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Birthday $16/11/1998$ Sample ID 1166835 Age at term 24.1 Sample Date $21/5/202$ Gestational age $12+5$ Sample Date $21/5/202$ Correction factorsFetuses1IVFunknownPrevious trisomy 21unknownWeight in kg 59 DiabetesNOPregnanciesunknownSmokerNOOriginAsianPregnanciesunknownBiochemical DataUltrasound DataUltrasound Data12+ParameterValueCorr MomMethodCRL (<>Robinson $lb-hCG$ 17.4 ng/ml 0.49 Scan date $21/5/202$ Risks at sampling date1:987Nuchal transluency1.Age Risk $1:987$ Nuchal transluency MoM 0.7					Date of Report PRISCA	23/5/2023 5.2.0.13
Birthday 16/11/1998 Age at term 21:1 Sample Date 21/5/202 Gestational age 12:5 Correction factors Fetuses 1 IVF unknown Previous trisony 21 unknow Weight in kg 59 Diabetes NO Smoker NO Origin Asian Biochemical Data Ultrasound Data Parameter Value Corr Mom PAPP-A 2.82 mIU/ml 0.47 BibCG 17.4 ng/ml 0.49 Risks at sampling date Nuchal translucency 1. Age Risk 1:987 Biochemical T21 risk 1:4412 Combined trisomy 21 risk 4:1:4412 Combined trisomy 21 risk 4:1:4000 Trisomy 13/18 + NT 4:10000 Risk The calculated risk for Trisomy 18/18 (with NT) is below the cut of the Trisomy 21 with NT test it is expected that among more than 1000 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please on the risk calculated risk have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment Calculated risk have no diagnostic value!	Patient Data					
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Biochemical T21 risk 1:4412 Nasal bone Preser Combined trisomy 21 risk <1:10000	Risks at sampling date			Nuchal translucency 1.2		
Combined trisomy 21 risk <1:10000	Age Risk 1:987		1:987	Nuchal translucency MoM 0.74		
Trisomy 13/18 + NT <1:10000	Biochemical T21 risk		1:4412	Nasal bone		Present
Risk Risk 1:10 1:1000 1:1000	Combined trisomy 21 risl	k	<1:10000			
Risk 1:10 1:10 The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. 1:100 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. 1:1000 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 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	Trisomy 13/18 + NT		<1:10000			
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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	1:10 1:10 1:250 Cut off 1:1000 1:1			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		
Risk Above Cut Off Risk above Age Risk Risk below Age risk	<1:10000 , which indicat	es a low risk		on the risk assessment! Calculated risks have no diagnostic		