

*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Risk below Age risk

 Date of Report
 19/3/2022

 PRISCA
 5.1.0.17

Birthday 1/1/1998 Sample ID 66 Age at term 24.08 Sample Date 16/ Gestational age 12+0 Correction factors Fetuses 1 IVF unknown Previous trisomy 21 unwing the result of the trisomy 21 unwing the result of the Trisomy 21 trisomy 21 unwing the result of the Trisomy 21 risk 1:1000 Risk 1:10000 Risk 1:100	Patient Data					
Age at term 24.08 Gestational age 12+0 Correction factors Fetuses 1 IVF unknown Previous trisomy 21 unwhown Pregnancies unwhom Previous trisomy 21 unwhom Previous	Name MRS. SHILPA DOGRA			Patient ID		012203160098
Gestational age Tetuses I IVF unknown Previous trisomy 21	Birthday		1/1/1998	Sample ID		6035984
Trisomy 13/18+NT Correction factors Fetuses 1 IVF unknown Previous trisomy 21 pregnancies Nuchal translucency Nown Nasal Bone Previous trisomy 21 t	Age at term		24.08	Sample Date		16/3/2022
Fetuses 1 IVF unknown Previous trisomy 21 um Weight in kg 79 Diabetes NO Pregnancies um Smoker NO Origin Asian Biochemical Data Parameter Value Corr Mom PAPP-A 3.48 mIU/ml 1.32 Gestational age Method CRL (⇔Rol Scan date 13/ Scan date 13/ Nuchal translucency Nuchal translucency MoM Nasal Bone P Combined trisomy 21 risk 1:1959 Nasal Bone P Combined trisomy 21 risk 1:8502 Trisomy 13/18+NT	Gestational age 12+0					
Weight in kg Top Diabetes NO Origin NO Origin NO Origin NO Pregnancies Ultrasound Data Parameter Value Corr Mom PAPP-A 3.48 mIU/ml 1.32 Risks at sampling date Nuchal translucency Nuchal translucency Nuchal translucency Mom Nasal Bone Parameter Nuchal translucency Nombined trisomy 21 risk 1:8502 Trisomy 13/18+NT The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate information provided by the referring physician. Ple note that the risk calculations are statistical aapproaches 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 im on the risk assessment! Calculated risks have no diagnostic value!	Correction factors					
Biochemical Data Parameter Value Corr Mom PAPP-A 3.48 mIU/ml 1.32 Risks at sampling date Age Risk 1:974 Biochemical T21 risk 1:1959 Combined trisomy 21 risk 1:18502 Trisomy 13/18+NT Cat off 1:1000 Risk Black 1:1000 Risk Down's Syndrome Risk (Trisomy 21 screening) The calculated risk for Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate information provided by the referring physician. Ple note that the risk calculations are statistical aapproaches 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 im on the risk assessment! Calculated risks have no diagnoon the risk assessment!	Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Parameter Value Corr Mom PAPP-A 3.48 mIU/ml 1.32 Method CRL (◇Rol Ib-hCG 92.6 ng/ml 2.03 Scan date 13/ Risks at sampling date Age Risk 1:974 Nuchal translucency Nuchal translucency MoM Biochemical T21 risk 1:1959 Nasal Bone P Combined trisomy 21 risk 1:8502 Trisomy 13/18+NT < <1:10000 Risk Risk 1:10 1:100 1:150 Cut off The calculated risk for Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate the information provided by the referring physician. Ple note that the risk calculations are statistical aapproaches 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 impont the risk assessment! Calculated risks have no diagnotic contents are the information provided by the referring physician. Ple note that the risk calculations are statistical aapproaches have no diagnostic value!	Weight in kg	79 Diabetes		NO	Pregnancies	unknown
Parameter Value Corr Mom PAPP-A 3.48 mIU/ml 1.32 Method CRL (≪Rol fb-hCG 92.6 ng/ml 2.03 Scan date 13/ Risks at sampling date Age Risk 1:974 Nuchal translucency Nuchal translucency MoM Nasal Bone P Combined trisomy 21 risk 1:8502 Trisomy 13/18+NT <1:1000 Risk Risk 1:100 1:	Smoker	NO Origin		Asian		
PAPP-A 3.48 mIU/ml 1.32 Method CRL (SRol pb-hCG 92.6 ng/ml 2.03 Scan date 13/ Risks at sampling date Age Risk 1:974 Nuchal translucency Nuchal translucency MoM Nasal Bone P Combined trisomy 21 risk 1:8502 Trisomy 13/18+NT <1:10000 Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate the information provided by the referring physician. Ple note that the risk calculations are statistical aapproaches have no diagnostic value! Trisomy 13/18+NT The calculated risk for Trisomy 13/18 with NT is on the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk assessment! Calculated risks have no diagnostic valued to the risk	Biochemical Data			Ultrasound Data		
