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					Date of Report PRISCA	31/5/2022 5.1.0.17
Patient Data						
Name			MRS. JUL	I Patient ID		012205290038
Birthday			12/6/199	8 Sample ID		11242320
Age at term			24.0	6 Sample Date		29/5/2022
Gestational age			12+	6		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy	21 unknown
Weight in kg	52.5	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	12+5
PAPP-A	5.29	mIU/ml	0.89	Method		CRL (<>Robinson)
fb-hCG	54.7	ng/ml	1.16	Scan date		28/05/2022
Risks at sampling date				Nuchal Translucency 1.5		
Age Risk			1:1012	Nuchal Transl	ucency MoM	0.9
Biochemical T21 risk		1:3486	Nasal Bone		Present	
Combined T21 risk			<1:10000			
Trisomy 13/18+NT			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:100 1:250 1:1000 1:250 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000 1:1000 1:1000 1:250 Cut off 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 test (with NT) 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 calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk				on the risk assessment! Calculated risks have no diagnostic values		
Risk Above Cut Off				Risk above Ag	e Risk	Risk below Age risk