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Birthday $29/05/1988$ Sample ID $1114758$ Age at delivery $34.1$ Sample Date $26/12/202$ Gestational age $12+5$ $26/12/202$ Correction factorsFetuses1IVFunknownPrevious trisomy 21unknownWeight in kg $60.2$ DiabetesNOPregnanciesunknownSmokerNOOriginAsianPregnanciesunknownBiochemical DataUltrasound DataParameterValueCorr MomGestational age124PAPP-A $7.04$ mIU/ml $1.51$ MethodCRL(<>Robinsonfb-hCG $29.2$ mg/ml $0.7$ Scan date $23/12/202$ Risks at sampling dateNuchal translucencyNuchal translucency $0.5$ Age Risk $1:513$ Nuchal translucency MoM $0.5$					Date of Report PRISCA	27/12/2021 5.2.0.13
Birthday 29/05/1988 Sample ID 1114758 Age at delivery 34.1 Sample Date 26/12/202 Cestational age 12*5 Correction factors Fetuses 1 IVF unknown Previous trisony 21 unknown Weight in kg 60.2 Diabetes NO Pregnancies unknow Smoker NO Origin Asia Biochemical Data Ultrasound Data Parameter Value Corr Mom PAPP-A 7.04 mIU/ml 1.51 Biochemical 721 risk 1.513 Biochemical T21 risk 4.1:0000 Age Risk 1:513 Nuchal translucency 0.0 Age Risk 4.1:10000 Trisony 13/18+NT 4.1:0000 Trisony 13/18+NT 4.1:0000 Risk Combined T21 risk 4.1:0000 Risk 4.1:0000 Risk 4.1:0000 Risk 4.1:0000 Risk 4.1:0000 Risk 7.1:0000 Risk 7.1:00	Patient Data					
Age at delivery       34.1       Sample Date       26/12/202         Gestational age       122-5       122-5       122-5         Correction factors         Fetuses       1       IVF       unknown       Previous trisony 21       unknown         Smoker       NO       Origin       Asian       NO       Pregnancies       unknown         Biochemical Data       Ultrasound Data       Corr Mom       Gestational age       124         PAPP-A       7.04 mIU/ml       1.51       Method       CRL( <robinso< td="">         fb-hCG       29.2 ng/ml       0.7       Scan date       23/12/202         Risks at sampling date       Nuchal translucency       0.       Age         Nichal translucency       0.       Nasal bone       Presention         Combined T21 risk       &lt;1:10000</robinso<>	Name	MR	S. SWEKSHA	Patient ID		12112260051
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Correction factors         Fetuses       1       IVF       unknown       Previous trisomy 21       unknow         Weight in kg       60.2       Diabetes       NO       Pregnancies       unknow         Smoker       NO       Origin       Asian       Intersection factors       unknow         Biochemical Data       Ultrasound Data       Intersection factors       Intersection factors       Intersection factors         PAPP.A       7.04 mIU/ml       1.51       Method       CRL( <robinso< th="">       CRL(<robinso< th="">         Ib-hCG       29.2 ng/ml       0.7       Scan date       23/12/202         Risk as asampling date       Nuchal translucency       0.0         Age Risk       1:513       Nuchal translucency MoM       0.3         Biochemical T21 risk       &lt;1:10000</robinso<></robinso<>	Age at delivery		34.1	Sample Date		26/12/2021
Fetuses       1       IVF       unknown       Previous trisomy 21       unknown         Weight in kg       60.2       Diabetes       NO       Pregnancies       unknown         Smoker       NO       Origin       Asian       Pregnancies       unknown         Biochemical Data       Ultrasound Data       Intersection       Intersection <t< td=""><td>Gestational age</td><td></td><td>12+5</td><td></td><td></td><td></td></t<>	Gestational age		12+5			
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Smoker       NO       Origin       Asian         Biochemical Data       Ultrasound Data         Parameter       Value       Corr Mom         PAPP-A       7.04 mIU/ml       1.51         Bio-hCG       29.2 ng/ml       0.7         Stask at sampling date       Nuchal translucency       0.         Age Risk       1:513       Nuchal translucency MoM       0.5         Biochemical T21 risk       <1:10000	Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
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Age Risk1:513Nuchal translucency MoM0.5Biochemical T21 risk<1:10000	fb-hCG	29.2 ng/ml	0.7	Scan date		23/12/2021
Biochemical T21 risk       <1:10000	Risks at sampling date			Nuchal translucency 0.8		
Combined T21 risk<1:10000Trisomy 13/18+NT<1:10000RiskDown'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 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 211:1000Cut off1:1000Cut off1:1000Cut off1:1000Cut off1:1000Age	Age Risk	isk 1:513		Nuchal translucency MoM 0.5		0.52
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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 21 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!	Combined T21 risk		<1:10000			
Risk 1:10 1:100 1:250 Cut off 1:100 1:250 Cut off 1:100 1:250 Cut off 1:100 1:250 Cut off 1:100 1:250 Cut off 1:100 1:250 Cut off 1:250 Cut off Age	Trisomy 13/18+NT		<1:10000			
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 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!	Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
The calculated risk for Trisomy 18/18 (with NT) is       on the risk assessment! Calculated risks have no diagnostic         <1:10000, which indicates a low risk	1:100 1:250 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000			<ul> <li>cut off, which represents a low risk.</li> <li>After the result of the Trisomy 21 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!</li> <li>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic</li> </ul>		