Risks at sampling date Age Risk 1:974 Biochemical T21 risk 1:1959 Combined trisomy 21 risk 1:8502 Trisomy 13/18+NT Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurace the information provided by the referring physician. Ple note that the risk calculations are statistical aapproaches 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 im on the risk assessment! Calculated risks have no diagnotic risks have no diagnotic risks assessment! Calculated risks have no diagnotic risks have no diagno	Parameter	Value	Corr Mom	Gestational age	2	11+4
Risks at sampling date Age Risk 1:974 Nuchal translucency Nuchal translucency Nuchal translucency MoM Nasal Bone P Combined trisomy 21 risk 1:8502 Trisomy 13/18+NT Cut off 1:100 1:100 Cut off Trisomy 13/18+NT Cut off Trisomy 13/18+NT The calculated risk for Trisomy 21 with NT is Nuchal translucency Person 13/18+NT The calculated risk for Trisomy 21 with NT is expected that among 8502 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate the information provided by the referring physician. Plenote that the risk calculations are statistical aapproaches have no diagnostic value! The calculated risk for Trisomy 21 wit	PAPP-A	$3.48~\mathrm{mIU/ml}$	1.32	Method		CRL (<>Robinson)
Age Risk Biochemical T21 risk 1:1959 Combined trisomy 21 risk 1:8502 Trisomy 13/18+NT Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate information provided by the referring physician. Plenote that the risk calculations are statistical aapproaches have no diagnostic value! Trisomy 13/18+NT The calculated risk for Trisomy 13/18 with NT is Nuchal translucency MoM Nasal Bone P Nasal Bone P The calculated risk (Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate the information provided by the referring physician. Plenote that the risk calculations are statistical aapproaches have no diagnostic value! Trisomy 13/18+NT The laboratory cannot be hold responsible for their importance of the provided by the referring physician. Plenote that the risk calculations are statistical aapproaches have no diagnostic value!	fb-hCG	92.6 ng/ml	2.03	Scan date		13/3/2022
Biochemical T21 risk 1:1959 Combined trisomy 21 risk 1:8502 Trisomy 13/18+NT Slisk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate information provided by the referring physician. Ple note that the risk calculations are statistical aapproaches 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 importance on the risk assessment! Calculated risks have no diagnostic valued.	Risks at sampling date			Nuchal translucency 1.2		
Combined trisomy 21 risk Trisomy 13/18+NT 1:10000 Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate the information provided by the referring physician. Plenote that the risk calculations are statistical aapproaches 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 impontance on the risk assessment! Calculated risks have no diagnostic valued.	Age Risk 1:974		Nuchal translucency MoM 0.94			
Risk Down's Syndrome Risk (Trisomy 21 Screening)	Biochemical T21 risk 1		1:1959	Nasal Bone Preso		Presesnt
Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate the information provided by the referring physician. Plenote that the risk calculations are statistical aapproaches have no diagnostic value! Trisomy 13/18+NT	Combined trisomy 21 risk 1:8502					
The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate the information provided by the referring physician. Plenote that the risk calculations are statistical aapproaches 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 importance on the risk assessment! Calculated risks have no diagnostic valued the risk assessment on the risk assessment.	Trisomy 13/18+NT		<1:10000			
The calculated risk for Trisomy 21 with NT is below off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 women with not affected pregnancies. The calculated risk by PRISCA depends on the accurate the information provided by the referring physician. Ple note that the risk calculations are statistical aapproaches 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 importance on the risk assessment! Calculated risks have no diagnostic value.				Down's Syndrome Risk (Trisomy 21 Screening)		
	1:1000 1:1000 1:10000 13 15 17 19 21 23 25 2 Trisomy 13/18+NT The calculated risk for 7	Trisomy 13/18 with	41 43 45 47 49 Age	off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 8502 women with the same data, there is one woman with a trisomy 21 pregnancy 8501 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		

Risk above Age Risk

Risk Above Cut Off