

\*Free Home Sample Collection 9999 778 778



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 Date of Report
 23-04-2024

 PRISCA
 5.2.0.13

Patient Data						
Name	MRS.HATHC	KIM	CHINGLOI F1	Patient ID		12404220124
Birthday			02-02-1997	Sample ID		12001550
Age at Sample date			27.2	Sample Date		22-04-2024
Gestational age 13+0						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	55	Diabete	es	NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+5
PAPP-A	6.6	mIU/m	d 0.49	Method		CRL (<>Robinson)
fb-hCG	93.4	ng/ml	1.28	Scan date		20-04-2024
Risks at sampling date				Crown rump length in mm 63.7		
Age Risk		1:860	Nuchal translucency MoM		0.73	
Biochemical T21 risk			1:560	Nasal bone Pres		Present
Combined trisomy 21 risk			1:3469	9 1		DR. DEEPIKA
Trisomy 13/18 + NT <1:10000  Risk				Qualifications in measuring NT MD  Down's Syndrome Risk (Trisomy 21 Screening)		
b				The calculated risk for Trisomy 21 (with NT) is below the		
1:100 1:1000 1:10000 1:10000 1:1010000			cut off, which represents a low risk.  After the result of the Trisomy 21 test (with NT) it is expected that among 3469 women with the same data, there is one woman with a trisomy 21 pregnancy and 3468 women with not affected pregnancies. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.  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;			
Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
which indicates a low	rick					<u> </u>
	Risk Above Cu	t Off		Risk above Age	e Risk	sk below Age risk