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					Date of Report PRISCA	09-03-19 5.0.2.37
Patient Data						0.0.2.07
Name	Μ	Irs.GEET	A YADAV	Patient ID		011903080151
Birthday			07-11-89	Sample ID		10396431
Age at delivery			29.3	Sample Date		08/03/19
Gestational age by			13+2			
Correction factors					1	
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	66	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	13+2
PAPP-A	5.41	mIU/ml	1.21	Method		CRL (<>Hadlock)
fb-hCG	32.2	ng/ml	0.81	Scan Date		08-03-19
				CRL measurm	ents	71
Risks at sampling date				Nuchal translucency MoM 0.62		
Age Risk			1:717	Nasal bone		Present
Biochemical T21 Risk			<1:10000	Sonographer		LPRAKASH LALCHANDANI
Combined Trisomy 21 Risk			<1:10000	Qualifications	in measuring NT	MD
Trisomy 13/18			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10				The calculated risk for Trisomy 21 (with nuchal		
1:100				translucency) is below the cut off, which indicates a low risk.		
				After the result of the Trisomy 21 test (with NT) it is expected that among 10000 women with the same data,		
1:250		/	Cut off		1	
				The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please		
1:1000				note that risk calculations are statistical approach and have no		
1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				diagnostic values! The patient combined risk presumes the NT		
				measurement was done according to accepted guidelines		
Trisomy 13/18 + NT		Age		-		
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.						
Risk /	Above C	Cut Off		Risk above Ag	ge Risk	Risk below Age risk