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						Date of Report	04/03/2020
Patient Data						PRISCA	5.0.2.37
Name Mrs Bhuvaneswari					Patient ID		062003010045
				Sample ID		10513185	
Age at delivery			,		Sample Date		01/03/2020
Gestational age 13+6				-			
Correction factors							
Fetuses	1	IVF			unknown	Previous trisomy 21	unknown
Weight in kg	46	Diabetes			no	Pregnancies	
Smoker	no	Origin			Asian		
Biochemical Data				Ultrasound Data			
Parameter	Value		Corr M	lom	Gestational age		12+5
PAPP-A	6.1	mIU/ml	0.6	64	Method		CRL (<>Robinson)
fb-hCG	42.1	ng/ml	0.9	96	Scan Date		24/02/2020
Risks at sampling date					Crown Rump Length (mm) 66		
Age Risk			1:1082		Nuchal translu	cency MoM	1.01
Biochemical Trisomy 21 Risk			1:2636		Nasal Bone		Present
Combined Trisomy 21 Risk			<1:1000	00	Sonographer		DR. A DHANADIA
Trisomy 13/18 + NT			<1:1000	00	Qualification in measuring NT		MD
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: 00 1:250 Out off 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 Age				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!			
Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.					The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		



Risk Above Cut Off



