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Sample Collection
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				Date of Report PRISCA	2/3/2022 5.2.0.13
Patient Data					
Name	MRS. NISH	IA KANWAR	Patient ID		012203010132
Birthday		29/8/1991	Sample ID		11307232
Age at term		30.5	Sample Date		1/3/2022
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	55 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+0
PAPP-A	3.29 mIU/ml	0.51	Method		CRL (<>Robinson)
fb-hCG	26.7 ng/ml	0.61	Scan date		27/2/2022
Risks at sampling date			Nuchal translucency 1.46		
Age Risk		1:622	Nuchal translucency MoM		0.86
Biochemical T21 risk		1:2173	Nasal Bone		Present
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10			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! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
	ates a low risk sk Above Cut Off		values Risk above Ag	e Risk 🛛 📕 B	lisk below Age risk