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				Date of Report PRISCA	24/08/19 5.0.2.37
Patient Data				THOCH	0.0.2.07
Name		Mrs Sweety	Patient ID		011908230044
Birthday		2/8/1994	Sample ID		10598016
Age at delivery		25.1	Sample Date		23/08/19
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	45 Diabetes		no	Pregnancies	
Smoker	no Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+2
PAPP-A	4.61 mIU/ml	0.56	Method		CRL (<>Robinson)
fb-hCG	33.05 ng/ml	0.70	Scan Date		23/08/2019
Risks at sampling date			Crown Rump Length (mm) 71.5		
Age Risk		1:984	Nuchal translu	Nuchal translucency MoM	
Biochemical Trisomy 21 Risk		1:3436	Nasal Bone	Nasal Bone pres	
Combined Trisomy 21 Risk		<1:10000	Sonographer	r DR.RUBY	
Trisomy 13/18 + NT		<1:10000	Qualification in	ation in measuring NT CON. RADIOLO	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.		
1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 Trisomy 13/18 + NT			After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 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		
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			on the risk assessment! Calculated risks have no diagnostic values		
Ris	sk Above Cut Off		Risk above Ag	e Risk	Risk below Age risk