelucion av) is < 1.10000, which nonnegonate a large size	1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (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 approach and have no diagnostic values! The patient combined risk presumes the NT		
transfucency) is < 1:10000, which represents a fow risk.	translucency) is < 1:10000, which represents a low risk.						