

*Free Home Sample Collection 9999 778 778



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Date of Report	07-03-19
PRISCA	5.0.2.37

				PF	RISCA	5.0.2.37
Patient Data						
Name		M	Irs Renuka	Patient ID		011903050245
Birthday			20-07-91	Sample ID		10436342
Age at delivery			28.6	Sample Date		05/03/19
Gestational age by			13+4			
Correction factors						
Fetuses	1 IV	/F		unknown Pr	revious trisomy 21	unknown
Weight in kg	63 D	iabetes		no Pr	regnancies	
Smoker	no O	rigin		Asian		
Biochemical Data				Ultrasound Data	1	
Parameter	Value		Corr Mom	Gestational age		13+3
PAPP-A	6.12 m	IU/ml	0.86	Method		CRL (⇔Hadlock)
fb-hCG	18.75 ng	g/ml	0.44	Scan Date		05-03-19
				CRL measurment	ts	72.4
Risks at sampling date			Nuchal translucency MoM 0.50			
Age Risk			1:780	Nasal bone		Present
Biochemical T21 Risk			<1:10000	Sonographer		DR.SAHIL LOOMBA
Combined Trisomy 21 Ris	k		<1:10000	Qualifications in	measuring NT	MBBS,DNB
Trisomy 13/18			<1:10000			
		Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a				
1:10						
				low risk.	, , , , , , , , , , , , , , , , , , ,	,
		,			f the Trisomy 21 test	
1.100		expected that among more than 10000 women with the same data, there is one woman with a trisomy 21				
1 2 2		pregnancy.	s one woman with a	u150111y 21		
Th		The calculated risk by PRISCA depends on the accuracy of				
		the information provided by the referring physician. Please note that risk calculations are statistical approach and have no				
		diagnostic values!				
10 10 11 10 21 20 20 21 20 01 00 00 01 00 11 10 10 11 10		The patient combined risk presumes the NT measurement was done according to accepted guidelines				
			measurement was	done according to ac	cepted guidelines	
Trisomy 13/18 + NT The calculated risk for trise	omy 13/18 (with nuc	chal			
	translucency) is < 1:10000, which represents a low risk.					