

 Date of Report
 10/6/2023

 PRISCA
 5.1.0.17

Patient Data					
Name	N	ARS. SUMAN	Patient ID		012306080457
Birthday		5/1/1989	Sample ID		11660552
Age at term		34.1	Sample Date		8/6/2023
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	74 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+6
PAPP-A	$4.82~\mathrm{mIU/ml}$	0.91	Method		CRL (<>Robinson)
fb-hCG	75.4  ng/ml	2.56	Scan date		7/6/2023
Risks at sampling date			Nuchal translucency (NT) 1.1		
Age Risk		1:315	Nuchal translucency MoM 0.6		0.66
Biochemical T21 risk 1:1		1:162	Nasal bone preser		present
Combined trisomy 21 r	isk	1:938			
Trisomy 13/18		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:1000 1:3 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			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 it is expected that among 938 women with the same data, there is one woman with a trisomy 21 pregnancy and 937 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		
	k Above Cut Off		Risk above Age	e Risk	Risk below Age risk