

*Free Home Sample Collection **9999 778 778** Download "MOLQ" App on

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				Date of Report	14-09-2024 5.2.0.13
Patient Data				PRISCA	3.2.0.13
Name	MRS. SAKS	HI KUAMARI	Patient ID		12409130204
Birthday			Sample ID		11966963
Age at Sample date	23.5		Sample Date		13-09-2024
Gestational age 13+4					
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58.3 Diabete	5	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+4
PAPP-A	7.3 mIU/ml	0.90	Method		CRL (<>Robinson)
fb-hCG	15.3 ng/ml	0.55	Scan date		13-09-2024
Risks at sampling date			Crown rump l	ength in mm	73.6
Age Risk	1:1050		Nuchal translucency MoM 1.0		
Biochemical T21 risk	<1:10000		Nasal bone		PRESENT
Combined trisomy 21 risk	sk <1:10000		Sonographer		DR. AMENDA
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
1:10 1:100 1:250 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:100		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 more 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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				Calculated risks have no o	diagnostic values



Risk Above Cut Off

