

*Free Home				
Sample Collection 9999 778 778				

Download "MOLQ" App on

Book a Test Online www.molq.in

Patient Data MRS. REKHA Patient ID 012201280344 Birthday 1.5/11/1994 Sample ID 11307448 Age at delivery 27.2 Sample Date 28/1/2022 Gestational age 13+2 Correction factors 28/1/2022 Fetuses 1 IVF unknown Previous trisomy 21 unknown Weight in kg 61 Diabetes NO Pregnancies unknown Smoker NO Origin Asian Asian 141 Parameter Value Corr Mom Gestational age 13+1 PAPP-A 2.78 mIU/ml 0.48 Method CRL (<>Robinson) fb-hCG 23.4 ng/ml 0.55 Scan date 27/01/2022 Risks at sampling date Nuchal translucency 2.1 Age Risk 1:869 Nuchal translucency MoM 1.15					Date of Report PRISCA	29/1/2022 5.2.0.13
Birthday 15/11/1994 Age at delivery 27.2 Sample Date 28/1/2022 Gestational age 13+2 Correction factors Fetuses 1 IVF unknown Previous trisomy 21 unknown Weight in kg 61 Diabetes NO Pregnancies unknown Smoker NO Origin Asian Biochemical Data Ultrasound Data Parameter Value Corr Mom B-hCG 23.4 ng/ml 0.48 Biochemical T21 risk 1:8213 Trisony 13/18+NT <1:10000 Risk Biochemical T21 risk 1:8213 Trisony 13/18+NT <1:10000 Risk The calculated risk (Trisony 21 Screening) 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 approaches and have no diagnostic value! The laboratory cannot be hold responsible for their inpact	Patient Data					
Age at delivery 27.2 Sample Date 28/1/2022 Gestational age 13+2 Correction factors Fetuses 1 Fetuses 1 IVF unknown Previous trisomy 21 unknown Weight in kg 61 Diabetes NO Pregnancies unknown Biochemical Data Ultrasound Data Eastional age 13+1 Parameter Value Corr Mom Gestational age 13+1 PAPP-A 2.78 mIU/ml 0.48 Method CRL (\$Robinson] hb-hCG 23.4 ng/ml 0.55 Scan date 27/01/2022 Risk at sampling date Nuchal translucency 2.1 Age Risk 1:869 Nuchal translucency MoM 1.15 Biochemical T21 risk 1:8213 Trisomy 13/18+NT <1:10000	Name]	MRS. REKHA	Patient ID		012201280344
Gestational age 13+2 Correction factors Fetuses 1 Weight in kg 61 Diabetes NO Smoker NO Origin Asian Biochemical Data Ultrasound Data Parameter Value Corr Mom Gestational age 13+1 PAPP-A 2.78 mIU/ml 0.48 Method CRL (<>Robinson) Biochemical T21 23.4 ng/ml 0.55 Risk at sampling date Nuchal translucency 2.1 Age Risk 1:869 Nuchal translucency MoM 1.15 Biochemical T21 risk 1:3249 Nasal Bone Presen Combined T21 risk 1:8213 Trisony 13/18+NT <1:10000	Birthday		15/11/1994	Sample ID		11307448
Correction factors Fetuses 1 IVF unknown Previous trisony 21 unknown Weight in kg 61 Diabetes NO Pregnancies unknown Smoker NO Origin Asiau Pregnancies unknown Biochemical Data Ultrasound Data Gestational age 13+1 PAPP-A 2.78 mIU/ml 0.48 Method CRL (\$Robinson] fb-hCG 23.4 mg/ml 0.55 Scan date 27/01/2022 Risks at sampling date Nuchal translucency 2.1 Age Risk 1:869 Nuchal translucency MoM 1.15 Biochemical T21 risk 1:8213 Trisony 13/18+NT <1:10000	Age at delivery		27.2	Sample Date		28/1/2022
Fetuses 1 IVF unknown Previous trisony 21 unknown Weight in kg 61 Diabetes NO Pregnancies unknown Smoker NO Origin Asian Pregnancies unknown Biochemical Data Ultrasound Data Image: Corr Mom Gestational age 13+1 PAPP-A 2.78 mIU/ml 0.48 Method CRL (\$Robinson] Biochemical Data Method CRL (\$Robinson] Scan date 27/01/2022 Risks at sampling date Nuchal translucency 2.1 Nuchal translucency MoM 1.15 Biochemical T21 risk 1:869 Nuchal translucency MoM 1.15 Biochemical T21 risk 1:8213 Trisony 13/18+NT <1:10000	Gestational age		13+2			
Weight in kg 61 Diabetes NO Pregnancies unknown Smoker NO Origin Asian	Correction factors					
Smoker NO Origin Asian Biochemical Data Ultrasound Data Parameter Value Corr Mom Gestational age 13+1 PAPP-A 2.78 mIU/ml 0.48 hchoG 23.4 ng/ml 0.55 Kisks at sampling date Nuchal translucency 2.1 Age Risk 1:869 Nuchal translucency MoM 1.19 Biochemical T21 risk 1:3249 Nasal Bone Presen Combined T21 risk 1:8213 Trisomy 13/18+NT <1:10000	Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Biochemical Data Ultrasound Data Parameter Value Corr Mom Gestational age 13+1 PAPP-A 2.78 mIU/ml 0.48 Method CRL (<>Robinson) fb-hCG 23.4 ng/ml 0.55 Scan date 27/01/2022 Risks at sampling date Nuchal translucency 2.1 Age Risk 1:869 Nuchal translucency MoM 1.15 Biochemical T21 risk 1:3249 Nasal Bone Presen Combined T21 risk 1:8213 Trisomy 13/18+NT <1:10000	Weight in kg	61 Diabetes		NO	Pregnancies	unknown
