



Book a Test Online www.molq.in

Date of Report 03 - 12 - 2020PRISCA 5.0.2.37

Name MRS SONIA Patient IID 0.52011300022					
Birthday 15-08-1992 Age at delivery 28.3 Sample Date 30/11/2020 Gestational age 13+0 Correction factors Fetuses 1 IVF unknown Weight in kg 64.9 Diabetes unknown Smoker Unknown Origin Biochemical Data Ultrasound Data Parameter Value Corr Mom PAPP-A 4.3 miU/ml 0.89 Biochemical T21 risk 1:9122 Combined Trisony 21 Risk 31:1000 Risks at sampling date Crown rump length in mm 65.8 Nuchal translucency MOM 0.48 Biochemical T21 risk 1:9122 Nample ID 10792905 Unknown Previous trisomy 21 unknown Pregnancies Unknown Pre	Patient Data				
Age at delivery Gestational age 13+0 Correction factors Fetuses 1 IVF unknown Weight in kg 64.9 Diabetes unknown Smoker Unknown Origin Asian Biochemical Data Parameter Value Corr Mom PAPP-A 4.3 mIU/ml 0.89 Bi-hCG 28.6 ng/ml 0.66 Risks at sampling date Age Risk 1:788 Biochemical T21 risk 1:9122 Combined Trisony 21 Risk 4:10000 Risks 1:788 Nuchal translucency MOM 0.48 Biochemical T21 risk 1:9122 Nasal bone Present Combined Trisony 21 Risk 4:10000 Risks Down's Syndrome Risk (Trisomy 21 test (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) is test and any provided by the referring physician. Please note that the risk calculations are statistical approaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The laboratory cannot be hold responsible for their impact on the risk assessment Calculated risks have no diagnostic value!	Name MRS SONIA		Patient ID		052011300022
Gestational age 13+0 Correction factors Fetuses 1 IVF unknown Previous trisomy 21 unknown Weight in kg 64.9 Diabetes unknown Pregnancies unknown Origin Asian Ultrasound Data Gestational age 12+6 Gestational age Crack Pregnancies unknown Pregnancies unknown Origin Asian Pregnancies unknown Pregnancies unknown Pregnancies unknown Origin Asian Origin Asian Origin Data Gestational age 12+6 Gestational age 12+6 Scan date 30-11-2020 Scan date 30-11-2020 Scan date 30-11-2020 Scan date 30-11-2020 Gestational age 12+6 Scan date 30-11-2020 Scan date 40-11-2020 Scan	Birthday 15-08-1992		Sample ID		10792905
Fetuses 1 IVF unknown Weight in kg 64.9 Diabetes unknown Woight in kg 64.9 Diabetes unknown Woight in kg 64.9 Diabetes unknown Workingh IVF unknown Working	Age at delivery 28.3		Sample Date 30/11/2020		
Fetuses 1 IVF unknown Previous trisomy 21 unknown Pregnancies unkn	Gestational age 13+0				
Weight in kg 64.9 Diabetes Unknown Origin Asia Origin Asia Origin Asia Origin Asia Origin Asia Ultrasound Data Gestational age 12+6 PAPP-A 4.3 mIU/ml 0.89 Method CRL(◇Robinson fb-hCG 28.6 ng/ml 0.66 Scan date Crown rump length in mm 65.8 Age Risk 1:788 Nuchal translucency MOM 0.48 Biochemical T21 risk 1:9122 Nasal bone Present Combined Trisomy 21 Risk 1:10000 Risk Own'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 and 9999 women with not affected pregnancies. 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risk bave no diagnostic	Correction factors				
Biochemical Data Parameter Value Corr Mom PAPP-A 4.3 mIU/ml 0.89 Risks at sampling date Crown rump length in mm 65.8 Age Risk 1:788 Nuchal translucency MOM 0.48 Biochemical T21 risk 1:9122 Nasal bone Present Combined Trisomy 21 Risk 1:10000 Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk py PRSCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic on the risk assessment! Calculated risk have no diagnostic on the risk assessment! Calculated risk shave no diagnostic on the risk assessment! Calculated risk shave no diagnostic	Fetuses 1 IVF		unknown	Previous trisomy 21	unknown
Parameter Value Corr Mom PAPP-A 4.3 mIU/ml 0.89 Method CRL(S>Robinson B-hCG 28.6 ng/ml 0.66 Scan date 30-11-2020 Risks at sampling date Crown rump length in mm 65.8 Age Risk 1:788 Nuchal translucency MOM 0.48 Biochemical T21 risk 1:9122 Sonographer DRMEENU SOLANKI Trisomy 13/18 + NT <1:10000 Risk 2 Down's Syndrome Risk (Trisomy 21 Cwith NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 wenen with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The calculated risk for Trisomy 13/18 (with NT) is The calculated risk for Trisomy 13/18 (with NT) is	Weight in kg 64.9 Diabo	etes	unknown	Pregnancies	unknown
Parameter Value Corr Mom PAPP-A 4.3 mIU/ml 0.89 Method CRL(⋄ Robinson Bo-hCG 28.6 ng/ml 0.66 Scan date 30-11-2020 Risks at sampling date Crown rump length in mm 65.8 Age Risk 1:788 Nuchal translucency MOM 0.48 Biochemical T21 risk 1:9122 Combined Trisomy 21 Risk 4:10000 Sonographer DR.MEENU SOLANKI Trisomy 13/18 + NT 4:10000 Risk Down's Syndrome Risk (Trisomy 21 test (with NT) is below the cut off, which represents a low risk more than 10000 women with the same data, there is one woman with a trisomy 21 Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is Trisomy 13/18 + NT The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic	Smoker Unknown Origin	n	Asian		
PAPP-A 4.3 mIU/ml 0.89 BiohCG 28.6 ng/ml 0.66 Scan date 30-11-2020	Biochemical Data		Ultrasound Data		
Risks at sampling date Crown rump length in mm Crown rump length in me Crowletted States Stat	Parameter Value	Corr Mom	Gestational age	2	12+6
Risks at sampling date Crown rump length in mm Crown rump length in me Present Age by Calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low risk. After	PAPP-A 4.3 mIU/	ml 0.89	Method CRL(<>Robinson		
Age Risk Biochemical T21 risk 1:9122 Combined Trisomy 21 Risk 1:10000 Sonographer Qualification in measuring NT HMC 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic	fb-hCG 28.6 ng/ml	0.66	Scan date		30-11-2020
Biochemical T21 risk Combined Trisomy 21 Risk 1:9122 Nasal bone Present Sonographer DR.MEENU SOLANKI Trisomy 13/18 + NT Cut off 1:1000 Cut off 1:1000 Cut off 1:1000 Cut off Trisomy 13/18 + NT The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The calculated risk for Trisomy 13/18 (with NT) is	Risks at sampling date		Crown rump length in mm 65.8		
Combined Trisomy 21 Risk Trisomy 13/18 + NT Al:10000 Qualification in measuring NT HMC Risk 1:10 Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is	Age Risk	1:788	Nuchal translucency MOM		0.48
Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The calculated risk for Trisomy 13/18 (with NT) is	Biochemical T21 risk	1:9122	Nasal bone		Present
Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic	Combined Trisomy 21 Risk <1:1000		Sonographer DR.MEENU SOL		DR.MEENU SOLANKI
The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic	Trisomy 13/18 + NT <1:10000		Qualification in measuring NT HM		
The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is The calculated risk for Trisomy 13/18 (with NT) is			Down's Syndrome Risk (Trisomy 21 Screening)		
	1:10 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 Trisomy 13/18 + NT	cut off, which represents 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 and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic			