

\*Free Home Sample Collection 9999 778 778



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Date of Report 28/05/2019
PRISCA 5.0.2.37

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Patient Data					
Name		Mrs Sadhavi	Patient ID		011905260130
Birthday	18/03/1991		Sample ID		10431280
Age at delivery		28.5	Sample Date		26/05/2019
Gestational age		14+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	45 Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	14+0
AFP	33.63 ng/ml	1	Method		CRL (<>Robinson)
uE3	0.36  mIU/ml	1.33	Scan Date		
hCG	62659.4 ng/ml	1.14	Crown rump length (mm)		
Risks at sampling date			Nuchal translucency 0.8		
Age Risk		1:1117	Nuchal translucency MoM		0.42
Biochemical T21 Risk		1:3692	Nasal Bone		Present
Combined Trisomy 21 Risk		<1:10000	Sonographer		DR. Prakash Lal Chandani
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT		MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.		
1:1000 1:10000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age  Trisomy 13/18 + NT  The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			After the result of the Trisomy 21 test (with NT) it is expected that among 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies.  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 on the risk assessment! Calculated risks have no diagnostic values		