

\*Free Home Sample Collection 9999 778 778



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Date of Report	21-02-19
PRISCA	5 0 2 37

Patient Data						
Name	Name Mrs Diksha w/o Dinesh			Patient ID		011902190262
Birthday			17-02-88	Sample ID		10422432
Age at delivery			31.0	Sample Date		19/02/19
Gestational age by			13+1			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	61.3	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound D	ata	
Parameter	Value		Corr Mom	Gestational ag	e	13+0
PAPP-A	4.26	mIU/ml	1.01	Method		CRL (<>Hadlock)
fb-hCG	29.7	ng/ml	0.63	Scan Date		19-02-19
				CRL measurm	ents	66.0 mm
Risks at sampling date			Nuchal translucency MoM 0.65			
Age Risk			1:579	Nasal bone		Present
Biochemical T21 Risk			1:4654	Sonographer		Dr. Krishan Jangra
Combined trisomy 21 risk			<1:10000	Qualifications	in measuring NT	MBBS, DMRD, DNB
Trisomy 13/18			<1:10000			
Risk				Down's Syndi	rome Risk (Trisomy 21	Screening)
Risk 1:10			The calculated risk for Trisomy 21 (with nuchal			
1				translucency) is below the cut off, which indicates a low risk.		
1700 Acres	After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the			,		
1.100			same data, there is one woman with a trisomy 21			
1.000		pregnancy.	to is one woman with a ti	21		
		The calculated risk by PRISCA depends on the accuracy				
		of the information provided by the referring physician.				
		Please note that risk calculations are statistical				
THE RESIDENCE OF THE PROPERTY		approaches and have no diagnostic value!				
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age		The patient combined risk presumes the NT measurement was done according to accepted guidelines				
			The laboratory can not be hold responsible for their impact			
Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal			on the risk assessment! Calculated risks have no diagnostic value!			
translucency) is < 1:10000, v				valu <del>c</del> !		