

\*Free Home Sample Collection 9999 778 778



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Date of Report 7/6/2024
PRISCA 5.2.0.13

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Patient Data					
Name ASHA RANI		Patient ID 012406060		012406060176	
Birthday 1.		15/4/1990	Sample ID 11900386		
Age at sample 34		34.1	1 Sample Date 6/6/2024		
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+5
PAPP-A	$5.20~\mathrm{mIU/ml}$	0.87	Method		CRL (<>Robinson)
fb-hCG	44.3 ng/ml	1.24	Scan date		6/6/2024
Risks at sampling date			Crown rump length in mm 61.4		
Age Risk		1:328	Nuchal translucency MoM		0.94
Biochemical T21 risk		1:926	Nasal bone Pres		Present
Combined trisomy 21 risk		1:4249	Sonographer DR.VI		DR.VIPUL GOYAL
Trisomy 13/18 + NT <1:		<1:10000	Qualifications in measuring NT		MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:1000 1:1000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is < 1:10000,			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.  After the result of the Trisomy 21 test (with NT) it is expected that among 4249 women with the same data, there is one woman with a trisomy 21 pregnancy and 4248 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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).  The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
which indicates a low risk					