

 Date of Report
 9/12/2022

 PRISCA
 5.1.0.17

Birthday 29/05/1989 Sample ID 113608 Age at term 33.11 Sample Date 8/12/20  Correction factors  Fetuses 1 IVF unknown Previous trisomy 21 unknown Weight in kg 54 Diabetes NO Pregnancies unknown Smoker NO Origin Asian  Biochemical Data  Parameter Value Corr Mom Gestational age 12/20  PAPP-A 4.9 mIU/ml 0.88 Method CRL (◇Robinse 16/12/20  Risks at sampling date Nuchal translucency 1.  Age Risk 1:373 Nuchal translucency MoM Nasal bone Press  Combined trisomy 21 risk 4:10000  Trisomy 13/18 + NT 4:10000  Risk  Risk 1:000  Risk 1:000  Risk 1:000  Risk 1:000  Risk 1:000  Trisomy 13/18 + NT 4:10000  Risk 1:000  Trisomy 13/18 + NT 5:10000  Risk 1:000  Trisomy 13/18 + NT 5:10000  Trisomy 13/18 + NT 5:10000  Risk 1:000  Trisomy 13/18 + NT 5:10000	Patient Data					
Age at term 33.11 Sample Date 8/12/20 Gestational age 12+6  Correction factors  Fetuses 1 IVF unknown Previous trisomy 21 unknown Smoker NO Origin Asian  Biochemical Data  Parameter Value Corr Mom PAPP-A 4.9 mIU/ml 0.88 Ib-hCG 26.3 mg/ml 0.79  Risks at sampling date  Age Risk 1:373 Nuchal translucency Nuchal translucency MoM Nasal bone Press  Combined trisomy 21 risk 1:10000  Trisomy 13/18 + NT	Name MRS. SANANDA DAS			Patient ID		012212080124
Gestational age  Tetuses  I IVF  Unknown  Previous trisony 21  Unkno  Weight in kg  54  Diabetes  NO  Origin  Asian  Biochemical Data  Parameter  Value  Corr Mom  PAPP-A  4.9 mIU/ml  0.88  Bi-bCG  26.3 ng/ml  0.79  Scan date  Nuchal translucency  Trisomy 13/18 + NT  1:1000  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 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low misk.  After the result of	Birthday		29/05/1989	Sample ID		11360895
Trisomy 13/18+NT    IVF	Age at term		33.11	Sample Date		8/12/2022
Fetuses 1 IVF unknown Previous trisomy 21 unknown Weight in kg 54 Diabetes NO Origin Asian  Biochemical Data    Parameter   Value   Corr Mom	Gestational age		12+6			
Weight in kg Smoker NO Origin    NO   Origin   Asian	Correction factors					
Biochemical Data    Parameter   Value   Corr Mom   Gestational age   12   PAPP-A   4.9 mIU/ml   0.88   Method   CRL (⋄ Robinse fb-hCG   26.3 ng/ml   0.79   Scan date   6/12/20   Risks at sampling date   Nuchal translucency   Nuchal transluce	Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Parameter Value Corr Mom  PAPP-A 4.9 mIU/ml 0.88  Bio-hCG 26.3 ng/ml 0.79  Risks at sampling date  Age Risk 1:373  Bio-chemical T21 risk 1:2901  Combined trisomy 21 risk 1:10000  Risk  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 result o	Weight in kg	54 Diabetes		NO	Pregnancies	unknown
Parameter Value Corr Mom PAPP-A 4.9 mIU/ml 0.88  fb-hCG 26.3 ng/ml 0.79  Risks at sampling date  Age Risk 1:373  Biochemical T21 risk 1:2901  Combined trisomy 21 risk 1:10000  Risk  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 with NT test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 preparately 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 laboratory cannot be hold responsible for their impact	Smoker	NO Origin		Asian		
PAPP-A 4.9 mIU/ml 0.88 Method CRL (Sobinsoft-hCG 26.3 ng/ml 0.79 Scan date 6/12/20  Risks at sampling date Nuchal translucency 1.  Age Risk 1:373 Nuchal translucency MoM Nasal bone Press  Combined trisomy 21 risk 1:2901 Nasal bone Press  Combined trisomy 21 risk 1:10000  Risk 1:10000  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 with NT test 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 laboratory cannot be hold responsible for their impact	Biochemical Data			Ultrasound Data		
Risks at sampling date  Age Risk  Biochemical T21 risk  Combined trisomy 21 risk  Trisomy 13/18 + NT  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 with NT test 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 laboratory cannot be hold responsible for their impact	Parameter	Value	Corr Mom	Gestational age	2	12+4
Risk 1:373  Biochemical T21 risk 1:2901  Combined trisomy 21 risk 4:10000  Trisomy 13/18 + NT	PAPP-A	$4.9~\mathrm{mIU/ml}$	0.88	Method		CRL (<>Robinson)
Age Risk  1:373  Nuchal translucency MoM  Nasal bone  Press  Combined trisomy 21 risk  1:10000  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 with NT test 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 laboratory cannot be hold responsible for their impact	fb-hCG	26.3 ng/ml	0.79	Scan date		6/12/2022
Biochemical T21 risk  Combined trisomy 21 risk  Trisomy 13/18 + NT    Syndrome Risk (Trisomy 21 Screening)	Risks at sampling date	e e		Nuchal translucency 1.12		
Combined trisomy 21 risk  Trisomy 13/18 + NT  S1:10000  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 21with NT test 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 laboratory cannot be hold responsible for their impact	Age Risk		1:373	Nuchal translucency MoM 0.3		
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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 21with NT test 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 laboratory cannot be hold responsible for their impact	Combined trisomy 21 risk <1:100					
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 with NT test 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 laboratory cannot be hold responsible for their impact	Trisomy 13/18 + NT <1:10000					
cut off, which represents a low risk.  After the result of the Trisomy 21 with NT test 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 laboratory cannot be hold responsible for their impact	Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
<1:10000, which indicates a low risk  values	1:1000 1:250 1:10000 13 15 17 19 21 23 2 Trisomy 13/18+NT The calculated risk for	or Trisomy 13/18 (with	41 43 45 47 49 Age	After the result of the Trisomy 21with NT test 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		