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Sample Collection				
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				Date of Report PRISCA	16/2/2022 5.2.0.13
Patient Data					
Name	MR	MRS. PARWATI			012202150111
Birthday		1/1/1997	Sample ID		11375726
Age at delivery		25.1	Sample Date		15/02.2022
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+6
PAPP-A	4.5 mIU/ml	0.89	Method		CRL(<>Robinson
fb-hCG	58.3 ng/ml	1.3	Scan date		15/2/2022
Risks at sampling date			Nuchal translucency 1.5		
Age Risk		1:967	Nuchal translucency MoM		0.9
Biochemical T21 risk		1:2549	Nasal Bone		Present
Combined T21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21 S	Screening)
Risk 1:10 1:250 Cut off 1:1000			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 approaches and		
Trisomy 13/18 + NT The calculated risk for <1:10000, which indica		Age	have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values Risk above Age Risk Risk below Age risk		