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				Date of Report PRISCA	1/9/2019 5.0.2.37
Patient Data					
Name		Mrs Jyoti	Patient ID		011908310111
Birthday		11/3/1992	Sample ID		10607832
Age at delivery		<b>27.</b> 5	Sample Date		31/08/19
Gestational age		11+4	9		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	65 Diabete	S	no	Pregnancies	
Smoker	no Origin		Asian		
<b>Biochemical Data</b>			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	11+4
PAPP-A	3.6 mIU/m	l 1.29	Method		CRL (<>Robinson)
fb-hCG	21.5 ng/ml	0.42	Scan Date		31/08/19
Risks at sampling date			Crown Rump Length (mm) 47		
Age Risk 1:801		1:801	Nuchal translucency MoM 0.86		
Biochemical Trisomy 21 Risk		<1:10000	Nasal Bone p		present
Combined Trisomy 21 Risk		<1:10000	Sonographer Dr. Prakash La		Dr. Prakash Lalchandani
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MD
Risk			Down's Syndro	ome Risk (Trisomy	21 Screening)
Risk 1:10 1:250 Cut off 1:250 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:1000 (which represents a low risk.			5		