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Sample Collection			
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<b>Patient Data</b> Name Birthday Age at delivery					
Birthday					
-		MKS. POOJA	Patient ID		012201310431
Age at delivery		10/3/2001	Sample ID		11178228
		20.9	Sample Date		31/1/2022
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	42 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data		Ultrasound Da	ata		
Parameter	Value	Corr Mom	Gestational age	2	12+4
PAPP-A	5.67 mIU/ml	0.78	Method		CRL (<>Robinson)
fb-hCG	78.2 ng/ml	1.5	Scan date		31/01/2022
Risks at sampling date			Nuchal translucency 1.3		
Age Risk		1:1082	Nuchal translucency MoM		0.83
Biochemical T21 risk		1:1509	Nasal Bone		Present
Combined T21 risk		1:8311			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 1:250 Cut off 1:1000 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 Trisomy 13/18 (with NT) The calculated risk for Trisomy 13/18 with NT is <1:1000, which indicates a low risk			<ul> <li>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</li> <li>After the result of the Trisomy 21 test it is expected that among 8311 women with the same data, there is one woman with a trisomy 21 pregnancy and 8310 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 values</li> </ul>		