ParameterValueCorr MomGestational age13+1PAPP-A2.78 mIU/ml0.48MethodCRL (<>Robinson)fb-hCG23.4 ng/ml0.55Scan date27/01/2022Risks at sampling dateNuchal translucency2.1Age Risk1:869Nuchal translucency MoM1.19Biochemical T21 risk1:3249Nasal BonePresenCombined T21 risk1:8213Trisony 13/18+NT<1:10000	Smoker	NO Origin		Asian		
PAPP-A 2.78 mIU/ml 0.48 fb-hCG 23.4 ng/ml 0.55 Risks at sampling date Scan date 27/01/2022 Risks at sampling date Nuchal translucency 2.1 Age Risk 1:869 Nuchal translucency MoM 1.19 Biochemical T21 risk 1:8213 Trisomy 13/18+NT <1:10000	Biochemical Data			Ultrasound Data		
fb-hCG23.4 ng/ml0.55Scan date27/01/2022Risks at sampling dateNuchal translucency2.1Age Risk1:869Nuchal translucency MoM1.19Biochemical T21 risk1:3249Nasal BonePresenCombined T21 risk1:8213Trisomy 13/18+NTTrisomy 13/18+NTDown's Syndrome Risk (Trisomy 21 Screening)RiskDown's Syndrome Risk (Trisomy 21 with NT) is below the cut off, which represents a low risk.After the result of the Trisomy 21 with the same data, there is one woman with a trisomy 21 pregnancy and 8213 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 (with NT)The laboratory cannot be hold responsible for their impact	Parameter	Value	Corr Mom	Gestational age	2	13+1
Risks at sampling dateNuchal translucency2.1Age Risk1:869Nuchal translucency MoM1.19Biochemical T21 risk1:3249Nasal BonePresenCombined T21 risk1:8213Trisomy 13/18+NTTrisomy 13/18+NT<1:10000	PAPP-A	2.78 mIU/ml	0.48	Method		CRL (<>Robinson)
Age Risk1:869Nuchal translucency MoM1.19Biochemical T21 risk1:3249Nasal BonePresenCombined T21 risk1:8213Trisomy 13/18+NT<1:10000	fb-hCG	23.4 ng/ml	0.55	Scan date		27/01/2022
Biochemical T21 risk 1:3249 Combined T21 risk 1:8213 Trisony 13/18+NT <1:10000 Risk Risk Risk 1:700 1:700 1:500 1:1000 Cut off 1:1000 Cut off 1:1000 The calculated risk for Trisony 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisony 21 test it is expected that among 8213 women with the same data, there is one woman with a trisony 21 pregnancy and 8212 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	Risks at sampling date			Nuchal translucency 2.1		
Combined T21 risk1:8213Trisomy 13/18+NT<1:10000	Age Risk	1:869		Nuchal translucency MoM 1.		1.19
Trisony 13/18+NT<1:10000RiskDown's Syndrome Risk (Trisomy 21 Screening)RiskThe calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.1:1000Cut off1:250Cut off1:1000Cut off1:1000First to the result of the Trisomy 21 test it is expected that among 8213 women with the same data, there is one woman with a trisomy 21 pregnancy and 8212 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 (with NT)The laboratory cannot be hold responsible for their impact	Biochemical T21 risk		1:3249	Nasal Bone		Present
RiskDown's Syndrome Risk (Trisomy 21 Screening)RiskThe calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.1:100Cut off1:250Cut off1:250Cut off1:1000First and the same data, there is one woman with a trisomy 21 pregnancy and 8212 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 (with NT)The laboratory cannot be hold responsible for their impact	Combined T21 risk		1:8213			
Trisomy 13/18 (with NT)	Trisomy 13/18+NT		<1:10000			
 1:100 1:250 Cut off 1:250 Cut off 1:1000 1:10000 1:10000	Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
<1:10000, which indicates a low risk values	1:100 1:250 1:100 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1			 cut off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8213 women with the same data, there is one woman with a trisomy 21 pregnancy and 8212 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 		