



				Date of Report PRISCA	1/1/2019 5.0.2.37
Patient Data				ТКІЗСА	3.0.2.37
Name	Mrs	SEEMA-22708	Patient ID		011812300131
Birthday		5/12/1987	Sample ID		10348229
Age at delivery		31.1	Sample Date		30/12/18
Gestational age		13+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknow
Weight in kg	55 Diabetes		unknown	Pregnancies	unknow
Smoker	Unknown Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	5	13+1
PAPP-A	5.16 mIU/ml	0.89	Method		CRL (⇔Robinson)
fb-hCG	32.9 ng/ml	0.79	Scan Date		29/12/2018
Risks at sampling date			Crown rump length (mm)		70.4
Age Risk		1:577	Nuchal translucency MoM		0.65
Biochemical T21 Risk		1:4640	Nasal Bone		Present
Combined Trisomy 21 Risk		<1:10000	Sonographer		DR. NAVNEET
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS,DMRD
Risk			Down's Syndro	ome Risk (Trisomy S	21 Screening)
Risk 1:10 1:100 1:450 1:1000 1:1500 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1			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 4640 women with the same data, there is one woman with a trisomy 21 pregnancy. 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		