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 Date of Report
 27/9/2022

 PRISCA
 5.1.0.17

Birthday  Age at term  30.11 Gestational age  13+6  Correction factors  Fetuses  1 IVF unknown Weight in kg 53 Diabetes NO Origin  Biochemical Data  Parameter  Value  Corr Mom PAPP-A  5.12 mIU/ml 0.51 Biochemical T21 risk 1:636  Trisomy 13/18+NT  The calculated risk for Trisomy 13/18 (with NT) is  Biochemical T21 risk tis expected that among 636 women with not affected pregnancies.  Data 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 on the risk assessment! Calculated risk have no diagnostic value!  Sample ID  1124590  Sample ID  1124590  Sample ID  1124590  Sample Date 26/9/202  Unknown Previous trisomy 21 unknown Previous trisomy 21 unknow Pregnancies  Unknown Previous trisomy 21 unknown Previous trisomy 21 unknow Previous trisomy 21 unknow Pregnancies  Unknown Previous trisomy 21 unknow Previous trisomy 21 unknown Prev	Patient Data					
Age at term 30.11   Sample Date 26/9/202   Correction factors   Fetuses 1   IVF   unknown   Previous trisomy 21   unknown   Pregnancies   unknown   Previous trisomy 21   unknown   Pregnancies   unknown   Previous trisomy 21   unknown   Pregnancies   unknown   Previous trisomy 21   unknown   Previous	Name	I	MRS. PREETI	Patient ID		012209260150
Gestational age 13+6  Correction factors  Fetuses 1 IVF unknown Weight in kg 53 Diabetes NO Origin Asian  Biochemical Data  Parameter Value Corr Mom Gestational age 13+6  PAPP-A 5.12 mIU/ml 0.51 Method CRL (≪Robinson Ib-ltCG 62.5 ng/ml 2.42 Scan date 26/9/202  Risks at sampling date Nuchal translucency 1.  Age Risk 1:635 Nuchal translucency MoM 0.  Biochemical T21 risk 1:96 Nasal bone Present Combined trisomy 21 risk 1:636  Trisomy 13/18 + NT < 1:10000  Risk 1:00	Birthday		2/4/1992	Sample ID		11245905
Fetuses 1 IVF unknown Previous trisomy 21 unknow Smoker NO Origin NO Origin Asian  Biochemical Data  Parameter Value Corr Mom Gestational age 13+ PAPP-A 5.12 mIU/ml 0.51 Method CRL (≪Robinsor Scan date 26/9/202  Risks at sampling date Nuchal translucency 1.  Age Risk 1:635 Nuchal translucency MoM 0.  Biochemical T21 risk 1:96 Nasal bone Present Combined trisomy 21 risk 1:636  Trisomy 13/18 + NT < 1:1000  1:1000  Trisomy 13/18+NT	Age at term		30.11	Sample Date		26/9/2022
Fetuses 1 IVF unknown Previous trisomy 21 unknown Weight in kg 53 Diabetes NO Origin Asian Biochemical Data  Parameter Value Corr Mom PAPP-A 5.12 mIU/ml 0.51 Method CRL (≪Robinsor Rb-hCG 62.5 ng/ml 2.42 Scan date 26/9/202 Risks at sampling date Nuchal translucency Mom 0. Risk 1:635 Nuchal translucency Mom 0. Nasal bone Present Combined trisomy 21 risk 1:636 Trisomy 13/18 + NT <a href="https://linearchy.com/state/documents/">1:636 Trisomy 13/18 + NT State/documents/</a> Down'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 NT test it is expected that among 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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 18/18 (with NT) is  The calculated risk sasessment! Calculated risks have no diagnostic on the risk assessment! Calculated risks have no diagnostic on the risk assessment! Calculated risks have no diagnostic on the risk assessment! Calculated risks have no diagnostic on the risk assessment! Calculated risks have no diagnostic on the risk assessment!	Gestational age		13+6			
Weight in kg Smoker NO Origin    Diabetes   NO   Origin   Asian	Correction factors					
Biochemical Data  Parameter Value Corr Mom PAPP-A 5.12 mIU/ml 0.51 Method CRL (≪Robinsor Scan date 26/9/202 Risks at sampling date  Nuchal translucency 1.  Age Risk 1:635 Nuchal translucency MoM 0.  Biochemical T21 risk 1:96 Combined trisomy 21 risk 1:636 Trisomy 13/18 + NT < 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 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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	Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Parameter Value Corr Mom PAPP-A 5.12 mIU/ml 0.51 Biochemical Total translucency Risks at sampling date  Age Risk 1:635 Biochemical T21 risk 1:96 Combined trisomy 21 risk 1:636 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 test it is expected that among 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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 18/18+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	Weight in kg	53 Diabetes		NO	Pregnancies	unknown
PAPP-A 5.12 mIU/ml 0.51 Method CRL (\$Robinsor CRL (	Smoker	NO Origin		Asian		
PAPP-A 5.12 mIU/ml 0.51   th-hCG 62.5 ng/ml 2.42   Scan date 26/9/202   Risks at sampling date	Biochemical Data			Ultrasound Data		
Risks at sampling date  Age Risk  1:635  Biochemical T21 risk  1:96  Combined trisomy 21 risk  1:636  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 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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	Parameter	Value	Corr Mom	Gestational age	2	13+6
Risks at sampling date  Nuchal translucency  Nuchal translucency MoM  O.  Biochemical T21 risk  1:96  Nasal bone  Preser  Combined trisomy 21 risk  1:636  Trisomy 13/18 + NT  Cut off  1:100  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 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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	PAPP-A	$5.12~\mathrm{mIU/ml}$	0.51	Method		CRL (<>Robinson)
Age Risk  Biochemical T21 risk  1:96  Combined trisomy 21 risk  1:636  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 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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	62.5 ng/ml	2.42	Scan date		26/9/2022
Biochemical T21 risk  Combined trisomy 21 risk  1:636  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 21with NT test it is expected that among 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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	Risks at sampling date	;		Nuchal translucency 1.5		
Combined trisomy 21 risk  Trisomy 13/18 + NT  Slisk  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 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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 on the risk assessment! Calculated risks have no diagnostic	Age Risk		1:635	Nuchal translucency MoM 0.		0.8
Trisomy 13/18 + NT    Some state of the provided by the referring physician. Please note that the risk calculated risk for 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 with NT test it is expected that among 636 women with a trisomy 21 pregnancy and 635 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	Biochemical T21 risk		1:96	Nasal bone Prese		Present
Risk  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 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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 on the risk assessment! Calculated risks have no diagnostic	Combined trisomy 21 risk 1:636					
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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 with NT test it is expected that among 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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	Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
<1:10000, which indicates a low risk values	1:10 1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 2: Trisomy 13/18+NT The calculated risk for	r Trisomy 13/18 (with	41 43 45 47 49 Age	cut off, which represents a low risk.  After the result of the Trisomy 21 with NT test it is expected that among 636 women with the same data, there is one woman with a trisomy 21 pregnancy and 635 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!		