

*Free Home Sample Collection 9999 778 778

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				Date of Report PRISCA	23-11-18 5.0.2.37
Patient Data					
Name		Mrs Priya	Patient ID		011811210145
Birthday		29-08-96	Sample ID		10357343
Age at delivery		22.2	Sample Date		22/11/18
Gestational age		11+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	46 Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown Origin		Asian		
Biochemical Data			Ultrasound D	ata	
Parameter	Value	Corr Mom	Gestational ag	e	11+5
PAPP-A	3.16 mIU/ml	0.78	Method		CRL (<>Hadlock)
fb-hCG	103.5 ng/ml	1.94	Scan Date		21-11-18
Risks at sampling date	•		CRL		49
Age Risk		1:1025	Nuchal translu	cency MoM	0.75
Biochemical T21 Risk		1:767	Nasal Bone		Present
Combined Trisomy 21 H	Risk	1:4459	Sonographer		DR.RAHUL
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	DNB
Risk Risk 1:10 1:100 1:50 1:50 1:100 1:100 1:100 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:10000 1:100000 1:10000 1:100000 1:10000 1:100000 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 low risk. After the result of the Trisomy 21 test it is expected that among 9533 women with the same data, there is one woman with a trisomy 21 pregnancy and 9532 women with not affected pregnancies. After the result of the Trisomy 21 test (with NT) it is expected that among 4459 women with the same data, there is one woman with a trisomy 21 pregnancy and 4458 women with not affected pregnancies. The patient combined risk presumes the NT measurement was done according to accepted guidelines The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value! Risk above Age Risk Risk below Age risk		